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Amylases, nucleic acids encoding them and methods for making and using them

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ABSTRACT

AMYLASES, NUCLEIC ACIDS ENCODING THEM AND
METHODS FOR MAKING AND USING THEM

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In one aspect, the invention is directed to polypeptides having an amylase activity, polynucleotides encoding the polypeptides, and methods for making and using these polynucleotides and polypeptides. In one aspect, the polypeptides of the invention can be used as amylases, for example, alpha amylases, to catalyze the hydrolysis of starch
10 into sugars. In one aspect, the invention provides delayed release compositions comprising an desired ingredient coated by a latex polymer coating.

AUSTRALIA

Patents Act 1990

COMPLETE SPECIFICATION STANDARD PATENT

Invention Title: **Amylases, nucleic acids encoding them and methods for making and using them**

The following statement is a full description of this invention, including the best method of performing it known to us:

AMYLASES, NUCLEIC ACIDS ENCODING THEM AND METHODS FOR MAKING AND USING THEM

REFERENCE TO SEQUENCE LISTING SUBMITTED ON A COMPACT DISC

This application includes a compact disc (submitted in quadruplicate) containing a sequence listing. The entire content of the sequence listing is herein incorporated by reference. The sequence listing is identified on the compact disc as follows.

File Name	Date of Creation	Size (bytes)
Sequence Listing.txt	March 4, 2004	1,798,144

TECHNICAL FIELD

This invention relates to molecular and cellular biology and biochemistry.

In one aspect, the invention is directed to polypeptides having an amylase activity, polynucleotides encoding the polypeptides, and methods for making and using these polynucleotides and polypeptides. In one aspect, the polypeptides of the invention can be used as amylases, for example, alpha amylases or glucoamylases, to catalyze the hydrolysis of starch into sugars. In one aspect, the invention is directed to polypeptides having thermostable amylase activity, such as alpha amylases or glucoamylase activity, e.g., a 1,4-alpha-D-glucan glucohydrolase activity. In one aspect, the polypeptides of the invention can be used as amylases, for example, alpha amylases or glucoamylases, to catalyze the hydrolysis of starch into sugars, such as glucose. The invention is also directed to nucleic acid constructs, vectors, and host cells comprising the nucleic acid sequences of the invention as well as recombinant methods for producing the polypeptides of the invention. The invention is also directed to the use of amylases of the invention in starch conversion processes, including production of high fructose corn syrup (HFCS), ethanol, dextrose, and dextrose syrups.

BACKGROUND

Starch is a complex carbohydrate often found in the human diet. The structure of starch is glucose polymers linked by α -1,4 and α -1,6 glucosidic bonds. Amylase is an enzyme that catalyzes the hydrolysis of starches into sugars. Amylases

hydrolyze internal α -1,4-glucosidic linkages in starch, largely at random, to produce smaller molecular weight malto-dextrins. The breakdown of starch is important in the digestive system and commercially. Amylases are of considerable commercial value, being used in the initial stages (liquefaction) of starch processing; in wet corn milling; in alcohol production; as cleaning agents in detergent matrices; in the textile industry for starch desizing; in baking applications; in the beverage industry; in oilfields in drilling processes; in inking of recycled paper; and in animal feed.

Amylases are produced by a wide variety of microorganisms including *Bacillus* and *Aspergillus*, with most commercial amylases being produced from bacterial sources such as *Bacillus licheniformis*, *Bacillus amyloliquefaciens*, *Bacillus subtilis*, or *Bacillus stearothermophilus*. In recent years, the enzymes in commercial use have been those from *Bacillus licheniformis* because of their heat stability and performance, at least at neutral and mildly alkaline pHs.

Commercially, glucoamylases are used to further hydrolyze cornstarch, which has already been partially hydrolyzed with an alpha-amylase. The glucose produced in this reaction may then be converted to a mixture of glucose and fructose by a glucose isomerase enzyme. This mixture, or one enriched with fructose, is the high fructose corn syrup commercialized throughout the world. In general, starch to fructose processing consists of four steps: liquefaction of granular starch, saccharification of the liquefied starch into dextrose, purification, and isomerization to fructose. The object of a starch liquefaction process is to convert a concentrated suspension of starch polymer granules into a solution of soluble shorter chain length dextrins of low viscosity.

The most widely utilized glucoamylase is produced from the fungus *Aspergillus niger*. One of the problems with the commercial use of this enzyme is its relatively low thermostability. A number of other fungal glucoamylases have been reported, including *Rizopus*, *Thielavia*, *Thermoascus* and *Talaromyces*, and a glucoamylase from the thermophilic fungus *Thermomyces lanuginosus*.

In general, starch to fructose processing consists of four steps: liquefaction of granular starch, saccharification of the liquefied starch into dextrose, purification, and isomerization to fructose. The object of a starch liquefaction process is to convert a concentrated suspension of starch polymer granules into a solution of soluble shorter chain length dextrins of low viscosity. This step is essential for convenient handling with standard equipment and for efficient conversion to glucose or other sugars. To liquefy

granular starch, it is necessary to gelatinize the granules by raising the temperature of the granular starch to over about 72°C. The heating process instantaneously disrupts the insoluble starch granules to produce a water soluble starch solution. The solubilized starch solution is then liquefied by amylase. A starch granule is composed of: 69-74% amylopectin, 26-31% amylose, 11-14% water, 0.2-0.4% protein, 0.5-0.9% lipid, 0.05-0.1% ash, 0.02-0.03% phosphorus, 0.1% pentosan. Approximately 70% of a granule is amorphous and 30% is crystalline.

A common enzymatic liquefaction process involves adjusting the pH of a granular starch slurry to between 6.0 and 6.5, the pH optimum of alpha-amylase derived from *Bacillus licheniformis*, with the addition of calcium hydroxide, sodium hydroxide or sodium carbonate. The addition of calcium hydroxide has the advantage of also providing calcium ions which are known to stabilize the alpha-amylase against inactivation. Upon addition of alpha-amylase, the suspension is pumped through a steam jet to instantaneously raise the temperature to between 80°C to 115°C. The starch is immediately gelatinized and, due to the presence of alpha-amylase, depolymerized through random hydrolysis of a (1-4) glycosidic bonds by alpha-amylase to a fluid mass which is easily pumped.

In a second variation to the liquefaction process, alpha-amylase is added to the starch suspension, the suspension is held at a temperature of 80-100°C to partially hydrolyze the starch granules, and the partially hydrolyzed starch suspension is pumped through a jet at temperatures in excess of about 105°C to thoroughly gelatinize any remaining granular structure. After cooling the gelatinized starch, a second addition of alpha-amylase can be made to further hydrolyze the starch.

A third variation of this process is called the dry milling process. In dry milling, whole grain is ground and combined with water. The germ is optionally removed by flotation separation or equivalent techniques. The resulting mixture, which contains starch, fiber, protein and other components of the grain, is liquefied using alpha-amylase. The general practice in the art is to undertake enzymatic liquefaction at a lower temperature when using the dry milling process. Generally, low temperature liquefaction is believed to be less efficient than high temperature liquefaction in converting starch to soluble dextrans.

Typically, after gelatinization the starch solution is held at an elevated temperature in the presence of alpha-amylase until a DE of 10-20 is achieved, usually a

period of 1-3 hours. Dextrose equivalent (DE) is the industry standard for measuring the concentration of total reducing sugars, calculated as D-glucose on a dry weight basis. Unhydrolyzed granular starch has a DE of virtually zero, whereas the DE of D-glucose is defined as 100.

5 Corn wet milling is a process which produces corn oil, gluten meal, gluten feed and starch. Alkaline-amylase is used in the liquefaction of starch and glucoamylase is used in saccharification, producing glucose. Corn, a kernel of which consists of a outer seed coat (fiber), starch, a combination of starch and glucose and the inner germ, is subjected to a four step process, which results in the production of starch. The corn is
10 steeped, de-germed, de-fibered, and finally the gluten is separated. In the steeping process, the solubles are taken out. The product remaining after removal of the solubles is de-germed, resulting in production of corn oil and production of an oil cake, which is added to the solubles from the steeping step. The remaining product is de-fibered and the fiber solids are added to the oil cake/solubles mixture. This mixture of fiber solids, oil
15 cake and solubles forms a gluten feed. After de-fibered, the remaining product is subjected to gluten separation. This separation results in a gluten meal and starch. The starch is then subjected to liquefaction and saccharification to produce glucose.

Staling of baked products (such as bread) has been recognized as a problem which becomes more serious as more time lies between the moment of
20 preparation of the bread product and the moment of consumption. The term staling is used to describe changes undesirable to the consumer in the properties of the bread product after leaving the oven, such as an increase of the firmness of the crumb, a decrease of the elasticity of the crumb, and changes in the crust, which becomes tough and leathery. The firmness of the bread crumb increases further during storage up to a
25 level, which is considered as negative. The increase in crumb firmness, which is considered as the most important aspect of staling, is recognized by the consumer a long time before the bread product has otherwise become unsuitable for consumption.

There is a need in the industry for the identification and optimization of amylases, useful for various uses, including commercial cornstarch liquefaction
30 processes. These second generation acid amylases will offer improved manufacturing and/or performance characteristics over the industry standard enzymes from *Bacillus licheniformis*, for example.

There is also a need for the identification and optimization of amylases having utility in automatic dish wash (ADW) products and laundry detergent. In ADW products, the amylase will function at pH 10-11 and at 45-60°C in the presence of calcium chelators and oxidative conditions. For laundry, activity at pH 9-10 and 40°C in the appropriate detergent matrix will be required. Amylases are also useful in textile desizing, brewing processes, starch modification in the paper and pulp industry and other processes described in the art.

Amylases can be used commercially in the initial stages (liquefaction) of starch processing; in wet corn milling; in alcohol production; as cleaning agents in detergent matrices; in the textile industry for starch desizing; in baking applications; in the beverage industry; in oilfields in drilling processes; in inking of recycled paper and in animal feed. Amylases are also useful in textile desizing, brewing processes, starch modification in the paper and pulp industry and other processes.

The publications discussed herein are provided solely for their disclosure prior to the filing date of the present application. Nothing herein is to be construed as an admission that the invention is not entitled to antedate such disclosure by virtue of prior invention.

SUMMARY

The invention provides isolated or recombinant nucleic acids comprising a nucleic acid sequence having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to a nucleic acid of the invention, e.g., an exemplary nucleic acid of the invention, over a region of at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more, residues. In one aspect, the nucleic acid encodes at least one polypeptide having an amylase activity, and the sequence identities are determined by analysis with a sequence comparison algorithm or by a visual inspection. In another aspect, the invention provides nucleic acids for use as probes, inhibitory molecules (e.g., antisense, iRNAs), transcriptional or translational regulation, and the like. Exemplary

nucleic acids of the invention include isolated or recombinant nucleic acids comprising a nucleic acid sequence as set forth in SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:9, SEQ ID NO:11, SEQ ID NO:13, SEQ ID NO:15, SEQ ID NO:17, SEQ ID NO:19, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:25, SEQ ID NO:27, SEQ ID NO:29, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:35, SEQ ID NO:37, SEQ ID NO:39, SEQ ID NO:41, SEQ ID NO:43, SEQ ID NO:45, SEQ ID NO:47, SEQ ID NO:49, SEQ ID NO:51, SEQ ID NO:53, SEQ ID NO:55, SEQ ID NO:57, SEQ ID NO:59, SEQ ID NO:61, SEQ ID NO:63, SEQ ID NO:65, SEQ ID NO:67, SEQ ID NO:69, SEQ ID NO:71, SEQ ID NO:73, SEQ ID NO:75, SEQ ID NO:77, SEQ ID NO:79, SEQ ID NO:81, SEQ ID NO:83, SEQ ID NO:85, SEQ ID NO:87, SEQ ID NO:89, SEQ ID NO:91, SEQ ID NO:93, SEQ ID NO:95, SEQ ID NO:97, SEQ ID NO:99, SEQ ID NO:101, SEQ ID NO:103, SEQ ID NO:105, SEQ ID NO:107, SEQ ID NO:109, SEQ ID NO:111, SEQ ID NO:113, SEQ ID NO:115, SEQ ID NO:117, SEQ ID NO:119, SEQ ID NO:121, SEQ ID NO:123, SEQ ID NO:125, SEQ ID NO:127, SEQ ID NO:129, SEQ ID NO:131, SEQ ID NO:133, SEQ ID NO:135, SEQ ID NO:137, SEQ ID NO:139, SEQ ID NO:141, SEQ ID NO:143, SEQ ID NO:145, SEQ ID NO:147, SEQ ID NO:149, SEQ ID NO:151, SEQ ID NO:153, SEQ ID NO:155, SEQ ID NO:157, SEQ ID NO:159, SEQ ID NO:161, SEQ ID NO:163, SEQ ID NO:165, SEQ ID NO:167, SEQ ID NO:189, SEQ ID NO:191, SEQ ID NO:193, SEQ ID NO:203, SEQ ID NO:205, SEQ ID NO:207, SEQ ID NO:209, SEQ ID NO:211, SEQ ID NO:322, SEQ ID NO:324, SEQ ID NO:326, SEQ ID NO:328, SEQ ID NO:330, SEQ ID NO:332, SEQ ID NO:334, SEQ ID NO:336, SEQ ID NO:338, SEQ ID NO:340, SEQ ID NO:342, SEQ ID NO:344, SEQ ID NO:346, SEQ ID NO:348, SEQ ID NO:350, SEQ ID NO:352, SEQ ID NO:354, SEQ ID NO:356, SEQ ID NO:358, SEQ ID NO:360, SEQ ID NO:362, SEQ ID NO:364, SEQ ID NO:366, SEQ ID NO:368, SEQ ID NO:370, SEQ ID NO:372, SEQ ID NO:374, SEQ ID NO:376, SEQ ID NO:378, SEQ ID NO:380, SEQ ID NO:382, SEQ ID NO:384, SEQ ID NO:386, SEQ ID NO:388, SEQ ID NO:390, SEQ ID NO:392, SEQ ID NO:394, SEQ ID NO:396, SEQ ID NO:398, SEQ ID NO:400, SEQ ID NO:402, SEQ ID NO:404, SEQ ID NO:406, SEQ ID NO:408, SEQ ID NO:410, SEQ ID NO:412, SEQ ID NO:414, SEQ ID NO:416, SEQ ID NO:418, SEQ ID NO:420, SEQ ID NO:422, SEQ ID NO:424, SEQ ID NO:426, SEQ ID NO:428, SEQ ID NO:430, SEQ ID NO:432, SEQ ID NO:434, SEQ ID NO:436, SEQ ID NO:438, SEQ ID NO:440, SEQ ID NO:442, SEQ ID NO:444, SEQ ID NO:446, SEQ ID NO:448, SEQ ID NO:450, SEQ ID NO:452, SEQ ID NO:454, SEQ ID

NO:456, SEQ ID NO:458, SEQ ID NO:460, SEQ ID NO:460, SEQ ID NO:462, SEQ ID NO:465, SEQ ID NO:467, SEQ ID NO:473, SEQ ID NO:475, SEQ ID NO:478, SEQ ID NO:480, SEQ ID NO:484, SEQ ID NO:486, SEQ ID NO:492, SEQ ID NO:494, SEQ ID NO:498, SEQ ID NO:500, SEQ ID NO:509, SEQ ID NO:511, SEQ ID NO:515, SEQ ID NO:517, SEQ ID NO:517, SEQ ID NO:519, SEQ ID NO:522, SEQ ID NO:524, SEQ ID NO:527, SEQ ID NO:529, SEQ ID NO:532, SEQ ID NO:534, SEQ ID NO:539, SEQ ID NO:541, SEQ ID NO:544, SEQ ID NO:546, SEQ ID NO:552, SEQ ID NO:554, SEQ ID NO:558, SEQ ID NO:560, SEQ ID NO:565, SEQ ID NO:567, SEQ ID NO:569, SEQ ID NO:571, SEQ ID NO:573, SEQ ID NO:575, SEQ ID NO:577, SEQ ID NO:579, SEQ ID NO:581, SEQ ID NO:583, SEQ ID NO:585, SEQ ID NO:587, SEQ ID NO:593, SEQ ID NO:603, SEQ ID NO:605, SEQ ID NO:607, SEQ ID NO:609, SEQ ID NO:611, SEQ ID NO:613, SEQ ID NO:615, SEQ ID NO:617, SEQ ID NO:619 or SEQ ID NO:621, and subsequences thereof, e.g., at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500 or more residues in length, or over the full length of a gene or transcript.

Exemplary nucleic acids of the invention also include isolated or recombinant nucleic acids encoding a polypeptide of the invention, e.g., an exemplary polypeptide having a sequence as set forth in SEQ ID NO:2, SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:10, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:16, SEQ ID NO:18, SEQ ID NO:20, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:40, SEQ ID NO:42, SEQ ID NO:44, SEQ ID NO:46, SEQ ID NO:48, SEQ ID NO:50, SEQ ID NO:52, SEQ ID NO:54, SEQ ID NO:56, SEQ ID NO:58, SEQ ID NO:60, SEQ ID NO:62, SEQ ID NO:64, SEQ ID NO:66, SEQ ID NO:68, SEQ ID NO:70, SEQ ID NO:72, SEQ ID NO:74, SEQ ID NO:76, SEQ ID NO:78, SEQ ID NO:80, SEQ ID NO:82, SEQ ID NO:84, SEQ ID NO:86, SEQ ID NO:88, SEQ ID NO:90, SEQ ID NO:92, SEQ ID NO:94, SEQ ID NO:96, SEQ ID NO:98, SEQ ID NO:100, SEQ ID NO:102, SEQ ID NO:104, SEQ ID NO:106, SEQ ID NO:108, SEQ ID NO:110, SEQ ID NO:112, SEQ ID NO:114, SEQ ID NO:116, SEQ ID NO:118, SEQ ID NO:120, SEQ ID NO:122, SEQ ID NO:124, SEQ ID NO:126, SEQ ID NO:128, SEQ ID NO:130, SEQ ID NO:132, SEQ ID NO:134, SEQ ID NO:136, SEQ ID NO:138, SEQ ID NO:140, SEQ ID NO:142, SEQ ID NO:144, SEQ ID NO:146, SEQ ID NO:148, SEQ ID

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In one aspect, the invention also provides amylase-encoding nucleic acids with a common novelty in that they are derived from mixed cultures. The invention provides amylase-encoding nucleic acids isolated from mixed cultures comprising a nucleic acid sequence having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an exemplary nucleic acid of the invention over a region of at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more, residues, wherein the nucleic acid encodes at least one polypeptide having an amylase activity, and the sequence identities are determined by analysis with a sequence comparison algorithm or by a visual inspection. In one aspect, the invention provides amylase-encoding nucleic acids isolated from mixed cultures comprising a nucleic acid of the invention, e.g., an exemplary nucleic acid of the invention, e.g., a sequence as set forth in SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:9, SEQ ID NO:11, etc., and subsequences thereof, e.g., at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500 or more residues in length, or over the full length of a gene or transcript; or, a nucleic acid encoding a polypeptide of the invention.

In one aspect, the invention also provides amylase-encoding nucleic acids with a common novelty in that they are derived from environmental sources, e.g., mixed

environmental sources. In one aspect, the invention provides amylase-encoding nucleic acids isolated from environmental sources, e.g., mixed environmental sources, comprising a nucleic acid sequence having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an exemplary nucleic acid of the invention over a region of at least about 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more, residues, wherein the nucleic acid encodes at least one polypeptide having an amylase activity, and the sequence identities are determined by analysis with a sequence comparison algorithm or by a visual inspection. In one aspect, the invention provides amylase-encoding nucleic acids isolated from environmental sources, e.g., mixed environmental sources, comprising a nucleic acid of the invention, e.g., an exemplary nucleic acid sequence of the invention as set forth in SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:9, SEQ ID NO:11, etc., SEQ ID NO:583, SEQ ID NO:585, and subsequences thereof, e.g., at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500 or more residues in length, or over the full length of a gene or transcript; or, a nucleic acid encoding a polypeptide of the invention.

In one aspect, the invention also provides amylases, and amylase-encoding nucleic acids, with a common novelty in that they are derived from archael sources, including the archael-derived amylases of SEQ ID NO:80 (encoded by SEQ ID NO:79), SEQ ID NO:82 (encoded by SEQ ID NO:81), SEQ ID NO:116 (encoded by SEQ ID NO:115), SEQ ID NO:323 (encoded by SEQ ID NO:322), SEQ NO: 570 (encoded by SEQ ID NO:169).

In one aspect, the sequence comparison algorithm is a BLAST version 2.2.2 algorithm where a filtering setting is set to blastall -p blastp -d "nr pataa" -F F, and all other options are set to default.

Another aspect of the invention is an isolated or recombinant nucleic acid including at least 10 consecutive bases of a nucleic acid sequence of the invention, sequences substantially identical thereto, and the sequences complementary thereto.

In one aspect, the amylase activity comprises α -amylase activity, including the ability to hydrolyze internal α -1,4-glucosidic linkages in starch to produce smaller molecular weight malto-dextrins. In one aspect, the α -amylase activity includes hydrolyzing internal α -1,4-glucosidic linkages in starch at random. The amylase activity can comprise an α -amylase activity, a β -amylase activity, a glucoamylase activity, a 1,4- α -D-glucan glucohydrolase activity, an exoamylase activity, a glucan α -maltotetrahydrolase activity, a maltase activity, an isomaltase activity, a glucan 1, 4, α -glucosidase activity, an α -glucosidase activity, a sucrase activity or an agarase activity (e.g., a β -agarase activity).

The amylase activity can comprise hydrolyzing glucosidic bonds. In one aspect, the glucosidic bonds comprise an α -1,4-glucosidic bond. In another aspect, the glucosidic bonds comprise an α -1,6-glucosidic bond. In one aspect, the amylase activity comprises hydrolyzing glucosidic bonds in starch, e.g., liquefied starch. The amylase activity can further comprise hydrolyzing glucosidic bonds into maltodextrins. In one aspect, the amylase activity comprises cleaving a maltose or a D-glucose unit from non-reducing end of the starch.

In one aspect, the isolated or recombinant nucleic acid encodes a polypeptide having an amylase activity which is thermostable. The polypeptide can retain an amylase activity under conditions comprising a temperature range of anywhere between about 0°C to about 37°C, or, between about 37°C to about 95°C or more, e.g., 98°C, 100°C or more; between about 55°C to about 85°C, between about 70°C to about 95°C, or, between about 90°C to about 95°C. For example, the exemplary polypeptide having a sequence as set forth in SEQ ID NO:437 is thermostable, retaining 50% activity after 25 minutes at 100°C in the absence of added calcium.

In another aspect, the isolated or recombinant nucleic acid encodes a polypeptide having an amylase activity which is thermotolerant. The polypeptide can retain an amylase activity after exposure to a temperature in the range from greater than 37°C to about 95°C or anywhere in the range from greater than 55°C to about 85°C. In one aspect, the polypeptide retains an amylase activity after exposure to a temperature in the range from greater than 90°C to about 95°C at pH 4.5.

The invention provides isolated or recombinant nucleic acids comprising a sequence that hybridizes under stringent conditions to a nucleic acid of the invention, e.g., an exemplary nucleic acid of the invention, a nucleic acid comprising a sequence as set

forth in SEQ ID NO:1, SEQ ID NO:3, SEQ ID NO:5, SEQ ID NO:9, SEQ ID NO:11,
 SEQ ID NO:13, SEQ ID NO:15, SEQ ID NO:17, SEQ ID NO:19, SEQ ID NO:21, SEQ
 ID NO:23, SEQ ID NO:25, SEQ ID NO:27, SEQ ID NO:29, SEQ ID NO:31, SEQ ID
 NO:33, SEQ ID NO:35, SEQ ID NO:37, SEQ ID NO:39, SEQ ID NO:41, SEQ ID
 NO:43, SEQ ID NO:45, SEQ ID NO:47, SEQ ID NO:49, SEQ ID NO:51, SEQ ID
 NO:53, SEQ ID NO:55, SEQ ID NO:57, SEQ ID NO:59, SEQ ID NO:61, SEQ ID
 NO:63, SEQ ID NO:65, SEQ ID NO:67, SEQ ID NO:69, SEQ ID NO:71, SEQ ID
 NO:73, SEQ ID NO:75, SEQ ID NO:77, SEQ ID NO:79, SEQ ID NO:81, SEQ ID
 NO:83, SEQ ID NO:85, SEQ ID NO:87, SEQ ID NO:89, SEQ ID NO:91, SEQ ID
 NO:93, SEQ ID NO:95, SEQ ID NO:97, SEQ ID NO:99, SEQ ID NO:101, SEQ ID
 NO:103, SEQ ID NO:105, SEQ ID NO:107, SEQ ID NO:109, SEQ ID NO:111, SEQ ID
 NO:113, SEQ ID NO:115, SEQ ID NO:117, SEQ ID NO:119, SEQ ID NO:121, SEQ ID
 NO:123, SEQ ID NO:125, SEQ ID NO:127, SEQ ID NO:129, SEQ ID NO:131, SEQ ID
 NO:133, SEQ ID NO:135, SEQ ID NO:137, SEQ ID NO:139, SEQ ID NO:141, SEQ ID
 NO:143, SEQ ID NO:145, SEQ ID NO:147, SEQ ID NO:149, SEQ ID NO:151, SEQ ID
 NO:153, SEQ ID NO:155, SEQ ID NO:157, SEQ ID NO:159, SEQ ID NO:161, SEQ ID
 NO:163, SEQ ID NO:165, SEQ ID NO:167, SEQ ID NO:189, SEQ ID NO:191, SEQ ID
 NO:193, SEQ ID NO:203, SEQ ID NO:205, SEQ ID NO:207, SEQ ID NO:209, SEQ ID
 NO:211, SEQ ID NO:322, SEQ ID NO:324, SEQ ID NO:326, SEQ ID NO:328, SEQ ID
 NO:330, SEQ ID NO:332, SEQ ID NO:334, SEQ ID NO:336, SEQ ID NO:338, SEQ ID
 NO:340, SEQ ID NO:342, SEQ ID NO:344, SEQ ID NO:346, SEQ ID NO:348, SEQ ID
 NO:350, SEQ ID NO:352, SEQ ID NO:354, SEQ ID NO:356, SEQ ID NO:358, SEQ ID
 NO:360, SEQ ID NO:362, SEQ ID NO:364, SEQ ID NO:366, SEQ ID NO:368, SEQ ID
 NO:370, SEQ ID NO:372, SEQ ID NO:374, SEQ ID NO:376, SEQ ID NO:378, SEQ ID
 NO:380, SEQ ID NO:382, SEQ ID NO:384, SEQ ID NO:386, SEQ ID NO:388, SEQ ID
 NO:390, SEQ ID NO:392, SEQ ID NO:394, SEQ ID NO:396, SEQ ID NO:398, SEQ ID
 NO:400, SEQ ID NO:402, SEQ ID NO:404, SEQ ID NO:406, SEQ ID NO:408, SEQ ID
 NO:410, SEQ ID NO:412, SEQ ID NO:414, SEQ ID NO:416, SEQ ID NO:418, SEQ ID
 NO:420, SEQ ID NO:422, SEQ ID NO:424, SEQ ID NO:426, SEQ ID NO:428, SEQ ID
 NO:430, SEQ ID NO:432, SEQ ID NO:434, SEQ ID NO:436, SEQ ID NO:438, SEQ ID
 NO:440, SEQ ID NO:442, SEQ ID NO:444, SEQ ID NO:446, SEQ ID NO:448, SEQ ID
 NO:450, SEQ ID NO:452, SEQ ID NO:454, SEQ ID NO:456, SEQ ID NO:458, SEQ ID
 NO:460, SEQ ID NO:460, SEQ ID NO:462, SEQ ID NO:465, SEQ ID NO:467, SEQ ID

NO:473, SEQ ID NO:475, SEQ ID NO:478, SEQ ID NO:480, SEQ ID NO:484, SEQ ID NO:486, SEQ ID NO:492, SEQ ID NO:494, SEQ ID NO:498, SEQ ID NO:500, SEQ ID NO:509, SEQ ID NO:511, SEQ ID NO:515, SEQ ID NO:517, SEQ ID NO:517, SEQ ID NO:519, SEQ ID NO:522, SEQ ID NO:524, SEQ ID NO:527, SEQ ID NO:529, SEQ ID NO:532, SEQ ID NO:534, SEQ ID NO:539, SEQ ID NO:541, SEQ ID NO:544, SEQ ID NO:546, SEQ ID NO:552, SEQ ID NO:554, SEQ ID NO:558, SEQ ID NO:560, SEQ ID NO:565, SEQ ID NO:567, SEQ ID NO:569, SEQ ID NO:571, SEQ ID NO:573, SEQ ID NO:575, SEQ ID NO:577, SEQ ID NO:579, SEQ ID NO:581, SEQ ID NO:583, SEQ ID NO:585, SEQ ID NO:587, SEQ ID NO:593, SEQ ID NO:603, SEQ ID NO:605, SEQ ID NO:607, SEQ ID NO:609, SEQ ID NO:611, SEQ ID NO:613, SEQ ID NO:615, SEQ ID NO:617, SEQ ID NO:619 or SEQ ID NO:621, or fragments or subsequences thereof. In one aspect, the nucleic acid encodes a polypeptide having an amylase activity. The nucleic acid can be at least about 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500 or more residues in length or the full length of the gene or transcript. In one aspect, the stringent conditions include a wash step comprising a wash in 0.2X SSC at a temperature of about 65°C for about 15 minutes.

The invention provides a nucleic acid probe for identifying a nucleic acid encoding a polypeptide having an amylase activity, wherein the probe comprises at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 95, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000 or more, consecutive bases of a sequence comprising a sequence of the invention, or fragments or subsequences thereof, wherein the probe identifies the nucleic acid by binding or hybridization. The probe can comprise an oligonucleotide comprising at least about 10 to 50, about 20 to 60, about 30 to 70, about 40 to 80, or about 60 to 100 consecutive bases of a sequence comprising a sequence of the invention, or fragments or subsequences thereof.

The invention provides a nucleic acid probe for identifying a nucleic acid encoding a polypeptide having an amylase activity, wherein the probe comprises a nucleic acid comprising a sequence at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 90, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000 or more residues having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%,

82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to a nucleic acid of the invention, wherein the sequence identities are determined by analysis with a sequence comparison algorithm or by visual inspection.

The probe can comprise an oligonucleotide comprising at least about 10 to 50, about 20 to 60, about 30 to 70, about 40 to 80, or about 60 to 100 consecutive bases of a nucleic acid sequence of the invention, or a subsequence thereof.

The invention provides an amplification primer sequence pair for amplifying a nucleic acid encoding a polypeptide having an amylase activity, wherein the primer pair is capable of amplifying a nucleic acid comprising a sequence of the invention, or fragments or subsequences thereof. One or each member of the amplification primer sequence pair can comprise an oligonucleotide comprising at least about 10 to 50 consecutive bases of the sequence.

The invention provides methods of amplifying a nucleic acid encoding a polypeptide having an amylase activity comprising amplification of a template nucleic acid with an amplification primer sequence pair capable of amplifying a nucleic acid sequence of the invention, or fragments or subsequences thereof.

The invention provides expression cassettes comprising a nucleic acid of the invention or a subsequence thereof. In one aspect, the expression cassette can comprise the nucleic acid that is operably linked to a promoter. The promoter can be a viral, bacterial, mammalian or plant promoter. In one aspect, the plant promoter can be a potato, rice, corn, wheat, tobacco or barley promoter. The promoter can be a constitutive promoter. The constitutive promoter can comprise CaMV35S. In another aspect, the promoter can be an inducible promoter. In one aspect, the promoter can be a tissue-specific promoter or an environmentally regulated or a developmentally regulated promoter. Thus, the promoter can be, e.g., a seed-specific, a leaf-specific, a root-specific, a stem-specific or an abscission-induced promoter. In one aspect, the expression cassette can further comprise a plant or plant virus expression vector.

The invention provides cloning vehicles comprising an expression cassette (e.g., a vector) of the invention or a nucleic acid of the invention. The cloning vehicle can be a viral vector, a plasmid, a phage, a phagemid, a cosmid, a fosmid, a bacteriophage or an artificial chromosome. The viral vector can comprise an adenovirus vector, a retroviral vector or an adeno-associated viral vector. The cloning vehicle can comprise a

bacterial artificial chromosome (BAC), a plasmid, a bacteriophage P1-derived vector (PAC), a yeast artificial chromosome (YAC), or a mammalian artificial chromosome (MAC).

5 The invention provides transformed cell comprising a nucleic acid of the invention or an expression cassette (e.g., a vector) of the invention, or a cloning vehicle of the invention. In one aspect, the transformed cell can be a bacterial cell, a mammalian cell, a fungal cell, a yeast cell, an insect cell or a plant cell. In one aspect, the plant cell can be a potato, wheat, rice, corn, tobacco or barley cell.

10 The invention provides transgenic non-human animals comprising a nucleic acid of the invention or an expression cassette (e.g., a vector) of the invention. In one aspect, the animal is a mouse.

The invention provides transgenic plants comprising a nucleic acid of the invention or an expression cassette (e.g., a vector) of the invention. The transgenic plant can be a corn plant, a potato plant, a tomato plant, a wheat plant, an oilseed plant, a rapeseed plant, a soybean plant, a rice plant, a barley plant or a tobacco plant.

15 The invention provides transgenic seeds comprising a nucleic acid of the invention or an expression cassette (e.g., a vector) of the invention. The transgenic seed can be a corn seed, a wheat kernel, an oilseed, a rapeseed, a soybean seed, a palm kernel, a sunflower seed, a sesame seed, a peanut or a tobacco plant seed.

20 The invention provides an antisense oligonucleotide comprising a nucleic acid sequence complementary to or capable of hybridizing under stringent conditions to a nucleic acid of the invention. The invention provides methods of inhibiting the translation of an amylase message in a cell comprising administering to the cell or expressing in the cell an antisense oligonucleotide comprising a nucleic acid sequence complementary to or capable of hybridizing under stringent conditions to a nucleic acid of the invention.

25 The invention provides an isolated or recombinant polypeptide comprising an amino acid sequence having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an exemplary polypeptide or peptide of the invention over a region of at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70,

75, 80, 90, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more residues, or over the full length of the polypeptide, and the sequence identities are determined by analysis with a sequence comparison algorithm or by a visual inspection. Exemplary polypeptide or peptide sequences of the invention include SEQ ID NO:2, SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:10, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:16, SEQ ID NO:18, SEQ ID NO:20, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:40, SEQ ID NO:42, SEQ ID NO:44, SEQ ID NO:46, SEQ ID NO:48, SEQ ID NO:50, SEQ ID NO:52, SEQ ID NO:54, SEQ ID NO:56, SEQ ID NO:58, SEQ ID NO:60, SEQ ID NO:62, SEQ ID NO:64, SEQ ID NO:66, SEQ ID NO:68, SEQ ID NO:70, SEQ ID NO:72, SEQ ID NO:74, SEQ ID NO:76, SEQ ID NO:78, SEQ ID NO:80, SEQ ID NO:82, SEQ ID NO:84, SEQ ID NO:86, SEQ ID NO:88, SEQ ID NO:90, SEQ ID NO:92, SEQ ID NO:94, SEQ ID NO:96, SEQ ID NO:98, SEQ ID NO:100, SEQ ID NO:102, SEQ ID NO:104, SEQ ID NO:106, SEQ ID NO:108, SEQ ID NO:110, SEQ ID NO:112, SEQ ID NO:114, SEQ ID NO:116, SEQ ID NO:118, SEQ ID NO:120, SEQ ID NO:122, SEQ ID NO:124, SEQ ID NO:126, SEQ ID NO:128, SEQ ID NO:130, SEQ ID NO:132, SEQ ID NO:134, SEQ ID NO:136, SEQ ID NO:138, SEQ ID NO:140, SEQ ID NO:142, SEQ ID NO:144, SEQ ID NO:146, SEQ ID NO:148, SEQ ID NO:150, SEQ ID NO:152, SEQ ID NO:154, SEQ ID NO:156, SEQ ID NO:158, SEQ ID NO:160, SEQ ID NO:162, SEQ ID NO:164, SEQ ID NO:166, SEQ ID NO:168, SEQ ID NO:190, SEQ ID NO:192, SEQ ID NO:194, SEQ ID NO:204, SEQ ID NO:206, SEQ ID NO:208, SEQ ID NO:210, SEQ ID NO:212, SEQ ID NO:323, SEQ ID NO:325, SEQ ID NO:327, SEQ ID NO:329, SEQ ID NO:331, SEQ ID NO:333, SEQ ID NO:335, SEQ ID NO:337, SEQ ID NO:339, SEQ ID NO:341, SEQ ID NO:343, SEQ ID NO:345, SEQ ID NO:347, SEQ ID NO:349, SEQ ID NO:351, SEQ ID NO:353, SEQ ID NO:355, SEQ ID NO:357, SEQ ID NO:359, SEQ ID NO:361, SEQ ID NO:363, SEQ ID NO:365, SEQ ID NO:367, SEQ ID NO:369, SEQ ID NO:371, SEQ ID NO:373, SEQ ID NO:375, SEQ ID NO:377, SEQ ID NO:379, SEQ ID NO:381, SEQ ID NO:383, SEQ ID NO:385, SEQ ID NO:387, SEQ ID NO:389, SEQ ID NO:391, SEQ ID NO:393, SEQ ID NO:395, SEQ ID NO:397, SEQ ID NO:399, SEQ ID NO:401, SEQ ID NO:403, SEQ ID NO:405, SEQ ID NO:407, SEQ ID NO:409, SEQ ID NO:411, SEQ ID NO:413, SEQ ID NO:415, SEQ ID NO:417, SEQ ID NO:419, SEQ ID NO:421, SEQ ID

NO:423, SEQ ID NO:425, SEQ ID NO:427, SEQ ID NO:429, SEQ ID NO:431, SEQ ID
 NO:433, SEQ ID NO:435, SEQ ID NO:437, SEQ ID NO:439, SEQ ID NO:441, SEQ ID
 NO:443, SEQ ID NO:445, SEQ ID NO:447, SEQ ID NO:449, SEQ ID NO:451, SEQ ID
 NO:453, SEQ ID NO:455, SEQ ID NO:457, SEQ ID NO:459, SEQ ID NO:461, SEQ ID
 NO:461, SEQ ID NO:463, SEQ ID NO:464, SEQ ID NO:466, SEQ ID NO:468, SEQ ID
 NO:469, SEQ ID NO:470, SEQ ID NO:471, SEQ ID NO:472, SEQ ID NO:474, SEQ ID
 NO:476, SEQ ID NO:477, SEQ ID NO:479, SEQ ID NO:481, SEQ ID NO:482, SEQ ID
 NO:483, SEQ ID NO:485, SEQ ID NO:487, SEQ ID NO:488, SEQ ID NO:489, SEQ ID
 NO:490, SEQ ID NO:491, SEQ ID NO:493, SEQ ID NO:495, SEQ ID NO:496, SEQ ID
 NO:497, SEQ ID NO:499, SEQ ID NO:501, SEQ ID NO:502, SEQ ID NO:503, SEQ ID
 NO:504, SEQ ID NO:505, SEQ ID NO:506, SEQ ID NO:507, SEQ ID NO:508, SEQ ID
 NO:510, SEQ ID NO:512, SEQ ID NO:513, SEQ ID NO:514, SEQ ID NO:516, SEQ ID
 NO:518, SEQ ID NO:518, SEQ ID NO:520, SEQ ID NO:521, SEQ ID NO:523, SEQ ID
 NO:525, SEQ ID NO:526, SEQ ID NO:528, SEQ ID NO:530, SEQ ID NO:531, SEQ ID
 NO:533, SEQ ID NO:535, SEQ ID NO:536, SEQ ID NO:537, SEQ ID NO:538, SEQ ID
 NO:540, SEQ ID NO:542, SEQ ID NO:543, SEQ ID NO:545, SEQ ID NO:547, SEQ ID
 NO:548, SEQ ID NO:549, SEQ ID NO:550, SEQ ID NO:551, SEQ ID NO:553, SEQ ID
 NO:555, SEQ ID NO:556, SEQ ID NO:557, SEQ ID NO:559, SEQ ID NO:561, SEQ ID
 NO:562, SEQ ID NO:563, SEQ ID NO:564, SEQ ID NO:566, SEQ ID NO:568, SEQ ID
 NO:570, SEQ ID NO:572, SEQ ID NO:574, SEQ ID NO:576, SEQ ID NO:578, SEQ ID
 NO:580, SEQ ID NO:582, SEQ ID NO:584, SEQ ID NO:586, SEQ ID NO:588, SEQ ID
 NO:589, SEQ ID NO:590, SEQ ID NO:591, SEQ ID NO:592, SEQ ID NO:594, SEQ ID
 NO:604, SEQ ID NO:606, SEQ ID NO:608, SEQ ID NO:610, SEQ ID NO:612, SEQ ID
 NO:614, SEQ ID NO:616, SEQ ID NO:618, SEQ ID NO:620 or SEQ ID NO:622, and
 25 subsequences thereof and variants thereof, e.g., at least about 10, 15, 20, 25, 30, 35, 40,
 45, 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800,
 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500 or
 more residues in length, or over the full length of an enzyme. Exemplary polypeptide or
 peptide sequences of the invention include sequence encoded by a nucleic acid of the
 30 invention. Exemplary polypeptide or peptide sequences of the invention include
 polypeptides or peptides specifically bound by an antibody of the invention. In one
 aspect, a polypeptide of the invention has at least one amylase activity, e.g., an alpha
 amylase activity.

Another aspect of the invention is an isolated or recombinant polypeptide or peptide including at least 10 consecutive bases of a polypeptide or peptide sequence of the invention, sequences substantially identical thereto, and the sequences complementary thereto.

In one aspect, the amylase activity of a polypeptide or peptide of the invention comprises an α -amylase activity, including the ability to hydrolyze internal α -1,4-glucosidic linkages in starch to produce smaller molecular weight malto-dextrins. In one aspect, the α -amylase activity includes hydrolyzing internal α -1,4-glucosidic linkages in starch at random. The amylase activity can comprise a glucoamylase activity, a 1,4- α -D-glucan glucohydrolase activity, an α -amylase activity, an exoamylase activity, or a β -amylase activity. The amylase activity can comprise hydrolyzing glucosidic bonds. In one aspect, the glucosidic bonds comprise an α -1,4-glucosidic bond. In another aspect, the glucosidic bonds comprise an α -1,6-glucosidic bond. In one aspect, the amylase activity comprises hydrolyzing glucosidic bonds in starch, e.g., liquefied starch. The amylase activity can further comprise hydrolyzing glucosidic bonds into maltodextrins. In one aspect, the amylase activity comprises cleaving a maltose or a D-glucose unit from non-reducing end of the starch.

In one aspect, the amylase activity of the invention comprises a glucoamylase activity, which can comprise catalysis of the hydrolysis of glucosidic bonds. The glucoamylase activity of the invention can comprise catalyzing the step-wise hydrolytic release of D-glucose from the non-reducing ends of starch or other related dextrins. The glucoamylase activity can comprise a 1,4- α -D-glucan glucohydrolase activity. The glucoamylase activity can comprise catalysis of the hydrolysis of malto-dextrins resulting in the generation of free glucose. The glucoamylase activity can comprise an exoamylase activity. The glucoamylase activity can comprise an α -amylase or a β -amylase activity. The hydrolyzed glucosidic bonds can comprise α -1,4-glucosidic bonds or α -1,6-glucosidic bonds. The glucoamylase activity can comprise hydrolyzing glucosidic bonds in a starch. The glucoamylase activity can further comprise hydrolyzing glucosidic bonds in the starch to produce maltodextrins. The glucoamylase activity can comprise cleaving a maltose or a D-glucose unit from non-reducing end of the starch.

In one aspect, the amylase activity can be thermostable. The polypeptide can retain an amylase activity under conditions comprising a temperature range of between about 37°C to about 95°C, between about 55°C to about 85°C, between about

70°C to about 95°C, or between about 90°C to about 95°C. In another aspect, the amylase activity can be thermotolerant. The polypeptide can retain an amylase activity after exposure to a temperature in the range from greater than 37°C to about 95°C, or in the range from greater than 55°C to about 85°C. In one aspect, the polypeptide can retain an amylase activity after exposure to a temperature in the range from greater than 90°C to about 95°C at pH 4.5.

In one aspect, the isolated or recombinant polypeptide can comprise the polypeptide of the invention that lacks a signal sequence. In one aspect, the isolated or recombinant polypeptide can comprise the polypeptide of the invention comprising a heterologous signal sequence, such as a heterologous amylase or non-amylase signal sequence.

In one aspect, the invention provides a signal sequence comprising a peptide as set forth in Table 3. In one aspect, the invention provides a signal sequence consisting of a peptide as set forth in Table 3. In one aspect, the invention provides chimeric proteins comprising a first domain comprising a signal sequence of the invention and at least a second domain. The protein can be a fusion protein. The second domain can comprise an enzyme. The enzyme can be an amylase (e.g., an amylase of the invention, or, another amylase).

In one aspect, the amylase activity comprises a specific activity at about 37°C in the range from about 10 to 10,000, or, 100 to about 1000 units per milligram of protein. In another aspect, the amylase activity comprises a specific activity from about 500 to about 750 units per milligram of protein. Alternatively, the amylase activity comprises a specific activity at 37°C in the range from about 500 to about 1200 units per milligram of protein. In one aspect, the amylase activity comprises a specific activity at 37°C in the range from about 750 to about 1000 units per milligram of protein. In another aspect, the thermotolerance comprises retention of at least half of the specific activity of the amylase at 37°C after being heated to the elevated temperature. Alternatively, the thermotolerance can comprise retention of specific activity at 37°C in the range from about 500 to about 1200 units per milligram of protein after being heated to the elevated temperature.

The invention provides isolated or recombinant polypeptides of the invention, wherein the polypeptide comprises at least one glycosylation site. In one aspect, glycosylation can be an N-linked glycosylation. In one aspect, the polypeptide

can be glycosylated after being expressed in a *P. pastoris* or a *S. pombe*. The invention also provides methods for adding glycosylation to a polypeptide, either post-translationally or chemically, to change the property of the polypeptides, e.g., its thermal stability, solubility, tendency to aggregate, and the like.

In one aspect, the polypeptide can retain an amylase activity under conditions comprising about pH 6.5, pH 6, pH 5.5, pH 5, pH 4.5 or pH 4. In another aspect, the polypeptide can retain an amylase activity under conditions comprising about pH 7, pH 7.5 pH 8.0, pH 8.5, pH 9, pH 9.5, pH 10, pH 10.5 or pH 11.

The invention provides protein preparations comprising a polypeptide of the invention, wherein the protein preparation comprises a liquid, a solid or a gel.

The invention provides heterodimers comprising a polypeptide of the invention and a second domain. In one aspect, the second domain can be a polypeptide and the heterodimer can be a fusion protein. In one aspect, the second domain can be an epitope or a tag. In one aspect, the invention provides homodimers comprising a polypeptide of the invention.

The invention provides immobilized polypeptides having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention, a polypeptide encoded by a nucleic acid of the invention, or a polypeptide comprising a polypeptide of the invention and a second domain. In one aspect, the polypeptide can be immobilized on a cell, a metal, a resin, a polymer, a ceramic, a glass, a microelectrode, a graphitic particle, a bead, a gel, a plate, an array or a capillary tube.

The invention provides arrays comprising an immobilized nucleic acid of the invention. The invention provides arrays comprising an antibody of the invention.

25 The invention provides isolated or recombinant antibodies that specifically bind to a polypeptide of the invention or to a polypeptide encoded by a nucleic acid of the invention. The antibody can be a monoclonal or a polyclonal antibody. The invention provides hybridomas comprising an antibody of the invention, e.g., an antibody that specifically binds to a polypeptide of the invention or to a polypeptide encoded by a nucleic acid of the invention.

30 The invention provides food supplements for an animal comprising a polypeptide of the invention, e.g., a polypeptide encoded by the nucleic acid of the invention. In one aspect, the polypeptide in the food supplement can be glycosylated. The invention provides edible enzyme delivery matrices comprising a polypeptide of the

invention, e.g., a polypeptide encoded by the nucleic acid of the invention. In one aspect, the delivery matrix comprises a pellet. In one aspect, the polypeptide can be glycosylated. In one aspect, the amylase activity is thermotolerant. In another aspect, the amylase activity is thermostable.

The invention provides method of isolating or identifying a polypeptide having an amylase activity comprising the steps of: (a) providing an antibody of the invention; (b) providing a sample comprising polypeptides; and (c) contacting the sample of step (b) with the antibody of step (a) under conditions wherein the antibody can specifically bind to the polypeptide, thereby isolating or identifying a polypeptide having an amylase activity.

The invention provides methods of making an anti-amylase antibody comprising administering to a non-human animal a nucleic acid of the invention or a polypeptide of the invention or subsequences thereof in an amount sufficient to generate a humoral immune response, thereby making an anti-amylase antibody. The invention provides methods of making an anti-amylase immune comprising administering to a non-human animal a nucleic acid of the invention or a polypeptide of the invention or subsequences thereof in an amount sufficient to generate an immune response.

The invention provides methods of producing a recombinant polypeptide comprising the steps of: (a) providing a nucleic acid of the invention operably linked to a promoter; and (b) expressing the nucleic acid of step (a) under conditions that allow expression of the polypeptide, thereby producing a recombinant polypeptide. In one aspect, the method can further comprise transforming a host cell with the nucleic acid of step (a) followed by expressing the nucleic acid of step (a), thereby producing a recombinant polypeptide in a transformed cell.

25 The invention provides methods for identifying a polypeptide having an amylase activity comprising the following steps: (a) providing a polypeptide of the invention; or a polypeptide encoded by a nucleic acid of the invention; (b) providing an amylase substrate; and (c) contacting the polypeptide or a fragment or variant thereof of step (a) with the substrate of step (b) and detecting a decrease in the amount of substrate
30 or an increase in the amount of a reaction product, wherein a decrease in the amount of the substrate or an increase in the amount of the reaction product detects a polypeptide having an amylase activity. In one aspect, the substrate can be a starch, e.g., a liquefied starch.

The invention provides methods for identifying an amylase substrate comprising the following steps: (a) providing a polypeptide of the invention; or a polypeptide encoded by a nucleic acid of the invention; (b) providing a test substrate; and (c) contacting the polypeptide of step (a) with the test substrate of step (b) and detecting a decrease in the amount of substrate or an increase in the amount of reaction product, wherein a decrease in the amount of the substrate or an increase in the amount of a reaction product identifies the test substrate as an amylase substrate.

The invention provides methods of determining whether a test compound specifically binds to a polypeptide comprising the following steps: (a) expressing a nucleic acid or a vector comprising the nucleic acid under conditions permissive for translation of the nucleic acid to a polypeptide, wherein the nucleic acid comprises a nucleic acid of the invention, or, providing a polypeptide of the invention; (b) providing a test compound; (c) contacting the polypeptide with the test compound; and (d) determining whether the test compound of step (b) specifically binds to the polypeptide.

The invention provides methods for identifying a modulator of an amylase activity comprising the following steps: (a) providing a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention; (b) providing a test compound; (c) contacting the polypeptide of step (a) with the test compound of step (b) and measuring an activity of the amylase, wherein a change in the amylase activity measured in the presence of the test compound compared to the activity in the absence of the test compound provides a determination that the test compound modulates the amylase activity. In one aspect, the amylase activity can be measured by providing an amylase substrate and detecting a decrease in the amount of the substrate or an increase in the amount of a reaction product, or, an increase in the amount of the substrate or a decrease in the amount of a reaction product. A decrease in the amount of the substrate or an increase in the amount of the reaction product with the test compound as compared to the amount of substrate or reaction product without the test compound identifies the test compound as an activator of amylase activity. An increase in the amount of the substrate or a decrease in the amount of the reaction product with the test compound as compared to the amount of substrate or reaction product without the test compound identifies the test compound as an inhibitor of amylase activity.

The invention provides computer systems comprising a processor and a data storage device wherein said data storage device has stored thereon a polypeptide

sequence or a nucleic acid sequence of the invention (e.g., a polypeptide encoded by a nucleic acid of the invention). In one aspect, the computer system can further comprise a sequence comparison algorithm and a data storage device having at least one reference sequence stored thereon. In another aspect, the sequence comparison algorithm
5 comprises a computer program that indicates polymorphisms. In one aspect, the computer system can further comprise an identifier that identifies one or more features in said sequence. The invention provides computer readable media having stored thereon a polypeptide sequence or a nucleic acid sequence of the invention. The invention provides methods for identifying a feature in a sequence comprising the steps of: (a) reading the
10 sequence using a computer program which identifies one or more features in a sequence, wherein the sequence comprises a polypeptide sequence or a nucleic acid sequence of the invention; and (b) identifying one or more features in the sequence with the computer program. The invention provides methods for comparing a first sequence to a second sequence comprising the steps of: (a) reading the first sequence and the second sequence
15 through use of a computer program which compares sequences, wherein the first sequence comprises a polypeptide sequence or a nucleic acid sequence of the invention; and (b) determining differences between the first sequence and the second sequence with the computer program. The step of determining differences between the first sequence and the second sequence can further comprise the step of identifying polymorphisms. In
20 one aspect, the method can further comprise an identifier that identifies one or more features in a sequence. In another aspect, the method can comprise reading the first sequence using a computer program and identifying one or more features in the sequence.

The invention provides methods for isolating or recovering a nucleic acid encoding a polypeptide having an amylase activity from an environmental sample
25 comprising the steps of: (a) providing an amplification primer sequence pair for amplifying a nucleic acid encoding a polypeptide having an amylase activity, wherein the primer pair is capable of amplifying a nucleic acid of the invention; (b) isolating a nucleic acid from the environmental sample or treating the environmental sample such that nucleic acid in the sample is accessible for hybridization to the amplification primer pair;
30 and, (c) combining the nucleic acid of step (b) with the amplification primer pair of step (a) and amplifying nucleic acid from the environmental sample, thereby isolating or recovering a nucleic acid encoding a polypeptide having an amylase activity from an environmental sample. One or each member of the amplification primer sequence pair

can comprise an oligonucleotide comprising at least about 10 to 50 consecutive bases of a sequence of the invention.

The invention provides methods for isolating or recovering a nucleic acid encoding a polypeptide having an amylase activity from an environmental sample comprising the steps of: (a) providing a polynucleotide probe comprising a nucleic acid of the invention or a subsequence thereof; (b) isolating a nucleic acid from the environmental sample or treating the environmental sample such that nucleic acid in the sample is accessible for hybridization to a polynucleotide probe of step (a); (c) combining the isolated nucleic acid or the treated environmental sample of step (b) with the polynucleotide probe of step (a); and (d) isolating a nucleic acid that specifically hybridizes with the polynucleotide probe of step (a), thereby isolating or recovering a nucleic acid encoding a polypeptide having an amylase activity from an environmental sample. The environmental sample can comprise a water sample, a liquid sample, a soil sample, an air sample or a biological sample. In one aspect, the biological sample can be derived from a bacterial cell, a protozoan cell, an insect cell, a yeast cell, a plant cell, a fungal cell or a mammalian cell.

The invention provides methods of generating a variant of a nucleic acid encoding a polypeptide having an amylase activity comprising the steps of: (a) providing a template nucleic acid comprising a nucleic acid of the invention; and (b) modifying, deleting or adding one or more nucleotides in the template sequence, or a combination thereof, to generate a variant of the template nucleic acid. In one aspect, the method can further comprise expressing the variant nucleic acid to generate a variant amylase polypeptide. The modifications, additions or deletions can be introduced by a method comprising error-prone PCR, shuffling, oligonucleotide-directed mutagenesis, assembly PCR, sexual PCR mutagenesis, *in vivo* mutagenesis, cassette mutagenesis, recursive ensemble mutagenesis, exponential ensemble mutagenesis, site-specific mutagenesis, gene reassembly, gene site saturated mutagenesis (GSSM), synthetic ligation reassembly (SLR) or a combination thereof. In another aspect, the modifications, additions or deletions are introduced by a method comprising recombination, recursive sequence recombination, phosphothioate-modified DNA mutagenesis, uracil-containing template mutagenesis, gapped duplex mutagenesis, point mismatch repair mutagenesis, repair-deficient host strain mutagenesis, chemical mutagenesis, radiogenic mutagenesis, deletion mutagenesis, restriction-selection mutagenesis, restriction-purification mutagenesis,

artificial gene synthesis, ensemble mutagenesis, chimeric nucleic acid multimer creation and a combination thereof.

In one aspect, the method can be iteratively repeated until an amylase having an altered or different activity or an altered or different stability from that of a polypeptide encoded by the template nucleic acid is produced. In one aspect, the variant amylase polypeptide is thermotolerant, and retains some activity after being exposed to an elevated temperature. In another aspect, the variant amylase polypeptide has increased glycosylation as compared to the amylase encoded by a template nucleic acid.

Alternatively, the variant amylase polypeptide has an amylase activity under a high temperature, wherein the amylase encoded by the template nucleic acid is not active under the high temperature. In one aspect, the method can be iteratively repeated until an amylase coding sequence having an altered codon usage from that of the template nucleic acid is produced. In another aspect, the method can be iteratively repeated until an amylase gene having higher or lower level of message expression or stability from that of the template nucleic acid is produced.

The invention provides methods for modifying codons in a nucleic acid encoding a polypeptide having an amylase activity to increase its expression in a host cell, the method comprising the following steps: (a) providing a nucleic acid of the invention encoding a polypeptide having an amylase activity; and, (b) identifying a non-preferred or a less preferred codon in the nucleic acid of step (a) and replacing it with a preferred or neutrally used codon encoding the same amino acid as the replaced codon, wherein a preferred codon is a codon over-represented in coding sequences in genes in the host cell and a non-preferred or less preferred codon is a codon under-represented in coding sequences in genes in the host cell, thereby modifying the nucleic acid to increase its expression in a host cell.

The invention provides methods for modifying codons in a nucleic acid encoding a polypeptide having an amylase activity; the method comprising the following steps: (a) providing a nucleic acid of the invention; and, (b) identifying a codon in the nucleic acid of step (a) and replacing it with a different codon encoding the same amino acid as the replaced codon, thereby modifying codons in a nucleic acid encoding an amylase.

The invention provides methods for modifying codons in a nucleic acid encoding a polypeptide having an amylase activity to increase its expression in a host

cell, the method comprising the following steps: (a) providing a nucleic acid of the invention encoding an amylase polypeptide; and, (b) identifying a non-preferred or a less preferred codon in the nucleic acid of step (a) and replacing it with a preferred or neutrally used codon encoding the same amino acid as the replaced codon, wherein a preferred codon is a codon over-represented in coding sequences in genes in the host cell and a non-preferred or less preferred codon is a codon under-represented in coding sequences in genes in the host cell, thereby modifying the nucleic acid to increase its expression in a host cell.

The invention provides methods for modifying a codon in a nucleic acid encoding a polypeptide having an amylase activity to decrease its expression in a host cell, the method comprising the following steps: (a) providing a nucleic acid of the invention; and (b) identifying at least one preferred codon in the nucleic acid of step (a) and replacing it with a non-preferred or less preferred codon encoding the same amino acid as the replaced codon, wherein a preferred codon is a codon over-represented in coding sequences in genes in a host cell and a non-preferred or less preferred codon is a codon under-represented in coding sequences in genes in the host cell, thereby modifying the nucleic acid to decrease its expression in a host cell. In one aspect, the host cell can be a bacterial cell, a fungal cell, an insect cell, a yeast cell, a plant cell or a mammalian cell.

The invention provides methods for producing a library of nucleic acids encoding a plurality of modified amylase active sites or substrate binding sites, wherein the modified active sites or substrate binding sites are derived from a first nucleic acid comprising a sequence encoding a first active site or a first substrate binding site the method comprising the following steps: (a) providing a first nucleic acid encoding a first active site or first substrate binding site, wherein the first nucleic acid sequence comprises a sequence that hybridizes under stringent conditions to a nucleic acid of the invention, and the nucleic acid encodes an amylase active site or an amylase substrate binding site; (b) providing a set of mutagenic oligonucleotides that encode naturally-occurring amino acid variants at a plurality of targeted codons in the first nucleic acid; and, (c) using the set of mutagenic oligonucleotides to generate a set of active site-encoding or substrate binding site-encoding variant nucleic acids encoding a range of amino acid variations at each amino acid codon that was mutagenized, thereby producing a library of nucleic acids encoding a plurality of modified amylase active sites or substrate binding sites. In one

aspect, the method comprises mutagenizing the first nucleic acid of step (a) by a method comprising an optimized directed evolution system, gene site-saturation mutagenesis (GSSM), synthetic ligation reassembly (SLR), error-prone PCR, shuffling, oligonucleotide-directed mutagenesis, assembly PCR, sexual PCR mutagenesis, in vivo mutagenesis, cassette mutagenesis, recursive ensemble mutagenesis, exponential ensemble mutagenesis, site-specific mutagenesis, gene reassembly, gene site saturated mutagenesis (GSSM), synthetic ligation reassembly (SLR) and a combination thereof. In another aspect, the method comprises mutagenizing the first nucleic acid of step (a) or variants by a method comprising recombination, recursive sequence recombination, phosphothioate-modified DNA mutagenesis, uracil-containing template mutagenesis, gapped duplex mutagenesis, point mismatch repair mutagenesis, repair-deficient host strain mutagenesis, chemical mutagenesis, radiogenic mutagenesis, deletion mutagenesis, restriction-selection mutagenesis, restriction-purification mutagenesis, artificial gene synthesis, ensemble mutagenesis, chimeric nucleic acid multimer creation and a combination thereof.

The invention provides methods for making a small molecule comprising the following steps: (a) providing a plurality of biosynthetic enzymes capable of synthesizing or modifying a small molecule, wherein one of the enzymes comprises an amylase enzyme encoded by a nucleic acid of the invention; (b) providing a substrate for at least one of the enzymes of step (a); and (c) reacting the substrate of step (b) with the enzymes under conditions that facilitate a plurality of biocatalytic reactions to generate a small molecule by a series of biocatalytic reactions. The invention provides methods for modifying a small molecule comprising the following steps: (a) providing an amylase enzyme, wherein the enzyme comprises a polypeptide of the invention, or, a polypeptide
25 encoded by a nucleic acid of the invention, or a subsequence thereof; (b) providing a small molecule; and (c) reacting the enzyme of step (a) with the small molecule of step (b) under conditions that facilitate an enzymatic reaction catalyzed by the amylase enzyme, thereby modifying a small molecule by an amylase enzymatic reaction. In one aspect, the method can comprise a plurality of small molecule substrates for the enzyme
30 of step (a), thereby generating a library of modified small molecules produced by at least one enzymatic reaction catalyzed by the amylase enzyme. In one aspect, the method can comprise a plurality of additional enzymes under conditions that facilitate a plurality of biocatalytic reactions by the enzymes to form a library of modified small molecules

produced by the plurality of enzymatic reactions. In another aspect, the method can further comprise the step of testing the library to determine if a particular modified small molecule which exhibits a desired activity is present within the library. The step of testing the library can further comprise the steps of systematically eliminating all but one of the biocatalytic reactions used to produce a portion of the plurality of the modified small molecules within the library by testing the portion of the modified small molecule for the presence or absence of the particular modified small molecule with a desired activity, and identifying at least one specific biocatalytic reaction that produces the particular modified small molecule of desired activity.

The invention provides methods for determining a functional fragment of an amylase enzyme comprising the steps of: (a) providing an amylase enzyme, wherein the enzyme comprises a polypeptide of the invention, or a polypeptide encoded by a nucleic acid of the invention, or a subsequence thereof; and (b) deleting a plurality of amino acid residues from the sequence of step (a) and testing the remaining subsequence for an amylase activity, thereby determining a functional fragment of an amylase enzyme. In one aspect, the amylase activity is measured by providing an amylase substrate and detecting a decrease in the amount of the substrate or an increase in the amount of a reaction product.

The invention provides methods for whole cell engineering of new or modified phenotypes by using real-time metabolic flux analysis, the method comprising the following steps: (a) making a modified cell by modifying the genetic composition of a cell, wherein the genetic composition is modified by addition to the cell of a nucleic acid of the invention; (b) culturing the modified cell to generate a plurality of modified cells; (c) measuring at least one metabolic parameter of the cell by monitoring the cell culture of step (b) in real time; and, (d) analyzing the data of step (c) to determine if the measured parameter differs from a comparable measurement in an unmodified cell under similar conditions, thereby identifying an engineered phenotype in the cell using real-time metabolic flux analysis. In one aspect, the genetic composition of the cell can be modified by a method comprising deletion of a sequence or modification of a sequence in the cell, or, knocking out the expression of a gene. In one aspect, the method can further comprise selecting a cell comprising a newly engineered phenotype. In another aspect, the method can comprise culturing the selected cell, thereby generating a new cell strain comprising a newly engineered phenotype.

The invention provides methods for hydrolyzing a starch comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a composition comprising a starch; and (c) contacting the polypeptide of step (a) with the composition of step (b) under conditions wherein the polypeptide hydrolyzes the starch. In one aspect, the composition comprising starch that comprises an α -1,4-glucosidic bond or an α -1,6-glucosidic bond. In one aspect, the amylase activity is an α -amylase activity. In one aspect, the α -amylase activity hydrolyzes internal bonds in a starch or other polysaccharide.

The invention provides methods for liquefying or removing a starch from a composition comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a composition comprising a starch; and (c) contacting the polypeptide of step (a) with the composition of step (b) under conditions wherein the polypeptide removes or liquefies the starch.

The invention provides methods of increasing thermotolerance or thermostability of an amylase polypeptide, the method comprising glycosylating an amylase polypeptide, wherein the polypeptide comprises at least thirty contiguous amino acids of a polypeptide of the invention; or a polypeptide encoded by a nucleic acid sequence of the invention, thereby increasing the thermotolerance or thermostability of the amylase polypeptide. In one aspect, the amylase specific activity can be thermostable or thermotolerant at a temperature in the range from greater than about 37°C to about 95°C.

25 The invention provides methods for overexpressing a recombinant amylase polypeptide in a cell comprising expressing a vector comprising a nucleic acid comprising a nucleic acid of the invention or a nucleic acid sequence of the invention, wherein the sequence identities are determined by analysis with a sequence comparison algorithm or by visual inspection, wherein overexpression is effected by use of a high activity promoter, a dicistronic vector or by gene amplification of the vector.

30 The invention provides detergent compositions comprising a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention, wherein the polypeptide comprises an amylase activity. In one aspect, the amylase can be a

nonsurface-active amylase. In another aspect, the amylase can be a surface-active amylase.

The invention provides methods for washing an object comprising the following steps: (a) providing a composition comprising a polypeptide having an amylase activity, wherein the polypeptide comprises: a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention; (b) providing an object; and (c) contacting the polypeptide of step (a) and the object of step (b) under conditions wherein the composition can wash the object.

The invention provides methods for hydrolyzing starch, e.g., in a feed or a food prior to consumption by an animal, comprising the following steps: (a) obtaining a composition, e.g., a feed material, comprising a starch, wherein the polypeptide comprises: a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention; and (b) adding the polypeptide of step (a) to the composition, e.g., the feed or food material, in an amount sufficient for a sufficient time period to cause hydrolysis of the starch, thereby hydrolyzing the starch. In one aspect, the food or feed comprises rice, corn, barley, wheat, legumes, or potato.

The invention provides methods for textile desizing comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention; (b) providing a fabric; and (c) contacting the polypeptide of step (a) and the fabric of step (b) under conditions wherein the amylase can desize the fabric.

The invention provides methods for deinking of paper or fibers comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a composition comprising paper or fiber; and (c) contacting the polypeptide of step (a) and the composition of step (b) under conditions wherein the polypeptide can deink the paper or fiber.

The invention provides methods for treatment of lignocellulosic fibers comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a lignocellulosic fiber; and (c) contacting the polypeptide of step (a) and the fiber of step

(b) under conditions wherein the polypeptide can treat the fiber thereby improving the fiber properties.

The invention provides methods for producing a high-maltose or a high-glucose syrup comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises an enzyme of the invention; (b) providing a composition comprising a starch; and (c) contacting the polypeptide of step (a) and the fabric of step (b) under conditions wherein the polypeptide of step (a) can liquefy the composition of step (b) thereby producing a soluble starch hydrolysate and saccharify the soluble starch hydrolysate thereby producing the syrup. In one aspect, the starch can be from rice, corn, barley, wheat, legumes, potato, or sweet potato.

The invention provides methods for improving the flow of the starch-containing production fluids comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing production fluid; and (c) contacting the polypeptide of step (a) and the production fluid of step (b) under conditions wherein the amylase can hydrolyze the starch in the production fluid thereby improving its flow by decreasing its density. In one aspect, the production fluid can be from a subterranean formation.

The invention provides anti-staling compositions comprising a polypeptide of the invention or a polypeptide encoded by a nucleic acid of the invention. The invention provides methods for preventing staling of the baked products comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a composition containing starch used for baking; (c) combining the polypeptide of step (a) with the composition of the step (b) under conditions wherein the polypeptide can hydrolyze the starch in the composition used for baking thereby preventing staling of the baked product. In one aspect, the baked product can be bread.

The invention provides methods for using amylase in brewing or alcohol production comprising the following steps: (a) providing a polypeptide having an amylase activity, wherein the polypeptide comprises a polypeptide of the invention; (b) providing a composition containing starch and used for brewing or in alcohol production; (c) combining the polypeptide of step (a) with the composition of the step (b) under conditions wherein the polypeptide can hydrolyze the starch in the composition used for

brewing or in alcohol production. In one aspect, the composition containing starch can be beer.

The invention provides methods of making a transgenic plant comprising the following steps: (a) introducing a heterologous nucleic acid sequence into the cell, wherein the heterologous nucleic sequence comprises a nucleic acid sequence of the invention, thereby producing a transformed plant cell; and (b) producing a transgenic plant from the transformed cell. In one aspect, the step (a) can further comprise introducing the heterologous nucleic acid sequence by electroporation or microinjection of plant cell protoplasts. In another aspect, the step (a) can further comprise introducing the heterologous nucleic acid sequence directly to plant tissue by DNA particle bombardment. Alternatively, the step (a) can further comprise introducing the heterologous nucleic acid sequence into the plant cell DNA using an *Agrobacterium tumefaciens* host. In one aspect, the plant cell can be a potato, corn, rice, wheat, tobacco, or barley cell.

The invention provides methods of expressing a heterologous nucleic acid sequence in a plant cell comprising the following steps: (a) transforming the plant cell with a heterologous nucleic acid sequence operably linked to a promoter, wherein the heterologous nucleic sequence comprises a nucleic acid of the invention; (b) growing the plant under conditions wherein the heterologous nucleic acids sequence is expressed in the plant cell.

The invention also provides a process for preparing a dough or a baked product prepared from the dough which comprises adding an amylase of the invention to the dough in an amount which is effective to retard the staling of the bread. The invention also provides a dough comprising said amylase and a premix comprising flour together with said amylase. Finally, the invention provides an enzymatic baking additive, which contains said amylase. The use of the amylase in accordance with the present invention provides an improved anti-staling effect as measured by, e.g. less crumb firming, retained crumb elasticity, improved slice-ability (e.g. fewer crumbs, non-gummy crumb), improved palatability or flavor.

The invention provides delayed release ("controlled release") compositions comprising an desired ingredient coated by a latex polymer (or equivalent) coating. In one aspect, the desired ingredient comprises an enzyme, e.g., an enzyme of the invention. In one aspect, the desired ingredient comprises a small molecule, a drug, a

polysaccharide, a lipid, a nucleic acid, a vitamin, an antibiotic or an insecticide. In one aspect, the desired ingredient comprises a pellet or a matrix, e. g. , a pellet or a matrix comprising an edible material (e. g. , as an animal food or feed or supplement or medicament). The invention also provides methods for the "controlled release" or "delayed release" of a composition, wherein
5 the composition is coated by a latex polymer (or equivalent) coating.

In one aspect, the latex polymer coating comprises a latex paint, or equivalent. The latex polymer coating can comprise a (meth) acrylate, a vinyl acetate, a styrene, an ethylene, a vinyl chloride, a butadiene, a vinylidene chloride, a vinyl versatate, a vinyl propionate, a t-butyl acrylate, an acrylonitrile, a neoprene, a maleate, a fumarate, equivalents thereof, combinations
10 thereof and/or derivatives thereof.

The details of one or more embodiments of the invention are set forth in the accompanying drawings and the description below. Other features, aspects, and advantages of the invention will be apparent from the description and drawings, and from the claims.

All publications, patents, patent applications, GenBank sequences and ATCC deposits,
15 cited herein are hereby expressly incorporated by reference for all purposes.

As used herein, except where the context requires otherwise, the term "comprise" and variations of the term, such as "comprising", "comprises" and "comprised", are not intended to exclude other additives, components, integers or steps.

Reference to any prior art in the specification is not, and should not be taken as, an
20 acknowledgment, or any form of suggestion, that this prior art forms part of the common general knowledge in Australia or any other jurisdiction or that this prior art could reasonably be expected to be ascertained, understood and regarded as relevant by a person skilled in the art.

DESCRIPTION OF DRAWINGS

Figure 1 is a block diagram of a computer system.

Figure 2 is a flow diagram illustrating one aspect of a process for comparing a new nucleotide or protein sequence with a database of sequences in order to determine the homology levels between the new sequence and the sequences in the database.

Figure 3 is a flow diagram illustrating one aspect of a process in a computer for
5 determining whether two sequences are homologous.

Figure 4 is a flow diagram illustrating one aspect of an identifier process 300 for detecting the presence of a feature in a sequence.

Figure 5 is a graph showing the Residual activity of various amylases following heating to 90°C for 10 min in Example 1.

Figure 6 is a graph showing the net percent starch removed versus enzyme concentration in ADW wash test with bleach and chelators.

Figure 7 is a graph showing the activity of parental amylases at pH 8, 40°C in ADW formulation at 55°C.

Figure 8 is a graph of data regarding the H₂O₂ tolerance of the novel enzymes in Example 4.

Figure 9 is a graph of the pH and temperature data for a selection of the amylases characterized. Figure 9a shows the data at pH 8 and 40°C and Figure 9b shows the data at pH 10 and 50°C.

Figure 10 sets forth the sequences to be used in reassembly experiments with the enzymes.

Figure 11 illustrates a sample Standard Curve of the assay of Example 5.

Figure 12 illustrates the pH rate profiles for SEQ ID NO.: 127, which has a neutral optimum pH and SEQ ID NO.: 211, which has an optimum around pH 10.

Figure 13 shows the stability of exemplary amylases vs. a commercial enzyme, as discussed in Example 2.

Figure 14 shows the sequence alignments of hypothermophilic α -amylases, as set forth in Example 8. Figure 14a shows an alignment of amylase sequences. SEQ ID NO.: 81= an environmental clone; pyro = *Pyrococcus sp.*

(strain:KOD1), Tachibana (1996) J. Ferment. Bioeng. 82:224-232; pyro2 = *Pyrococcus furiosus*, Appl. Environ. Microbiol. 63 (9):3569-3576, 1997; Thermo = *Thermococcus sp.*; Thermo2 = *Thermococcus hydrothermalis*, Leveque, E. et al. Patent: France 98.05655 05-MAY-1998. Figure 14b shows the amino acid sequence alignment of identified sequences: SEQ ID NO.: 81; pyro; SEQ ID NO.:75; SEQ ID NO.: 77; SEQ ID NO.: 83; SEQ ID NO.: 85; thermo2; SEQ ID NO.: 79 ; thermo ; pyro2 ; clone A; thermo3. Figure 14c shows the nucleic acid sequence alignment corresponding to the polypeptide sequence of Figures 5 and 6. SEQ ID NO.: 81; SEQ ID NO.:75; SEQ ID NO.: 77; SEQ ID NO.: 83; SEQ ID NO.: 85; SEQ ID NO.: 79; clone A; and SEQ ID NO.: 73.

Figure 15 is a neighbor-joining tree for *Thermococcales*.

Figure 16 shows sequences of exemplary sequences of the invention.

Figure 17 illustrates methods of the invention for liquefaction saccharification of starch, as described in detail, below.

Figure 18 illustrates Table 7, which lists the relative percent identities of exemplary sequences of the invention, as described in Example 8, below.

Figure 19 shows the pH profile of tested amylases of the invention and a commercial benchmark enzyme, as described in Example 15, below.

Figure 20 shows the temperature activity profiles of exemplary amylases of the invention, as described in Example 15, below.

Figure 21 shows enzyme activity (of exemplary amylases of the invention) in the presence of EDTA, as described in Example 15, below.

Figure 22 shows enzyme activity (of exemplary amylases of the invention) in the presence of peroxide hydroxide, as described in Example 15, below.

Figure 23 shows enzyme activity (of exemplary amylases of the invention) in the ADW solution (distilled water, hardening solution, bleach, chelators, surfactants) with soluble substrate (BODIPY-starch), as described in Example 15, below.

Figure 24 shows the results of the wash tests with starch-coated slides using exemplary amylases of the invention, as described in Example 15, below.

Figure 25 illustrates an exemplary corn wet milling process of the invention (using at least one enzyme of the invention).

Figure 26, Figure 27 and Figure 28 illustrate alternative exemplary starch processes, including starch liquefaction processes, of the invention (using at least one enzyme of the invention), as described in detail, below.

Figure 29 shows data summarizing these findings comparing amylase SEQ ID NO:437 with TERMAMYL™ SC (Novozymes A/S, Denmark) amylase in dry mill ethanol processing, as described in Example 1, below.

Figure 30 illustrates a pH activity profile of an exemplary enzyme of the invention (SEQ ID NO:594) in acetate buffer and phosphate buffer to determine the relative rate for the glucoamylase at each pH, as discussed in detail in Example 16, below.

Figure 31 illustrates a temperature activity profile of an exemplary enzyme of the invention (SEQ ID NO:594) in acetate buffer, as discussed in detail in Example 16, below.

Figure 32 illustrates a temperature stability profile of an exemplary enzyme of the invention (SEQ ID NO:594), as discussed in detail in Example 16, below.

Figure 33 illustrates a substrate utilization activity profile of an exemplary enzyme of the invention (SEQ ID NO:594) using the dextrans maltose (G2), maltotriose

(G3), panose (Pan), maltotetraose (G4), and maltoheptaose (G7), as discussed in detail in Example 16, below.

Figure 34 illustrates an exemplary glucoamylase-encoding nucleic acid of the invention, the genomic sequence set forth in SEQ ID NO:587. Coding sequences (exons) are denoted with the single-letter amino acid below it. Intron sequences are underlined.

Figure 35 is a chart describing selected characteristics of exemplary nucleic acids and polypeptides of the invention, as described in further detail, below.

Like reference symbols in the various drawings indicate like elements.

DETAILED DESCRIPTION

The invention provides amylase enzymes, e.g., an alpha amylases, polynucleotides encoding the enzymes, methods of making and using these polynucleotides and polypeptides. The invention is directed to novel polypeptides having an amylase activity, e.g., an alpha amylase activity, nucleic acids encoding them and antibodies that bind to them. The polypeptides of the invention can be used in a variety of diagnostic, therapeutic, and industrial contexts. The polypeptides of the invention can be used as, e.g., an additive for a detergent, for processing foods and for chemical synthesis utilizing a reverse reaction. Additionally, the polypeptides of the invention can be used in fabric treatment, alcohol production, and as additives to food or animal feed.

In one aspect, the amylases of the invention are active at a high and/or at a low temperature, or, over a wide range of temperature. For example, they can be active in the temperatures ranging between 20°C to 90°C, between 30°C to 80°C, or between 40°C to 70°C. The invention also provides amylases that have activity at alkaline pHs or at acidic pHs, e.g., low water acidity. In alternative aspects, the amylases of the invention can have activity in acidic pHs as low as pH 5.0, pH 4.5, pH 4.0, and pH 3.5. In alternative aspects, the amylases of the invention can have activity in alkaline pHs as high as pH 9.5, pH 10, pH 10.5, and pH 11. In one aspect, the amylases of the invention are active in the temperature range of between about 40°C to about 70°C under conditions of low water activity (low water content).

The invention also provides methods for further modifying the exemplary amylases of the invention to generate proteins with desirable properties. For example,

amylases generated by the methods of the invention can have altered enzymatic activity, thermal stability, pH/activity profile, pH/stability profile (such as increased stability at low, e.g. pH<6 or pH<5, or high, e.g. pH>9, pH values), stability towards oxidation, Ca²⁺ dependency, specific activity and the like. The invention provides for altering any property of interest. For instance, the alteration may result in a variant which, as compared to a parent enzyme, has altered enzymatic activity, or, pH or temperature activity profiles.

Definitions

The term "amylase" includes all polypeptides, e.g., enzymes, which catalyze the hydrolysis of a polysaccharide, e.g., a starch. The term "amylase" includes polypeptides having an α -amylase activity, a β -amylase activity, a glucoamylase activity, a 1,4- α -D-glucan glucohydrolase activity, an exoamylase activity, a glucan α -maltotetrahydrolase activity, a maltase activity, an isomaltase activity, a glucan 1, 4, α -glucosidase activity, an α -glucosidase activity, a sucrase activity or an agarase activity (e.g., a β -agarase activity). For example, an amylase activity of the invention includes α -amylase activity, including the ability to hydrolyze internal α -1,4-glucosidic linkages in starch to produce smaller molecular weight malto-dextrins. In one aspect, the α -amylase activity includes hydrolyzing internal α -1,4-glucosidic linkages in starch at random. An amylase activity of the invention includes polypeptides having glucoamylase activity, such as the ability to hydrolyze glucose polymers linked by α -1,4- and α -1,6-glucosidic bonds. In one aspect, the polypeptides of the invention have glucoamylase activity, hydrolyzing internal α -1,4-glucosidic linkages to yield smaller molecular weight malto-dextrins. An amylase activity of the invention also includes glucan 1,4- α -glucosidase activity, or, 1,4- α -D-glucan glucohydrolase, commonly called glucoamylase but also called amyloglucosidase and β -amylase that, in one aspect, releases β -D-glucose from 1,4- α -, 1,6- α - and 1,3- α -linked glucans. An amylase activity of the invention also includes exo-amylase activity.

In one aspect, the glucoamylase activity comprises catalysis of the hydrolysis of glucosidic bonds. The glucoamylase activity can comprise catalyzing the step-wise hydrolytic release of D-glucose from the non-reducing ends of starch or other related dextrins. The glucoamylase activity can comprise a 1,4- α -D-glucan glucohydrolase activity. The glucoamylase activity can comprise catalysis of the

hydrolysis of malto-dextrins resulting in the generation of free glucose. The glucoamylase activity can comprise an exoamylase activity. The glucoamylase activity can comprise an α -amylase or a β -amylase activity. The hydrolyzed glucosidic bonds can comprise α -1,4-glucosidic bonds or α -1,6-glucosidic bonds. The glucoamylase activity can comprise hydrolyzing glucosidic bonds in a starch. The glucoamylase activity can further comprise hydrolyzing glucosidic bonds in the starch to produce maltodextrines. The glucoamylase activity can comprise cleaving a maltose or a D-glucose unit from non-reducing end of the starch.

An amylase activity of the invention also includes hydrolyzing a polysaccharide, e.g., a starch, at high temperatures, low temperatures, alkaline pHs and at acidic pHs. For example, in one aspect, the invention provides polypeptides, and nucleic acids encoding them, having an amylase, e.g., a glucoamylase, activity which is thermostable. The polypeptide can retain an amylase activity under conditions comprising a temperature range of between about 37°C to about 95°C; between about 55°C to about 85°C, between about 70°C to about 95°C, or, between about 90°C to about 95°C. In another aspect, a polypeptide of the invention can have a glucoamylase activity which is thermotolerant. The polypeptide can retain an amylase, e.g., a glucoamylase, activity after exposure to a temperature in the range from greater than 37°C to about 95°C or anywhere in the range from greater than 55°C to about 85°C. In one aspect, the polypeptide retains an amylase activity after exposure to a temperature in the range from greater than 90°C to about 95°C at pH 4.5.

An "amylase variant" comprises an amino acid sequence which is derived from the amino acid sequence of a "precursor amylase". The precursor amylase can include naturally-occurring amylases and recombinant amylases. The amino acid sequence of the amylase variant can be "derived" from the precursor amylase amino acid sequence by the substitution, deletion or insertion of one or more amino acids of the precursor amino acid sequence. Such modification can be of the "precursor DNA sequence" which encodes the amino acid sequence of the precursor amylase rather than manipulation of the precursor amylase enzyme per se. Suitable methods for such manipulation of the precursor DNA sequence include methods disclosed herein, as well as methods known to those skilled in the art.

The term "antibody" includes a peptide or polypeptide derived from, modeled after or substantially encoded by an immunoglobulin gene or immunoglobulin

genes, or fragments thereof, capable of specifically binding an antigen or epitope, see, e.g. Fundamental Immunology, Third Edition, W.E. Paul, ed., Raven Press, N.Y. (1993); Wilson (1994) J. Immunol. Methods 175:267-273; Yarmush (1992) J. Biochem. Biophys. Methods 25:85-97. The term antibody includes antigen-binding portions, i.e.,
5 "antigen binding sites," (e.g., fragments, subsequences, complementarity determining regions (CDRs)) that retain capacity to bind antigen, including (i) a Fab fragment, a monovalent fragment consisting of the VL, VH, CL and CH1 domains; (ii) a F(ab')₂ fragment, a bivalent fragment comprising two Fab fragments linked by a disulfide bridge at the hinge region; (iii) a Fd fragment consisting of the VH and CH1 domains; (iv) a Fv
10 fragment consisting of the VL and VH domains of a single arm of an antibody, (v) a dAb fragment (Ward et al., (1989) Nature 341:544-546), which consists of a VH domain; and (vi) an isolated complementarity determining region (CDR). Single chain antibodies are also included by reference in the term "antibody."

The terms "array" or "microarray" or "biochip" or "chip" as used herein is
15 a plurality of target elements, each target element comprising a defined amount of one or more polypeptides (including antibodies) or nucleic acids immobilized onto a defined area of a substrate surface, as discussed in further detail, below.

As used herein, the terms "computer," "computer program" and
"processor" are used in their broadest general contexts and incorporate all such devices,
20 as described in detail, below. A "coding sequence of" or a "sequence encodes" a particular polypeptide or protein, is a nucleic acid sequence which is transcribed and translated into a polypeptide or protein when placed under the control of appropriate regulatory sequences.

The term "expression cassette" as used herein refers to a nucleotide
25 sequence which is capable of affecting expression of a structural gene (i.e., a protein coding sequence, such as an amylase of the invention) in a host compatible with such sequences. Expression cassettes include at least a promoter operably linked with the polypeptide coding sequence; and, optionally, with other sequences, e.g., transcription termination signals. Additional factors necessary or helpful in effecting expression may
30 also be used, e.g., enhancers. Thus, expression cassettes also include plasmids, expression vectors, recombinant viruses, any form of recombinant "naked DNA" vector, and the like.

"Operably linked" as used herein refers to a functional relationship between two or more nucleic acid (e.g., DNA) segments. Typically, it refers to the functional relationship of transcriptional regulatory sequence to a transcribed sequence. For example, a promoter is operably linked to a coding sequence, such as a nucleic acid of the invention, if it stimulates or modulates the transcription of the coding sequence in an appropriate host cell or other expression system. Generally, promoter transcriptional regulatory sequences that are operably linked to a transcribed sequence are physically contiguous to the transcribed sequence, i.e., they are *cis*-acting. However, some transcriptional regulatory sequences, such as enhancers, need not be physically contiguous or located in close proximity to the coding sequences whose transcription they enhance.

A "vector" comprises a nucleic acid which can infect, transfect, transiently or permanently transduce a cell. It will be recognized that a vector can be a naked nucleic acid, or a nucleic acid complexed with protein or lipid. The vector optionally comprises viral or bacterial nucleic acids and/or proteins, and/or membranes (e.g., a cell membrane, a viral lipid envelope, etc.). Vectors include, but are not limited to replicons (e.g., RNA replicons, bacteriophages) to which fragments of DNA may be attached and become replicated. Vectors thus include, but are not limited to RNA, autonomous self-replicating circular or linear DNA or RNA (e.g., plasmids, viruses, and the like, see, e.g., U.S. Patent No. 5,217,879), and include both the expression and non-expression plasmids. Where a recombinant microorganism or cell culture is described as hosting an "expression vector" this includes both extra-chromosomal circular and linear DNA and DNA that has been incorporated into the host chromosome(s). Where a vector is being maintained by a host cell, the vector may either be stably replicated by the cells during mitosis as an autonomous structure, or is incorporated within the host's genome.

As used herein, the term "promoter" includes all sequences capable of driving transcription of a coding sequence in a cell, e.g., a plant cell. Thus, promoters used in the constructs of the invention include *cis*-acting transcriptional control elements and regulatory sequences that are involved in regulating or modulating the timing and/or rate of transcription of a gene. For example, a promoter can be a *cis*-acting transcriptional control element, including an enhancer, a promoter, a transcription terminator, an origin of replication, a chromosomal integration sequence, 5' and 3' untranslated regions, or an intronic sequence, which are involved in transcriptional

regulation. These cis-acting sequences typically interact with proteins or other biomolecules to carry out (turn on/off, regulate, modulate, etc.) transcription.

“Constitutive” promoters are those that drive expression continuously under most environmental conditions and states of development or cell differentiation. “Inducible” or “regulatable” promoters direct expression of the nucleic acid of the invention under the influence of environmental conditions or developmental conditions. Examples of environmental conditions that may affect transcription by inducible promoters include anaerobic conditions, elevated temperature, drought, or the presence of light.

“Tissue-specific” promoters are transcriptional control elements that are only active in particular cells or tissues or organs, e.g., in plants or animals. Tissue-specific regulation may be achieved by certain intrinsic factors which ensure that genes encoding proteins specific to a given tissue are expressed. Such factors are known to exist in mammals and plants so as to allow for specific tissues to develop.

The term “plant” includes whole plants, plant parts (e.g., leaves, stems, flowers, roots, etc.), plant protoplasts, seeds and plant cells and progeny of same. The class of plants which can be used in the method of the invention is generally as broad as the class of higher plants amenable to transformation techniques, including angiosperms (monocotyledonous and dicotyledonous plants), as well as gymnosperms. It includes plants of a variety of ploidy levels, including polyploid, diploid, haploid and hemizygous states. As used herein, the term “transgenic plant” includes plants or plant cells into which a heterologous nucleic acid sequence has been inserted, e.g., the nucleic acids and various recombinant constructs (e.g., expression cassettes) of the invention.

“Plasmids” can be commercially available, publicly available on an unrestricted basis, or can be constructed from available plasmids in accord with published procedures. Equivalent plasmids to those described herein are known in the art and will be apparent to the ordinarily skilled artisan.

The term “gene” includes a nucleic acid sequence comprising a segment of DNA involved in producing a transcription product (e.g., a message), which in turn is translated to produce a polypeptide chain, or regulates gene transcription, reproduction or stability. Genes can include regions preceding and following the coding region, such as leader and trailer, promoters and enhancers, as well as, where applicable, intervening sequences (introns) between individual coding segments (exons).

The phrases "nucleic acid" or "nucleic acid sequence" includes oligonucleotide, nucleotide, polynucleotide, or to a fragment of any of these, to DNA or RNA (e.g., mRNA, rRNA, tRNA) of genomic or synthetic origin which may be single-stranded or double-stranded and may represent a sense or antisense strand, to peptide nucleic acid (PNA), or to any DNA-like or RNA-like material, natural or synthetic in origin, including, e.g., iRNA, ribonucleoproteins (e.g., iRNPs). The term encompasses nucleic acids, i.e., oligonucleotides, containing known analogues of natural nucleotides. The term also encompasses nucleic-acid-like structures with synthetic backbones, see e.g., Mata (1997) *Toxicol. Appl. Pharmacol.* 144:189-197; Strauss-Soukup (1997) *Biochemistry* 36:8692-8698; Samstag (1996) *Antisense Nucleic Acid Drug Dev* 6:153-156.

"Amino acid" or "amino acid sequence" include an oligopeptide, peptide, polypeptide, or protein sequence, or to a fragment, portion, or subunit of any of these, and to naturally occurring or synthetic molecules. The terms "polypeptide" and "protein" include amino acids joined to each other by peptide bonds or modified peptide bonds, i.e., peptide isosteres, and may contain modified amino acids other than the 20 gene-encoded amino acids. The term "polypeptide" also includes peptides and polypeptide fragments, motifs and the like. The term also includes glycosylated polypeptides. The peptides and polypeptides of the invention also include all "mimetic" and "peptidomimetic" forms, as described in further detail, below.

The term "isolated" includes a material removed from its original environment, e.g., the natural environment if it is naturally occurring. For example, a naturally occurring polynucleotide or polypeptide present in a living animal is not isolated, but the same polynucleotide or polypeptide, separated from some or all of the coexisting materials in the natural system, is isolated. Such polynucleotides could be part of a vector and/or such polynucleotides or polypeptides could be part of a composition, and still be isolated in that such vector or composition is not part of its natural environment. As used herein, an isolated material or composition can also be a "purified" composition, i.e., it does not require absolute purity; rather, it is intended as a relative definition. Individual nucleic acids obtained from a library can be conventionally purified to electrophoretic homogeneity. In alternative aspects, the invention provides nucleic acids which have been purified from genomic DNA or from other sequences in a

library or other environment by at least one, two, three, four, five or more orders of magnitude.

As used herein, the term "recombinant" can include nucleic acids adjacent to a "backbone" nucleic acid to which it is not adjacent in its natural environment. In one aspect, nucleic acids represent 5% or more of the number of nucleic acid inserts in a population of nucleic acid "backbone molecules." "Backbone molecules" according to the invention include nucleic acids such as expression vectors, self-replicating nucleic acids, viruses, integrating nucleic acids, and other vectors or nucleic acids used to maintain or manipulate a nucleic acid insert of interest. In one aspect, the enriched nucleic acids represent 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more of the number of nucleic acid inserts in the population of recombinant backbone molecules. "Recombinant" polypeptides or proteins refer to polypeptides or proteins produced by recombinant DNA techniques; e.g., produced from cells transformed by an exogenous DNA construct encoding the desired polypeptide or protein. "Synthetic" polypeptides or protein are those prepared by chemical synthesis, as described in further detail, below.

A promoter sequence can be "operably linked to" a coding sequence when RNA polymerase which initiates transcription at the promoter will transcribe the coding sequence into mRNA, as discussed further, below.

"Oligonucleotide" includes either a single stranded polydeoxynucleotide or two complementary polydeoxynucleotide strands which may be chemically synthesized. Such synthetic oligonucleotides have no 5' phosphate and thus will not ligate to another oligonucleotide without adding a phosphate with an ATP in the presence of a kinase. A synthetic oligonucleotide can ligate to a fragment that has not been dephosphorylated.

The phrase "substantially identical" in the context of two nucleic acids or polypeptides, can refer to two or more sequences that have, e.g., at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more nucleotide or amino acid residue (sequence) identity, when

compared and aligned for maximum correspondence, as measured using one any known sequence comparison algorithm, as discussed in detail below, or by visual inspection. In alternative aspects, the invention provides nucleic acid and polypeptide sequences having substantial identity to an exemplary sequence of the invention over a region of at least about 10, 20, 30, 40, 50, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000 or more residues, or a region ranging from between about 50 residues to the full length of the nucleic acid or polypeptide. Nucleic acid sequences of the invention can be substantially identical over the entire length of a polypeptide coding region.

A "substantially identical" amino acid sequence also can include a sequence that differs from a reference sequence by one or more conservative or non-conservative amino acid substitutions, deletions, or insertions, particularly when such a substitution occurs at a site that is not the active site of the molecule, and provided that the polypeptide essentially retains its functional properties. A conservative amino acid substitution, for example, substitutes one amino acid for another of the same class (e.g., substitution of one hydrophobic amino acid, such as isoleucine, valine, leucine, or methionine, for another, or substitution of one polar amino acid for another, such as substitution of arginine for lysine, glutamic acid for aspartic acid or glutamine for asparagine). One or more amino acids can be deleted, for example, from an amylase, resulting in modification of the structure of the polypeptide, without significantly altering its biological activity. For example, amino- or carboxyl-terminal amino acids that are not required for amylase activity can be removed.

"Hybridization" includes the process by which a nucleic acid strand joins with a complementary strand through base pairing. Hybridization reactions can be sensitive and selective so that a particular sequence of interest can be identified even in samples in which it is present at low concentrations. Stringent conditions can be defined by, for example, the concentrations of salt or formamide in the prehybridization and hybridization solutions, or by the hybridization temperature, and are well known in the art. For example, stringency can be increased by reducing the concentration of salt, increasing the concentration of formamide, or raising the hybridization temperature, altering the time of hybridization, as described in detail, below. In alternative aspects, nucleic acids of the invention are defined by their ability to hybridize under various stringency conditions (e.g., high, medium, and low), as set forth herein.

“Variant” includes polynucleotides or polypeptides of the invention modified at one or more base pairs, codons, introns, exons, or amino acid residues (respectively) yet still retain the biological activity of an amylase of the invention. Variants can be produced by any number of means included methods such as, for example, error-prone PCR, shuffling, oligonucleotide-directed mutagenesis, assembly PCR, sexual PCR mutagenesis, in vivo mutagenesis, cassette mutagenesis, recursive ensemble mutagenesis, exponential ensemble mutagenesis, site-specific mutagenesis, gene reassembly, GSSM and any combination thereof. Techniques for producing variant amylase having activity at a pH or temperature, for example, that is different from a wild-type amylase, are included herein.

The term “saturation mutagenesis” or “GSSM” includes a method that uses degenerate oligonucleotide primers to introduce point mutations into a polynucleotide, as described in detail, below.

The term “optimized directed evolution system” or “optimized directed evolution” includes a method for reassembling fragments of related nucleic acid sequences, e.g., related genes, and explained in detail, below.

The term “synthetic ligation reassembly” or “SLR” includes a method of ligating oligonucleotide fragments in a non-stochastic fashion, and explained in detail, below.

Generating and Manipulating Nucleic Acids

In one aspect, the invention provides isolated or recombinant nucleic acids comprising a nucleic acid sequence having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an exemplary nucleic acid of the invention over a region of at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more, residues. In one aspect, the nucleic acid encodes at least one polypeptide having an amylase activity, e.g., an alpha amylase activity.

For example, the following table describes some exemplary amylase-encoding nucleic acids of the invention, e.g., the invention provides an amylase having a sequence as set forth in SEQ ID NO:474, having an exemplary coding sequence as set forth in SEQ ID NO:473, and in one aspect is encoded by a gene, including introns and exons, having a sequence as set forth in SEQ ID NO:467 (including exons having sequences as set forth in SEQ ID NO:468, SEQ ID NO:469, SEQ ID NO:470, SEQ ID NO:471 and SEQ ID NO:472); etc.:

Amylase	SEQ ID NO: of full gene (exons and introns)	SEQ ID NOS: of exon sequences	SEQ ID NO: of DNA sequence of coding sequence (exons only)	SEQ ID NO: of protein sequence of coding sequence (exons only)	TOTAL
A	460, 461	N/A	460	461	460, 461
B	462	463, 464	465	466	462-466
C	467	468-472	473	474	467-474
D	475	476-477	478	479	475-479
E	480	481-483	484	485	480-485
F	486	487-491	492	493	486-493
G	494	495-497	498	499	494-499
H	500	501-508	509	510	500-510
I	511	512-514	515	516	511-516
J	517, 518	N/A	517	518	517, 518
K	519	520-521	522	523	519-523
L	524	525-526	527	528	524-528
M	529	530-531	532	533	529-533
N	534	535-538	539	540	534-540
O	541	542-543	544	545	541-545
P	546	547-551	552	553	546-553
Q	554	555-557	558	559	554-559
R	560	561-564	565	566	560-566
S	587	588-592	593	594	587-594

The above listed amylases (described as A thru S) and the nucleic acids that encode them have a common novelty in that they were initially isolated/ derived from fungal sources.

The invention also provides glucoamylases, such as the enzyme having a sequence as set forth in SEQ ID NO:594 encoded by the 4111 residues of the genomic SEQ ID NO:587, or, the 1854 residue long cDNA of SEQ ID NO:593). The genomic SEQ ID NO:587, comprises introns and exons, and the exons can be described as encoding polypeptide fragments having a sequence as set forth in SEQ ID NO:588, SEQ

ID NO:589, SEQ ID NO:590, SEQ ID NO:591, SEQ ID NO:592. In one aspect, the “mature” processed glucoamylase consisting of residues 32 to 617 of SEQ ID NO: 594.

The invention provides isolated and recombinant nucleic acids, including expression cassettes such as expression vectors encoding the polypeptides of the invention. The invention provides probes comprising or consisting of nucleic acids of the invention. The invention also includes methods for discovering new amylase sequences using the nucleic acids of the invention. The invention also includes methods for inhibiting the expression of amylase genes, transcripts and polypeptides using the nucleic acids of the invention. Also provided are methods for modifying the nucleic acids of the invention by, e.g., synthetic ligation reassembly, optimized directed evolution system and/or gene site saturation mutagenesis (GSSM™).

The nucleic acids of the invention can be made, isolated and/or manipulated by, e.g., cloning and expression of cDNA libraries, amplification of message or genomic DNA by PCR, and the like. In practicing the methods of the invention, homologous genes can be modified by manipulating a template nucleic acid, as described herein. The invention can be practiced in conjunction with any method or protocol or device known in the art, which are well described in the scientific and patent literature.

General Techniques

The nucleic acids used to practice this invention, whether RNA, iRNA, antisense nucleic acid, cDNA, genomic DNA, vectors, viruses or hybrids thereof, may be isolated from a variety of sources, genetically engineered, amplified, and/or expressed/generated recombinantly. Recombinant polypeptides generated from these nucleic acids can be individually isolated or cloned and tested for a desired activity. Any recombinant expression system can be used, including bacterial, mammalian, yeast, insect or plant cell expression systems.

Alternatively, these nucleic acids can be synthesized *in vitro* by well-known chemical synthesis techniques, as described in, e.g., Adams (1983) J. Am. Chem. Soc. 105:661; Belousov (1997) Nucleic Acids Res. 25:3440-3444; Frenkel (1995) Free Radic. Biol. Med. 19:373-380; Blommers (1994) Biochemistry 33:7886-7896; Narang (1979) Meth. Enzymol. 68:90; Brown (1979) Meth. Enzymol. 68:109; Beaucage (1981) Tetra. Lett. 22:1859; U.S. Patent No. 4,458,066.

Techniques for the manipulation of nucleic acids, such as, e.g., subcloning, labeling probes (e.g., random-primer labeling using Klenow polymerase, nick translation, amplification), sequencing, hybridization and the like are well described in the scientific and patent literature, see, e.g., Sambrook, ed., MOLECULAR CLONING: A

- 5 LABORATORY MANUAL (2ND ED.), Vols. 1-3, Cold Spring Harbor Laboratory, (1989); CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, Ausubel, ed. John Wiley & Sons, Inc., New York (1997); LABORATORY TECHNIQUES IN BIOCHEMISTRY AND MOLECULAR BIOLOGY: HYBRIDIZATION WITH NUCLEIC ACID PROBES, Part I. Theory and Nucleic Acid Preparation, Tijssen, ed. Elsevier, N.Y. (1993).

- Another useful means of obtaining and manipulating nucleic acids used to practice the methods of the invention is to clone from genomic samples, and, if desired, screen and re-clone inserts isolated or amplified from, e.g., genomic clones or cDNA clones. Sources of nucleic acid used in the methods of the invention include genomic or
- 15 cDNA libraries contained in, e.g., mammalian artificial chromosomes (MACs), see, e.g., U.S. Patent Nos. 5,721,118; 6,025,155; human artificial chromosomes, see, e.g., Rosenfeld (1997) Nat. Genet. 15:333-335; yeast artificial chromosomes (YAC); bacterial artificial chromosomes (BAC); P1 artificial chromosomes, see, e.g., Woon (1998) Genomics 50:306-316; P1-derived vectors (PACs), see, e.g., Kern (1997) Biotechniques
- 20 23:120-124; cosmids, recombinant viruses, phages or plasmids.

In one aspect, a nucleic acid encoding a polypeptide of the invention is assembled in appropriate phase with a leader sequence capable of directing secretion of the translated polypeptide or fragment thereof.

- The invention provides fusion proteins and nucleic acids encoding them.
- 25 A polypeptide of the invention can be fused to a heterologous peptide or polypeptide, such as N-terminal identification peptides which impart desired characteristics, such as increased stability or simplified purification. Peptides and polypeptides of the invention can also be synthesized and expressed as fusion proteins with one or more additional domains linked thereto for, e.g., producing a more immunogenic peptide, to more readily
- 30 isolate a recombinantly synthesized peptide, to identify and isolate antibodies and antibody-expressing B cells, and the like. Detection and purification facilitating domains include, e.g., metal chelating peptides such as polyhistidine tracts and histidine-tryptophan modules that allow purification on immobilized metals, protein A domains

that allow purification on immobilized immunoglobulin, and the domain utilized in the FLAGS extension/affinity purification system (Immunex Corp, Seattle WA). The inclusion of a cleavable linker sequences such as Factor Xa or enterokinase (Invitrogen, San Diego CA) between a purification domain and the motif-comprising peptide or polypeptide to facilitate purification. For example, an expression vector can include an epitope-encoding nucleic acid sequence linked to six histidine residues followed by a thioredoxin and an enterokinase cleavage site (see e.g., Williams (1995) *Biochemistry* 34:1787-1797; Dobeli (1998) *Protein Expr. Purif.* 12:404-414). The histidine residues facilitate detection and purification while the enterokinase cleavage site provides a means for purifying the epitope from the remainder of the fusion protein. Technology pertaining to vectors encoding fusion proteins and application of fusion proteins are well described in the scientific and patent literature, see e.g., Kroll (1993) *DNA Cell. Biol.*, 12:441-53.

Transcriptional and translational control sequences

The invention provides nucleic acid (e.g., DNA) sequences of the invention operatively linked to expression (e.g., transcriptional or translational) control sequence(s), e.g., promoters or enhancers, to direct or modulate RNA synthesis/ expression. The expression control sequence can be in an expression vector. Exemplary bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, PL and trp. Exemplary eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein I.

Promoters suitable for expressing a polypeptide in bacteria include the *E. coli* lac or trp promoters, the lacI promoter, the lacZ promoter, the T3 promoter, the T7 promoter, the gpt promoter, the lambda PR promoter, the lambda PL promoter, promoters from operons encoding glycolytic enzymes such as 3-phosphoglycerate kinase (PGK), and the acid phosphatase promoter. Eukaryotic promoters include the CMV immediate early promoter, the HSV thymidine kinase promoter, heat shock promoters, the early and late SV40 promoter, LTRs from retroviruses, and the mouse metallothionein-I promoter. Other promoters known to control expression of genes in prokaryotic or eukaryotic cells or their viruses may also be used.

Tissue-Specific Plant Promoters

The invention provides expression cassettes that can be expressed in a tissue-specific manner, e.g., that can express an amylase of the invention in a tissue-

specific manner. The invention also provides plants or seeds that express an amylase of the invention in a tissue-specific manner. The tissue-specificity can be seed specific, stem specific, leaf specific, root specific, fruit specific and the like.

In one aspect, a constitutive promoter such as the CaMV 35S promoter can be used for expression in specific parts of the plant or seed or throughout the plant. For example, for overexpression, a plant promoter fragment can be employed which will direct expression of a nucleic acid in some or all tissues of a plant, e.g., a regenerated plant. Such promoters are referred to herein as "constitutive" promoters and are active under most environmental conditions and states of development or cell differentiation. Examples of constitutive promoters include the cauliflower mosaic virus (CaMV) 35S transcription initiation region, the 1'- or 2'- promoter derived from T-DNA of *Agrobacterium tumefaciens*, and other transcription initiation regions from various plant genes known to those of skill. Such genes include, e.g., *ACT11* from *Arabidopsis* (Huang (1996) *Plant Mol. Biol.* 33:125-139); *Cat3* from *Arabidopsis* (GenBank No. U43147, Zhong (1996) *Mol. Gen. Genet.* 251:196-203); the gene encoding stearyl-acyl carrier protein desaturase from *Brassica napus* (Genbank No. X74782, Solocombe (1994) *Plant Physiol.* 104:1167-1176); *GPc1* from maize (GenBank No. X15596; Martinez (1989) *J. Mol. Biol.* 208:551-565); the *Gpc2* from maize (GenBank No. U45855, Manjunath (1997) *Plant Mol. Biol.* 33:97-112); plant promoters described in U.S. Patent Nos. 4,962,028; 5,633,440.

The invention uses tissue-specific or constitutive promoters derived from viruses which can include, e.g., the tobamovirus subgenomic promoter (Kumagai (1995) *Proc. Natl. Acad. Sci. USA* 92:1679-1683; the rice tungro bacilliform virus (RTBV), which replicates only in phloem cells in infected rice plants, with its promoter which drives strong phloem-specific reporter gene expression; the cassava vein mosaic virus (CVMV) promoter, with highest activity in vascular elements, in leaf mesophyll cells, and in root tips (Verdaguer (1996) *Plant Mol. Biol.* 31:1129-1139).

Alternatively, the plant promoter may direct expression of amylase-expressing nucleic acid in a specific tissue, organ or cell type (i.e. tissue-specific promoters) or may be otherwise under more precise environmental or developmental control or under the control of an inducible promoter. Examples of environmental conditions that may affect transcription include anaerobic conditions, elevated temperature, the presence of light, or sprayed with chemicals/hormones. For example, the

invention incorporates the drought-inducible promoter of maize (Busk (1997) supra); the cold, drought, and high salt inducible promoter from potato (Kirch (1997) Plant Mol. Biol. 33:897 909).

Tissue-specific promoters can promote transcription only within a certain time frame of developmental stage within that tissue. See, e.g., Blazquez (1998) Plant Cell 10:791-800, characterizing the *Arabidopsis* LEAFY gene promoter. See also Cardon (1997) *Plant J* 12:367-77, describing the transcription factor SPL3, which recognizes a conserved sequence motif in the promoter region of the *A. thaliana* floral meristem identity gene AP1; and Mandel (1995) Plant Molecular Biology, Vol. 29, pp 995-1004, describing the meristem promoter eIF4. Tissue specific promoters which are active throughout the life cycle of a particular tissue can be used. In one aspect, the nucleic acids of the invention are operably linked to a promoter active primarily only in cotton fiber cells. In one aspect, the nucleic acids of the invention are operably linked to a promoter active primarily during the stages of cotton fiber cell elongation, e.g., as described by Rinehart (1996) supra. The nucleic acids can be operably linked to the Fbl2A gene promoter to be preferentially expressed in cotton fiber cells (Ibid) . See also, John (1997) Proc. Natl. Acad. Sci. USA 89:5769-5773; John, et al., U.S. Patent Nos. 5,608,148 and 5,602,321, describing cotton fiber-specific promoters and methods for the construction of transgenic cotton plants. Root-specific promoters may also be used to express the nucleic acids of the invention. Examples of root-specific promoters include the promoter from the alcohol dehydrogenase gene (DeLisle (1990) Int. Rev. Cytol. 123:39-60). Other promoters that can be used to express the nucleic acids of the invention include, e.g., ovule-specific, embryo-specific, endosperm-specific, integument-specific, seed coat-specific promoters, or some combination thereof; a leaf-specific promoter (see, e.g., Busk (1997) Plant J. 11:1285 1295, describing a leaf-specific promoter in maize); the ORF13 promoter from *Agrobacterium rhizogenes* (which exhibits high activity in roots, see, e.g., Hansen (1997) supra); a maize pollen specific promoter (see, e.g., Guerrero (1990) Mol. Gen. Genet. 224:161 168); a tomato promoter active during fruit ripening, senescence and abscission of leaves and, to a lesser extent, of flowers can be used (see, e.g., Blume (1997) Plant J. 12:731 746); a pistil-specific promoter from the potato SK2 gene (see, e.g., Ficker (1997) Plant Mol. Biol. 35:425 431); the Blec4 gene from pea, which is active in epidermal tissue of vegetative and floral shoot apices of transgenic alfalfa making it a useful tool to target the expression of

foreign genes to the epidermal layer of actively growing shoots or fibers; the ovule-specific BEL1 gene (see, e.g., Reiser (1995) Cell 83:735-742, GenBank No. U39944); and/or, the promoter in Klee, U.S. Patent No. 5,589,583, describing a plant promoter region is capable of conferring high levels of transcription in meristematic tissue and/or rapidly dividing cells.

Alternatively, plant promoters which are inducible upon exposure to plant hormones, such as auxins, are used to express the nucleic acids of the invention. For example, the invention can use the auxin-response elements E1 promoter fragment (AuxREs) in the soybean (*Glycine max* L.) (Liu (1997) Plant Physiol. 115:397-407); the auxin-responsive *Arabidopsis* GST6 promoter (also responsive to salicylic acid and hydrogen peroxide) (Chen (1996) Plant J. 10: 955-966); the auxin-inducible parC promoter from tobacco (Sakai (1996) 37:906-913); a plant biotin response element (Streit (1997) Mol. Plant Microbe Interact. 10:933-937); and, the promoter responsive to the stress hormone abscisic acid (Sheen (1996) Science 274:1900-1902).

5 The nucleic acids of the invention can also be operably linked to plant promoters which are inducible upon exposure to chemicals reagents which can be applied to the plant, such as herbicides or antibiotics. For example, the maize In2-2 promoter, activated by benzenesulfonamide herbicide safeners, can be used (De Veylder (1997) Plant Cell Physiol. 38:568-577); application of different herbicide safeners induces
20 distinct gene expression patterns, including expression in the root, hydathodes, and the shoot apical meristem. Coding sequence can be under the control of, e.g., a tetracycline-inducible promoter, e.g., as described with transgenic tobacco plants containing the *Avena sativa* L. (oat) arginine decarboxylase gene (Masgrau (1997) Plant J. 11:465-473); or, a salicylic acid-responsive element (Stange (1997) Plant J.
25 11:1315-1324). Using chemically- (e.g., hormone- or pesticide-) induced promoters, i.e., promoter responsive to a chemical which can be applied to the transgenic plant in the field, expression of a polypeptide of the invention can be induced at a particular stage of development of the plant. Thus, the invention also provides for transgenic plants containing an inducible gene encoding for polypeptides of the invention whose host range
30 is limited to target plant species, such as corn, rice, barley, wheat, potato or other crops, inducible at any stage of development of the crop.

One of skill will recognize that a tissue-specific plant promoter may drive expression of operably linked sequences in tissues other than the target tissue. Thus, a

tissue-specific promoter is one that drives expression preferentially in the target tissue or cell type, but may also lead to some expression in other tissues as well.

The nucleic acids of the invention can also be operably linked to plant promoters which are inducible upon exposure to chemicals reagents. These reagents include, e.g., herbicides, synthetic auxins, or antibiotics which can be applied, e.g., sprayed, onto transgenic plants. Inducible expression of the amylase-producing nucleic acids of the invention will allow the grower to select plants with the optimal starch / sugar ratio. The development of plant parts can thus controlled. In this way the invention provides the means to facilitate the harvesting of plants and plant parts. For example, in various embodiments, the maize In2-2 promoter, activated by benzenesulfonamide herbicide safeners, is used (De Veylder (1997) *Plant Cell Physiol.* 38:568-577); application of different herbicide safeners induces distinct gene expression patterns, including expression in the root, hydathodes, and the shoot apical meristem. Coding sequences of the invention are also under the control of a tetracycline-inducible promoter, e.g., as described with transgenic tobacco plants containing the *Avena sativa* L. (oat) arginine decarboxylase gene (Masgrau (1997) *Plant J.* 11:465-473); or, a salicylic acid-responsive element (Stange (1997) *Plant J.* 11:1315-1324).

If proper polypeptide expression is desired, a polyadenylation region at the 3'-end of the coding region should be included. The polyadenylation region can be derived from the natural gene, from a variety of other plant genes, or from genes in the *Agrobacterium* T-DNA.

Expression vectors and cloning vehicles

The invention provides expression vectors and cloning vehicles comprising nucleic acids of the invention, e.g., sequences encoding the amylases of the invention. Expression vectors and cloning vehicles of the invention can comprise viral particles, baculovirus, phage, plasmids, phagemids, cosmids, fosmids, bacterial artificial chromosomes, viral DNA (e.g., vaccinia, adenovirus, fowl pox virus, pseudorabies and derivatives of SV40), P1-based artificial chromosomes, yeast plasmids, yeast artificial chromosomes, and any other vectors specific for specific hosts of interest (such as bacillus, *Aspergillus* and yeast). Vectors of the invention can include chromosomal, non-chromosomal and synthetic DNA sequences. Large numbers of suitable vectors are known to those of skill in the art, and are commercially available. Exemplary vectors are

include: bacterial: pQE vectors (Qiagen), pBluescript plasmids, pNH vectors, (lambda-ZAP vectors (Stratagene); ptrc99a, pKK223-3, pDR540, pRIT2T (Pharmacia); Eukaryotic: pXT1, pSG5 (Stratagene), pSVK3, pBPV, pMSG, pSVLSV40 (Pharmacia). However, any other plasmid or other vector may be used so long as they are replicable and viable in the host. Low copy number or high copy number vectors may be employed with the present invention.

The expression vector can comprise a promoter, a ribosome binding site for translation initiation and a transcription terminator. The vector may also include appropriate sequences for amplifying expression. Mammalian expression vectors can comprise an origin of replication, any necessary ribosome binding sites, a polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking non-transcribed sequences. In some aspects, DNA sequences derived from the SV40 splice and polyadenylation sites may be used to provide the required non-transcribed genetic elements.

In one aspect, the expression vectors contain one or more selectable marker genes to permit selection of host cells containing the vector. Such selectable markers include genes encoding dihydrofolate reductase or genes conferring neomycin resistance for eukaryotic cell culture, genes conferring tetracycline or ampicillin resistance in *E. coli*, and the *S. cerevisiae* TRP1 gene. Promoter regions can be selected from any desired gene using chloramphenicol transferase (CAT) vectors or other vectors with selectable markers.

Vectors for expressing the polypeptide or fragment thereof in eukaryotic cells can also contain enhancers to increase expression levels. Enhancers are cis-acting elements of DNA, usually from about 10 to about 300 bp in length that act on a promoter to increase its transcription. Examples include the SV40 enhancer on the late side of the replication origin bp 100 to 270, the cytomegalovirus early promoter enhancer, the polyoma enhancer on the late side of the replication origin, and the adenovirus enhancers.

A nucleic acid sequence can be inserted into a vector by a variety of procedures. In general, the sequence is ligated to the desired position in the vector following digestion of the insert and the vector with appropriate restriction endonucleases. Alternatively, blunt ends in both the insert and the vector may be ligated. A variety of cloning techniques are known in the art, e.g., as described in Ausubel and

Sambrook. Such procedures and others are deemed to be within the scope of those skilled in the art.

The vector can be in the form of a plasmid, a viral particle, or a phage. Other vectors include chromosomal, non-chromosomal and synthetic DNA sequences, derivatives of SV40; bacterial plasmids, phage DNA, baculovirus, yeast plasmids, vectors derived from combinations of plasmids and phage DNA, viral DNA such as vaccinia, adenovirus, fowl pox virus, and pseudorabies. A variety of cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described by, e.g., Sambrook.

Particular bacterial vectors which can be used include the commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017), pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden), GEM1 (Promega Biotec, Madison, WI, USA) pQE70, pQE60, pQE-9 (Qiagen), pD10, psiX174 pBluescript II KS, pNH8A, pNH16a, pNH18A, pNH46A (Stratagene), ptrc99a, pKK223-3, pKK233-3, DR540, pRIT5 (Pharmacia), pKK232-8 and pCM7. Particular eukaryotic vectors include pSV2CAT, pOG44, pXT1, pSG (Stratagene) pSVK3, pBPV, pMSG, and pSVL (Pharmacia). However, any other vector may be used as long as it is replicable and viable in the host cell.

The nucleic acids of the invention can be expressed in expression cassettes, vectors or viruses and transiently or stably expressed in plant cells and seeds.

One exemplary transient expression system uses episomal expression systems, e.g., cauliflower mosaic virus (CaMV) viral RNA generated in the nucleus by transcription of an episomal mini-chromosome containing supercoiled DNA, see, e.g., Covey (1990) Proc. Natl. Acad. Sci. USA 87:1633-1637. Alternatively, coding sequences, i.e., all or sub-fragments of sequences of the invention can be inserted into a plant host cell genome becoming an integral part of the host chromosomal DNA. Sense or antisense transcripts can be expressed in this manner. A vector comprising the sequences (e.g., promoters or coding regions) from nucleic acids of the invention can comprise a marker gene that confers a selectable phenotype on a plant cell or a seed. For example, the marker may encode biocide resistance, particularly antibiotic resistance, such as resistance to kanamycin, G418, bleomycin, hygromycin, or herbicide resistance, such as resistance to chlorosulfuron or Basta.

Expression vectors capable of expressing nucleic acids and proteins in plants are well known in the art, and can include, e.g., vectors from *Agrobacterium* spp.,

potato virus X (see, *e.g.*, Angell (1997) EMBO J. 16:3675-3684), tobacco mosaic virus (see, *e.g.*, Casper (1996) Gene 173:69-73), tomato bushy stunt virus (see, *e.g.*, Hillman (1989) Virology 169:42-50), tobacco etch virus (see, *e.g.*, Dolja (1997) Virology 234:243-252), bean golden mosaic virus (see, *e.g.*, Morinaga (1993) Microbiol Immunol. 37:471-476), cauliflower mosaic virus (see, *e.g.*, Cecchini (1997) Mol. Plant Microbe Interact. 10:1094-1101), maize Ac/Ds transposable element (see, *e.g.*, Rubin (1997) Mol. Cell. Biol. 17:6294-6302; Kunze (1996) Curr. Top. Microbiol. Immunol. 204:161-194), and the maize suppressor-mutator (Spm) transposable element (see, *e.g.*, Schlappi (1996) Plant Mol. Biol. 32:717-725); and derivatives thereof.

In one aspect, the expression vector can have two replication systems to allow it to be maintained in two organisms, for example in mammalian or insect cells for expression and in a prokaryotic host for cloning and amplification. Furthermore, for integrating expression vectors, the expression vector can contain at least one sequence homologous to the host cell genome. It can contain two homologous sequences which flank the expression construct. The integrating vector can be directed to a specific locus in the host cell by selecting the appropriate homologous sequence for inclusion in the vector. Constructs for integrating vectors are well known in the art.

Expression vectors of the invention may also include a selectable marker gene to allow for the selection of bacterial strains that have been transformed, *e.g.*, genes which render the bacteria resistant to drugs such as ampicillin, chloramphenicol, erythromycin, kanamycin, neomycin and tetracycline. Selectable markers can also include biosynthetic genes, such as those in the histidine, tryptophan and leucine biosynthetic pathways.

Host cells and transformed cells

25 The invention also provides a transformed cell comprising a nucleic acid sequence of the invention, *e.g.*, a sequence encoding an amylase of the invention, or a vector of the invention. The host cell may be any of the host cells familiar to those skilled in the art, including prokaryotic cells, eukaryotic cells, such as bacterial cells, fungal cells, yeast cells, mammalian cells, insect cells, or plant cells. Exemplary bacterial
30 cells include *E. coli*, any *Streptomyces* or *Bacillus* (*e.g.*, *Bacillus cereus*, *Bacillus subtilis*), *Salmonella typhimurium* and various species within the genera *Bacillus*, *Streptomyces*, and *Staphylococcus*. Exemplary insect cells include *Drosophila S2* and

Spodoptera Sf9. Exemplary animal cells include CHO, COS or Bowes melanoma or any mouse or human cell line. The selection of an appropriate host is within the abilities of those skilled in the art. Techniques for transforming a wide variety of higher plant species are well known and described in the technical and scientific literature. See, e.g., Weising (1988) Ann. Rev. Genet. 22:421-477, U.S. Patent No. 5,750,870.

The vector can be introduced into the host cells using any of a variety of techniques, including transformation, transfection, transduction, viral infection, gene guns, or Ti-mediated gene transfer. Particular methods include calcium phosphate transfection, DEAE-Dextran mediated transfection, lipofection, or electroporation (Davis, L., Dibner, M., Battey, I., Basic Methods in Molecular Biology, (1986)).

In one aspect, the nucleic acids or vectors of the invention are introduced into the cells for screening, thus, the nucleic acids enter the cells in a manner suitable for subsequent expression of the nucleic acid. The method of introduction is largely dictated by the targeted cell type. Exemplary methods include CaPO_4 precipitation, liposome fusion, lipofection (e.g., LIPOFECTINTM), electroporation, viral infection, etc. The candidate nucleic acids may stably integrate into the genome of the host cell (for example, with retroviral introduction) or may exist either transiently or stably in the cytoplasm (i.e. through the use of traditional plasmids, utilizing standard regulatory sequences, selection markers, etc.). As many pharmaceutically important screens require human or model mammalian cell targets, retroviral vectors capable of transfecting such targets are preferred.

Where appropriate, the engineered host cells can be cultured in conventional nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes of the invention. Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter may be induced by appropriate means (e.g., temperature shift or chemical induction) and the cells may be cultured for an additional period to allow them to produce the desired polypeptide or fragment thereof.

Cells can be harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract is retained for further purification. Microbial cells employed for expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents. Such methods are well known to those skilled in the art. The expressed

polypeptide or fragment thereof can be recovered and purified from recombinant cell cultures by methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity chromatography, hydroxylapatite chromatography and lectin chromatography. Protein refolding steps can be used, as necessary, in completing configuration of the polypeptide. If desired, high performance liquid chromatography (HPLC) can be employed for final purification steps.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts and other cell lines capable of expressing proteins from a compatible vector, such as the C127, 3T3, CHO, HeLa and BHK cell lines.

The constructs in host cells can be used in a conventional manner to produce the gene product encoded by the recombinant sequence. Depending upon the host employed in a recombinant production procedure, the polypeptides produced by host cells containing the vector may be glycosylated or may be non-glycosylated. Polypeptides of the invention may or may not also include an initial methionine amino acid residue.

Cell-free translation systems can also be employed to produce a polypeptide of the invention. Cell-free translation systems can use mRNAs transcribed from a DNA construct comprising a promoter operably linked to a nucleic acid encoding the polypeptide or fragment thereof. In some aspects, the DNA construct may be linearized prior to conducting an in vitro transcription reaction. The transcribed mRNA is then incubated with an appropriate cell-free translation extract, such as a rabbit reticulocyte extract, to produce the desired polypeptide or fragment thereof.

The expression vectors can contain one or more selectable marker genes to provide a phenotypic trait for selection of transformed host cells such as dihydrofolate reductase or neomycin resistance for eukaryotic cell culture, or such as tetracycline or ampicillin resistance in *E. coli*.

Amplification of Nucleic Acids

In practicing the invention, nucleic acids of the invention and nucleic acids encoding the polypeptides of the invention, or modified nucleic acids of the invention, can be reproduced by amplification. Amplification can also be used to clone or modify

the nucleic acids of the invention. Thus, the invention provides amplification primer sequence pairs for amplifying nucleic acids of the invention. One of skill in the art can design amplification primer sequence pairs for any part of or the full length of these sequences.

Amplification reactions can also be used to quantify the amount of nucleic acid in a sample (such as the amount of message in a cell sample), label the nucleic acid (e.g., to apply it to an array or a blot), detect the nucleic acid, or quantify the amount of a specific nucleic acid in a sample. In one aspect of the invention, message isolated from a cell or a cDNA library are amplified.

The skilled artisan can select and design suitable oligonucleotide amplification primers. Amplification methods are also well known in the art, and include, e.g., polymerase chain reaction, PCR (see, e.g., PCR PROTOCOLS, A GUIDE TO METHODS AND APPLICATIONS, ed. Innis, Academic Press, N.Y. (1990) and PCR STRATEGIES (1995), ed. Innis, Academic Press, Inc., N.Y., ligase chain reaction (LCR) (see, e.g., Wu (1989) Genomics 4:560; Landegren (1988) Science 241:1077; Barringer (1990) Gene 89:117); transcription amplification (see, e.g., Kwoh (1989) Proc. Natl. Acad. Sci. USA 86:1173); and, self-sustained sequence replication (see, e.g., Guatelli (1990) Proc. Natl. Acad. Sci. USA 87:1874); Q Beta replicase amplification (see, e.g., Smith (1997) J. Clin. Microbiol. 35:1477-1491), automated Q-beta replicase amplification assay (see, e.g., Burg (1996) Mol. Cell. Probes 10:257-271) and other RNA polymerase mediated techniques (e.g., NASBA, Cangene, Mississauga, Ontario); see also Berger (1987) Methods Enzymol. 152:307-316; Sambrook; Ausubel; U.S. Patent Nos. 4,683,195 and 4,683,202; Sooknanan (1995) Biotechnology 13:563-564.

Determining the degree of sequence identity

25 The invention provides nucleic acids comprising sequences having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an
30 exemplary nucleic acid of the invention over a region of at least about 50, 75, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, 1050, 1100, 1150, 1200, 1250, 1300, 1350, 1400, 1450, 1500, 1550 or more, residues.

The invention provides polypeptides comprising sequences having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity to an exemplary polypeptide of the invention. The extent of sequence identity (homology) may be determined using any computer program and associated parameters, including those described herein, such as BLAST 2.2.2. or FASTA version 3.0t78, with the default parameters.

Figure 35 is a chart describing selected characteristics of exemplary nucleic acids and polypeptides of the invention, including sequence identity comparison of the exemplary sequences to public databases. All sequences described in Figure 35 have been subject to a BLAST search (as described in detail, below) against two sets of databases. The first database set is available through NCBI (National Center for Biotechnology Information). All results from searches against these databases are found in the columns entitled "NR Description", "NR Accession Code", "NR Evalue" or "NR Organism". "NR" refers to the Non-Redundant nucleotide database maintained by NCBI. This database is a composite of GenBank, GenBank updates, and EMBL updates. The entries in the column "NR Description" refer to the definition line in any given NCBI record, which includes a description of the sequence, such as the source organism, gene name/protein name, or some description of the function of the sequence. The entries in the column "NR Accession Code" refer to the unique identifier given to a sequence record. The entries in the column "NR Evalue" refer to the Expect value (Evalue), which represents the probability that an alignment score as good as the one found between the query sequence (the sequences of the invention) and a database sequence would be found in the same number of comparisons between random sequences as was done in the present BLAST search. The entries in the column "NR Organism" refer to the source organism of the sequence identified as the closest BLAST hit. The second set of databases is collectively known as the GeneseqTM database, which is available through Thomson Derwent (Philadelphia, PA). All results from searches against this database are found in the columns entitled "Geneseq Protein Description", "Geneseq Protein Accession Code", "Geneseq Protein Evalue", "Geneseq DNA Description", "Geneseq DNA Accession Code" or "Geneseq DNA Evalue". The information found in these

columns is comparable to the information found in the NR columns described above, except that it was derived from BLAST searches against the Geneseq™ database instead of the NCBI databases. In addition, this table includes the column “Predicted EC No.”. An EC number is the number assigned to a type of enzyme according to a scheme of standardized enzyme nomenclature developed by the Enzyme Commission of the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology (IUBMB). The results in the “Predicted EC No.” column are determined by a BLAST search against the Kegg (Kyoto Encyclopedia of Genes and Genomes) database. If the top BLAST match has an Evalue equal to or less than e^{-6} , the EC number assigned to the top match is entered into the table. The EC number of the top hit is used as a guide to what the EC number of the sequence of the invention might be. The columns “Query DNA Length” and “Query Protein Length” refer to the number of nucleotides or the number amino acids, respectively, in the sequence of the invention that was searched or queried against either the NCBI or Geneseq databases. The columns “Geneseq or NR DNA Length” and “Geneseq or NR Protein Length” refer to the number of nucleotides or the number amino acids, respectively, in the sequence of the top match from the BLAST search. The results provided in these columns are from the search that returned the lower Evalue, either from the NCBI databases or the Geneseq database. The columns “Geneseq or NR %ID Protein” and “Geneseq or NR %ID DNA” refer to the percent sequence identity between the sequence of the invention and the sequence of the top BLAST match. The results provided in these columns are from the search that returned the lower Evalue, either from the NCBI databases or the Geneseq database.

Homologous sequences also include RNA sequences in which uridines replace the thymines in the nucleic acid sequences. The homologous sequences may be obtained using any of the procedures described herein or may result from the correction of a sequencing error. It will be appreciated that the nucleic acid sequences as set forth herein can be represented in the traditional single character format (see, e.g., Stryer, Lubert. Biochemistry, 3rd Ed., W. H Freeman & Co., New York) or in any other format which records the identity of the nucleotides in a sequence.

Various sequence comparison programs identified herein are used in this aspect of the invention. Protein and/or nucleic acid sequence identities (homologies) may be evaluated using any of the variety of sequence comparison algorithms and programs known in the art. Such algorithms and programs include, but are not limited to,

TBLASTN, BLASTP, FASTA, TFASTA, and CLUSTALW (Pearson and Lipman, Proc. Natl. Acad. Sci. USA 85(8):2444-2448, 1988; Altschul et al., J. Mol. Biol. 215(3):403-410, 1990; Thompson et al., Nucleic Acids Res. 22(2):4673-4680, 1994; Higgins et al., Methods Enzymol. 266:383-402, 1996; Altschul et al., J. Mol. Biol. 215(3):403-410, 1990; Altschul et al., Nature Genetics 3:266-272, 1993).

Homology or sequence identity can be measured using sequence analysis software (e.g., Sequence Analysis Software Package of the Genetics Computer Group, University of Wisconsin Biotechnology Center, 1710 University Avenue, Madison, WI 53705). Such software matches similar sequences by assigning degrees of homology to various deletions, substitutions and other modifications. The terms "homology" and "identity" in the context of two or more nucleic acids or polypeptide sequences, refer to two or more sequences or subsequences that are the same or have a specified percentage of amino acid residues or nucleotides that are the same when compared and aligned for maximum correspondence over a comparison window or designated region as measured using any number of sequence comparison algorithms or by manual alignment and visual inspection. For sequence comparison, one sequence can act as a reference sequence, e.g., a sequence of the invention, to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are entered into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. Default program parameters can be used, or alternative parameters can be designated. The sequence comparison algorithm then calculates the percent sequence identities for the test sequences relative to the reference sequence, based on the program parameters.

A "comparison window", as used herein, includes reference to a segment of any one of the numbers of contiguous residues. For example, in alternative aspects of the invention, contiguous residues ranging anywhere from 20 to the full length of an exemplary polypeptide or nucleic acid sequence of the invention are compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned. If the reference sequence has the requisite sequence identity to an exemplary polypeptide or nucleic acid sequence of the invention, e.g., 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%,

98%, 99%, or more sequence identity to a sequence of the invention, that sequence is within the scope of the invention. In alternative embodiments, subsequences ranging from about 20 to 600, about 50 to 200, and about 100 to 150 are compared to a reference sequence of the same number of contiguous positions after the two sequences are
 5 optimally aligned.

Methods of alignment of sequence for comparison are well known in the art. In alternative aspects, optimal alignment of sequences for comparison can be conducted, e.g., by the local homology algorithm of Smith & Waterman, Adv. Appl. Math. 2:482, 1981, by the homology alignment algorithm of Needleman & Wunsch, J.
 10 Mol. Biol. 48:443, 1970, by the search for similarity method of person & Lipman, Proc. Nat'l. Acad. Sci. USA 85:2444, 1988, by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI), or by manual alignment and visual inspection. Other algorithms for determining homology or identity
 15 include, for example, in addition to a BLAST program (Basic Local Alignment Search Tool at the National Center for Biological Information), ALIGN, AMAS (Analysis of Multiply Aligned Sequences), AMPS (Protein Multiple Sequence Alignment), ASSET (Aligned Segment Statistical Evaluation Tool), BANDS, BESTSCOR, BIOSCAN (Biological Sequence Comparative Analysis Node), BLIMPS (BLOCKS IMPROVED
 20 Searcher), FASTA, Intervals & Points, BMB, CLUSTAL V, CLUSTAL W, CONSENSUS, LCONSENSUS, WCONSENSUS, Smith-Waterman algorithm, DARWIN, Las Vegas algorithm, FNAT (Forced Nucleotide Alignment Tool), Framealign, Framesearch, DYNAMIC, FILTER, FSAP (Fristensky Sequence Analysis Package), GAP (Global Alignment Program), GENAL, GIBBS, GenQuest, ISSC
 25 (Sensitive Sequence Comparison), LALIGN (Local Sequence Alignment), LCP (Local Content Program), MACAW (Multiple Alignment Construction & Analysis Workbench), MAP (Multiple Alignment Program), MBLKP, MBLKN, PIMA (Pattern-Induced Multi-sequence Alignment), SAGA (Sequence Alignment by Genetic Algorithm) and WHAT-IF. Such alignment programs can also be used to screen genome databases to identify
 30 polynucleotide sequences having substantially identical sequences. A number of genome databases are available, for example, a substantial portion of the human genome is available as part of the Human Genome Sequencing Project (Gibbs, 1995). Several genomes have been sequenced, e.g., *M. genitalium* (Fraser et al., 1995), *M. jannaschii*

(Bult et al., 1996), *H. influenzae* (Fleischmann et al., 1995), *E. coli* (Blattner et al., 1997), and yeast (*S. cerevisiae*) (Mewes et al., 1997), and *D. melanogaster* (Adams et al., 2000). Significant progress has also been made in sequencing the genomes of model organism, such as mouse, *C. elegans*, and *Arabidopsis* sp. Databases containing genomic information annotated with some functional information are maintained by different organization, and are accessible via the internet.

BLAST, BLAST 2.0 and BLAST 2.2.2 algorithms also can be used to practice the invention. They are described, e.g., in Altschul (1977) Nuc. Acids Res. 25:3389-3402; Altschul (1990) J. Mol. Biol. 215:403-410. Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information. This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul (1990) supra). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always >0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W , T , and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, an expectation (E) of 10, $M=5$, $N=-4$ and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength of 3, and expectations (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff & Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915) alignments (B) of 50, expectation (E) of 10, $M=5$, $N=-4$, and a comparison of both strands. The BLAST algorithm also performs a statistical analysis of the similarity between two sequences (see, e.g., Karlin & Altschul (1993) Proc. Natl. Acad. Sci. USA 90:5873). One measure of similarity provided by BLAST algorithm is the smallest sum

probability ($P(N)$), which provides an indication of the probability by which a match between two nucleotide or amino acid sequences would occur by chance. For example, a nucleic acid is considered similar to a references sequence if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.2, more preferably less than about 0.01, and most preferably less than about 0.001. In one aspect, protein and nucleic acid sequence homologies are evaluated using the Basic Local Alignment Search Tool ("BLAST"). For example, five specific BLAST programs can be used to perform the following task: (1) BLASTP and BLAST3 compare an amino acid query sequence against a protein sequence database; (2) BLASTN compares a nucleotide query sequence against a nucleotide sequence database; (3) BLASTX compares the six-frame conceptual translation products of a query nucleotide sequence (both strands) against a protein sequence database; (4) TBLASTN compares a query protein sequence against a nucleotide sequence database translated in all six reading frames (both strands); and, (5) TBLASTX compares the six-frame translations of a nucleotide query sequence against the six-frame translations of a nucleotide sequence database. The BLAST programs identify homologous sequences by identifying similar segments, which are referred to herein as "high-scoring segment pairs," between a query amino or nucleic acid sequence and a test sequence which is preferably obtained from a protein or nucleic acid sequence database. High-scoring segment pairs are preferably identified (i.e., aligned) by means of a scoring matrix, many of which are known in the art. Preferably, the scoring matrix used is the BLOSUM62 matrix (Gonnet et al., Science 256:1443-1445, 1992; Henikoff and Henikoff, Proteins 17:49-61, 1993). Less preferably, the PAM or PAM250 matrices may also be used (see, e.g., Schwartz and Dayhoff, eds., 1978, Matrices for Detecting Distance Relationships: Atlas of Protein Sequence and Structure, Washington: National Biomedical Research Foundation).

In one aspect of the invention, to determine if a nucleic acid has the requisite sequence identity to be within the scope of the invention, the NCBI BLAST 2.2.2 programs is used, default options to blastp. There are about 38 setting options in the BLAST 2.2.2 program. In this exemplary aspect of the invention, all default values are used except for the default filtering setting (i.e., all parameters set to default except filtering which is set to OFF); in its place a "-F F" setting is used, which disables filtering. Use of default filtering often results in Karlin-Altschul violations due to short length of sequence.

The default values used in this exemplary aspect of the invention, and to determine the values in Figure 35, as discussed above, include:

"Filter for low complexity: ON

Word Size: 3

Matrix: Blosom62

Gap Costs: Existence:11

Extension:1"

Other default settings can be: filter for low complexity OFF, word size of 3 for protein, BLOSUM62 matrix, gap existence penalty of -11 and a gap extension penalty of -1. An exemplary NCBI BLAST 2.2.2 program setting has the "-W" option default to 0. This means that, if not set, the word size defaults to 3 for proteins and 11 for nucleotides.

Computer systems and computer program products

- To determine and identify sequence identities, structural homologies, motifs and the like in silico, the sequence of the invention can be stored, recorded, and manipulated on any medium which can be read and accessed by a computer. Accordingly, the invention provides computers, computer systems, computer readable mediums, computer programs products and the like recorded or stored thereon the nucleic acid and polypeptide sequences of the invention. As used herein, the words "recorded" and "stored" refer to a process for storing information on a computer medium. A skilled artisan can readily adopt any known methods for recording information on a computer readable medium to generate manufactures comprising one or more of the nucleic acid and/or polypeptide sequences of the invention.

- Another aspect of the invention is a computer readable medium having recorded thereon at least one nucleic acid and/or polypeptide sequence of the invention. Computer readable media include magnetically readable media, optically readable media, electronically readable media and magnetic/optical media. For example, the computer readable media may be a hard disk, a floppy disk, a magnetic tape, CD-ROM, Digital Versatile Disk (DVD), Random Access Memory (RAM), or Read Only Memory (ROM) as well as other types of other media known to those skilled in the art.

Aspects of the invention include systems (e.g., internet based systems), particularly computer systems, which store and manipulate the sequences and sequence

information described herein. One example of a computer system 100 is illustrated in block diagram form in Figure 1. As used herein, "a computer system" refers to the hardware components, software components, and data storage components used to analyze a nucleotide or polypeptide sequence of the invention. The computer system 100
5 can include a processor for processing, accessing and manipulating the sequence data. The processor 105 can be any well-known type of central processing unit, such as, for example, the Pentium III from Intel Corporation, or similar processor from Sun, Motorola, Compaq, AMD or International Business Machines. The computer system 100 is a general purpose system that comprises the processor 105 and one or more internal
10 data storage components 110 for storing data, and one or more data retrieving devices for retrieving the data stored on the data storage components. A skilled artisan can readily appreciate that any one of the currently available computer systems are suitable.

In one aspect, the computer system 100 includes a processor 105 connected to a bus which is connected to a main memory 115 (preferably implemented as
15 RAM) and one or more internal data storage devices 110, such as a hard drive and/or other computer readable media having data recorded thereon. The computer system 100 can further include one or more data retrieving device 118 for reading the data stored on the internal data storage devices 110. The data retrieving device 118 may represent, for example, a floppy disk drive, a compact disk drive, a magnetic tape drive, or a modem
20 capable of connection to a remote data storage system (e.g., via the internet) etc. In some embodiments, the internal data storage device 110 is a removable computer readable medium such as a floppy disk, a compact disk, a magnetic tape, etc. containing control logic and/or data recorded thereon. The computer system 100 may advantageously include or be programmed by appropriate software for reading the control logic and/or the
25 data from the data storage component once inserted in the data retrieving device. The computer system 100 includes a display 120 which is used to display output to a computer user. It should also be noted that the computer system 100 can be linked to other computer systems 125a-c in a network or wide area network to provide centralized access to the computer system 100. Software for accessing and processing the nucleotide
30 or amino acid sequences of the invention can reside in main memory 115 during execution. In some aspects, the computer system 100 may further comprise a sequence comparison algorithm for comparing a nucleic acid sequence of the invention. The algorithm and sequence(s) can be stored on a computer readable medium. A "sequence

comparison algorithm” refers to one or more programs which are implemented (locally or remotely) on the computer system 100 to compare a nucleotide sequence with other nucleotide sequences and/or compounds stored within a data storage means. For example, the sequence comparison algorithm may compare the nucleotide sequences of the invention stored on a computer readable medium to reference sequences stored on a computer readable medium to identify homologies or structural motifs.

The parameters used with the above algorithms may be adapted depending on the sequence length and degree of homology studied. In some aspects, the parameters may be the default parameters used by the algorithms in the absence of instructions from the user. Figure 2 is a flow diagram illustrating one aspect of a process 200 for comparing a new nucleotide or protein sequence with a database of sequences in order to determine the homology levels between the new sequence and the sequences in the database. The database of sequences can be a private database stored within the computer system 100, or a public database such as GENBANK that is available through the Internet. The process 200 begins at a start state 201 and then moves to a state 202 wherein the new sequence to be compared is stored to a memory in a computer system 100. As discussed above, the memory could be any type of memory, including RAM or an internal storage device. The process 200 then moves to a state 204 wherein a database of sequences is opened for analysis and comparison. The process 200 then moves to a state 206 wherein the first sequence stored in the database is read into a memory on the computer. A comparison is then performed at a state 210 to determine if the first sequence is the same as the second sequence. It is important to note that this step is not limited to performing an exact comparison between the new sequence and the first sequence in the database. Well-known methods are known to those of skill in the art for comparing two nucleotide or protein sequences, even if they are not identical. For example, gaps can be introduced into one sequence in order to raise the homology level between the two tested sequences. The parameters that control whether gaps or other features are introduced into a sequence during comparison are normally entered by the user of the computer system. Once a comparison of the two sequences has been performed at the state 210, a determination is made at a decision state 210 whether the two sequences are the same. Of course, the term “same” is not limited to sequences that are absolutely identical. Sequences that are within the homology parameters entered by the user will be marked as “same” in the process 200. If a determination is made that the

two sequences are the same, the process 200 moves to a state 214 wherein the name of the sequence from the database is displayed to the user. This state notifies the user that the sequence with the displayed name fulfills the homology constraints that were entered. Once the name of the stored sequence is displayed to the user, the process 200 moves to a decision state 218 wherein a determination is made whether more sequences exist in the database. If no more sequences exist in the database, then the process 200 terminates at an end state 220. However, if more sequences do exist in the database, then the process 200 moves to a state 224 wherein a pointer is moved to the next sequence in the database so that it can be compared to the new sequence. In this manner, the new sequence is aligned and compared with every sequence in the database. It should be noted that if a determination had been made at the decision state 212 that the sequences were not homologous, then the process 200 would move immediately to the decision state 218 in order to determine if any other sequences were available in the database for comparison. Accordingly, one aspect of the invention is a computer system comprising a processor, a data storage device having stored thereon a nucleic acid sequence of the invention and a sequence comparer for conducting the comparison. The sequence comparer may indicate a homology level between the sequences compared or identify structural motifs, or it may identify structural motifs in sequences which are compared to these nucleic acid codes and polypeptide codes. Figure 3 is a flow diagram illustrating one embodiment of a process 250 in a computer for determining whether two sequences are homologous. The process 250 begins at a start state 252 and then moves to a state 254 wherein a first sequence to be compared is stored to a memory. The second sequence to be compared is then stored to a memory at a state 256. The process 250 then moves to a state 260 wherein the first character in the first sequence is read and then to a state 262 wherein the first character of the second sequence is read. It should be understood that if the sequence is a nucleotide sequence, then the character would normally be either A, T, C, G or U. If the sequence is a protein sequence, then it can be a single letter amino acid code so that the first and sequence sequences can be easily compared. A determination is then made at a decision state 264 whether the two characters are the same. If they are the same, then the process 250 moves to a state 268 wherein the next characters in the first and second sequences are read. A determination is then made whether the next characters are the same. If they are, then the process 250 continues this loop until two characters are not the same. If a determination is made that the next two characters are not the same, the

process 250 moves to a decision state 274 to determine whether there are any more characters either sequence to read. If there are not any more characters to read, then the process 250 moves to a state 276 wherein the level of homology between the first and second sequences is displayed to the user. The level of homology is determined by calculating the proportion of characters between the sequences that were the same out of the total number of sequences in the first sequence. Thus, if every character in a first 100 nucleotide sequence aligned with an every character in a second sequence, the homology level would be 100%.

Alternatively, the computer program can compare a reference sequence to a sequence of the invention to determine whether the sequences differ at one or more positions. The program can record the length and identity of inserted, deleted or substituted nucleotides or amino acid residues with respect to the sequence of either the reference or the invention. The computer program may be a program which determines whether a reference sequence contains a single nucleotide polymorphism (SNP) with respect to a sequence of the invention, or, whether a sequence of the invention comprises a SNP of a known sequence. Thus, in some aspects, the computer program is a program which identifies SNPs. The method may be implemented by the computer systems described above and the method illustrated in Figure 3. The method can be performed by reading a sequence of the invention and the reference sequences through the use of the computer program and identifying differences with the computer program.

In other aspects the computer based system comprises an identifier for identifying features within a nucleic acid or polypeptide of the invention. An "identifier" refers to one or more programs which identifies certain features within a nucleic acid sequence. For example, an identifier may comprise a program which identifies an open reading frame (ORF) in a nucleic acid sequence. Figure 4 is a flow diagram illustrating one aspect of an identifier process 300 for detecting the presence of a feature in a sequence. The process 300 begins at a start state 302 and then moves to a state 304 wherein a first sequence that is to be checked for features is stored to a memory 115 in the computer system 100. The process 300 then moves to a state 306 wherein a database of sequence features is opened. Such a database would include a list of each feature's attributes along with the name of the feature. For example, a feature name could be "Initiation Codon" and the attribute would be "ATG". Another example would be the feature name "TAATAA Box" and the feature attribute would be "TAATAA". An

example of such a database is produced by the University of Wisconsin Genetics Computer Group. Alternatively, the features may be structural polypeptide motifs such as alpha helices, beta sheets, or functional polypeptide motifs such as enzymatic active sites, helix-turn-helix motifs or other motifs known to those skilled in the art. Once the database of features is opened at the state 306, the process 300 moves to a state 308 wherein the first feature is read from the database. A comparison of the attribute of the first feature with the first sequence is then made at a state 310. A determination is then made at a decision state 316 whether the attribute of the feature was found in the first sequence. If the attribute was found, then the process 300 moves to a state 318 wherein the name of the found feature is displayed to the user. The process 300 then moves to a decision state 320 wherein a determination is made whether more features exist in the database. If no more features do exist, then the process 300 terminates at an end state 324. However, if more features do exist in the database, then the process 300 reads the next sequence feature at a state 326 and loops back to the state 310 wherein the attribute of the next feature is compared against the first sequence. If the feature attribute is not found in the first sequence at the decision state 316, the process 300 moves directly to the decision state 320 in order to determine if any more features exist in the database. Thus, in one aspect, the invention provides a computer program that identifies open reading frames (ORFs).

A polypeptide or nucleic acid sequence of the invention can be stored and manipulated in a variety of data processor programs in a variety of formats. For example, a sequence can be stored as text in a word processing file, such as MicrosoftWORD or WORDPERFECT or as an ASCII file in a variety of database programs familiar to those of skill in the art, such as DB2, SYBASE, or ORACLE. In addition, many computer programs and databases may be used as sequence comparison algorithms, identifiers, or sources of reference nucleotide sequences or polypeptide sequences to be compared to a nucleic acid sequence of the invention. The programs and databases used to practice the invention include, but are not limited to: MacPattern (EMBL), DiscoveryBase (Molecular Applications Group), GeneMine (Molecular Applications Group), Look (Molecular Applications Group), MacLook (Molecular Applications Group), BLAST and BLAST2 (NCBI), BLASTN and BLASTX (Altschul et al, J. Mol. Biol. 215: 403, 1990), FASTA (Pearson and Lipman, Proc. Natl. Acad. Sci. USA, 85: 2444, 1988), FASTDB (Brutlag et al. Comp. App. Biosci. 6:237-245, 1990), Catalyst (Molecular Simulations Inc.),

Catalyst/SHAPE (Molecular Simulations Inc.), Cerius2.DBAccess (Molecular Simulations Inc.), HypoGen (Molecular Simulations Inc.), Insight II, (Molecular Simulations Inc.), Discover (Molecular Simulations Inc.), CHARMM (Molecular Simulations Inc.), Felix (Molecular Simulations Inc.), DelPhi, (Molecular Simulations Inc.), QuanteMM, (Molecular Simulations Inc.), Homology (Molecular Simulations Inc.), Modeler (Molecular Simulations Inc.), ISIS (Molecular Simulations Inc.), Quanta/Protein Design (Molecular Simulations Inc.), WebLab (Molecular Simulations Inc.), WebLab Diversity Explorer (Molecular Simulations Inc.), Gene Explorer (Molecular Simulations Inc.), SeqFold (Molecular Simulations Inc.), the MDL Available Chemicals Directory database, the MDL Drug Data Report data base, the Comprehensive Medicinal Chemistry database, Derwent's World Drug Index database, the BioByteMasterFile database, the Genbank database, and the Genseqn database. Many other programs and data bases would be apparent to one of skill in the art given the present disclosure.

Motifs which may be detected using the above programs include sequences encoding leucine zippers, helix-turn-helix motifs, glycosylation sites, ubiquitination sites, alpha helices, and beta sheets, signal sequences encoding signal peptides which direct the secretion of the encoded proteins, sequences implicated in transcription regulation such as homeoboxes, acidic stretches, enzymatic active sites, substrate binding sites, and enzymatic cleavage sites.

Hybridization of nucleic acids

The invention provides isolated or recombinant nucleic acids that hybridize under stringent conditions to an exemplary sequence of the invention, or a nucleic acid that encodes a polypeptide of the invention. The stringent conditions can be highly stringent conditions, medium stringent conditions, low stringent conditions, including the high and reduced stringency conditions described herein. In one aspect, it is the stringency of the wash conditions that set forth the conditions which determine whether a nucleic acid is within the scope of the invention, as discussed below.

In alternative embodiments, nucleic acids of the invention as defined by their ability to hybridize under stringent conditions can be between about five residues and the full length of nucleic acid of the invention; e.g., they can be at least 5, 10, 15, 20, 25, 30, 35, 40, 50, 55, 60, 65, 70, 75, 80, 90, 100, 150, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, or more, residues in length. Nucleic

acids shorter than full length are also included. These nucleic acids can be useful as, e.g., hybridization probes, labeling probes, PCR oligonucleotide probes, iRNA, antisense or sequences encoding antibody binding peptides (epitopes), motifs, active sites and the like.

In one aspect, nucleic acids of the invention are defined by their ability to
5 hybridize under high stringency comprising conditions of about 50% formamide at about 37°C to 42°C. In one aspect, nucleic acids of the invention are defined by their ability to hybridize under reduced stringency comprising conditions in about 35% to 25% formamide at about 30°C to 35°C.

Alternatively, nucleic acids of the invention are defined by their ability to
10 hybridize under high stringency comprising conditions at 42°C in 50% formamide, 5X SSPE, 0.3% SDS, and a repetitive sequence blocking nucleic acid, such as cot-1 or salmon sperm DNA (e.g., 200 n/ml sheared and denatured salmon sperm DNA). In one aspect, nucleic acids of the invention are defined by their ability to hybridize under reduced stringency conditions comprising 35% formamide at a reduced temperature of
15 35°C.

Following hybridization, the filter may be washed with 6X SSC, 0.5% SDS at 50°C. These conditions are considered to be "moderate" conditions above 25% formamide and "low" conditions below 25% formamide. A specific example of
"moderate" hybridization conditions is when the above hybridization is conducted at 30%
20 formamide. A specific example of "low stringency" hybridization conditions is when the above hybridization is conducted at 10% formamide.

The temperature range corresponding to a particular level of stringency can be further narrowed by calculating the purine to pyrimidine ratio of the nucleic acid of interest and adjusting the temperature accordingly. Nucleic acids of the invention are
25 also defined by their ability to hybridize under high, medium, and low stringency conditions as set forth in Ausubel and Sambrook. Variations on the above ranges and conditions are well known in the art. Hybridization conditions are discussed further, below.

The above procedure may be modified to identify nucleic acids having
30 decreasing levels of homology to the probe sequence. For example, to obtain nucleic acids of decreasing homology to the detectable probe, less stringent conditions may be used. For example, the hybridization temperature may be decreased in increments of 5°C from 68°C to 42°C in a hybridization buffer having a Na⁺ concentration of approximately

1M. Following hybridization, the filter may be washed with 2X SSC, 0.5% SDS at the temperature of hybridization. These conditions are considered to be "moderate" conditions above 50°C and "low" conditions below 50°C. A specific example of "moderate" hybridization conditions is when the above hybridization is conducted at 55°C. A specific example of "low stringency" hybridization conditions is when the above hybridization is conducted at 45°C.

Alternatively, the hybridization may be carried out in buffers, such as 6X SSC, containing formamide at a temperature of 42°C. In this case, the concentration of formamide in the hybridization buffer may be reduced in 5% increments from 50% to 0% to identify clones having decreasing levels of homology to the probe. Following hybridization, the filter may be washed with 6X SSC, 0.5% SDS at 50°C. These conditions are considered to be "moderate" conditions above 25% formamide and "low" conditions below 25% formamide. A specific example of "moderate" hybridization conditions is when the above hybridization is conducted at 30% formamide. A specific example of "low stringency" hybridization conditions is when the above hybridization is conducted at 10% formamide.

However, the selection of a hybridization format is not critical - it is the stringency of the wash conditions that set forth the conditions which determine whether a nucleic acid is within the scope of the invention. Wash conditions used to identify nucleic acids within the scope of the invention include, e.g.: a salt concentration of about 0.02 molar at pH 7 and a temperature of at least about 50°C or about 55°C to about 60°C; or, a salt concentration of about 0.15 M NaCl at 72°C for about 15 minutes; or, a salt concentration of about 0.2X SSC at a temperature of at least about 50°C or about 55°C to about 60°C for about 15 to about 20 minutes; or, the hybridization complex is washed twice with a solution with a salt concentration of about 2X SSC containing 0.1% SDS at room temperature for 15 minutes and then washed twice by 0.1X SSC containing 0.1% SDS at 68°C for 15 minutes; or, equivalent conditions. See Sambrook, Tijssen and Ausubel for a description of SSC buffer and equivalent conditions.

These methods may be used to isolate nucleic acids of the invention.

Oligonucleotides probes and methods for using them

The invention also provides nucleic acid probes that can be used, e.g., for identifying nucleic acids encoding a polypeptide with an amylase activity or fragments thereof or for identifying amylase genes. In one aspect, the probe comprises at least 10 consecutive bases of a nucleic acid of the invention. Alternatively, a probe of the invention can be at least about 5, 6, 7, 8, 9, 10, 15, 20, 25, 30, 35, 40, 45, 50, 60, 70, 80, 90, 100, 110, 120, 130, 150 or about 10 to 50, about 20 to 60 about 30 to 70, consecutive bases of a sequence as set forth in a nucleic acid of the invention. The probes identify a nucleic acid by binding and/or hybridization. The probes can be used in arrays of the invention, see discussion below, including, e.g., capillary arrays. The probes of the invention can also be used to isolate other nucleic acids or polypeptides.

The probes of the invention can be used to determine whether a biological sample, such as a soil sample, contains an organism having a nucleic acid sequence of the invention or an organism from which the nucleic acid was obtained. In such procedures, a biological sample potentially harboring the organism from which the nucleic acid was isolated is obtained and nucleic acids are obtained from the sample. The nucleic acids are contacted with the probe under conditions which permit the probe to specifically hybridize to any complementary sequences present in the sample. Where necessary, conditions which permit the probe to specifically hybridize to complementary sequences may be determined by placing the probe in contact with complementary sequences from samples known to contain the complementary sequence, as well as control sequences which do not contain the complementary sequence. Hybridization conditions, such as the salt concentration of the hybridization buffer, the formamide concentration of the hybridization buffer, or the hybridization temperature, may be varied to identify conditions which allow the probe to hybridize specifically to complementary nucleic acids (see discussion on specific hybridization conditions).

If the sample contains the organism from which the nucleic acid was isolated, specific hybridization of the probe is then detected. Hybridization may be detected by labeling the probe with a detectable agent such as a radioactive isotope, a fluorescent dye or an enzyme capable of catalyzing the formation of a detectable product. Many methods for using the labeled probes to detect the presence of complementary nucleic acids in a sample are familiar to those skilled in the art. These include Southern

Blots, Northern Blots, colony hybridization procedures, and dot blots. Protocols for each of these procedures are provided in Ausubel and Sambrook.

Alternatively, more than one probe (at least one of which is capable of specifically hybridizing to any complementary sequences which are present in the nucleic acid sample), may be used in an amplification reaction to determine whether the sample contains an organism containing a nucleic acid sequence of the invention (e.g., an organism from which the nucleic acid was isolated). In one aspect, the probes comprise oligonucleotides. In one aspect, the amplification reaction may comprise a PCR reaction. PCR protocols are described in Ausubel and Sambrook (see discussion on amplification reactions). In such procedures, the nucleic acids in the sample are contacted with the probes, the amplification reaction is performed, and any resulting amplification product is detected. The amplification product may be detected by performing gel electrophoresis on the reaction products and staining the gel with an intercalator such as ethidium bromide. Alternatively, one or more of the probes may be labeled with a radioactive isotope and the presence of a radioactive amplification product may be detected by autoradiography after gel electrophoresis.

Probes derived from sequences near the 3' or 5' ends of a nucleic acid sequence of the invention can also be used in chromosome walking procedures to identify clones containing additional, e.g., genomic sequences. Such methods allow the isolation of genes which encode additional proteins of interest from the host organism.

In one aspect, nucleic acid sequences of the invention are used as probes to identify and isolate related nucleic acids. In some aspects, the so-identified related nucleic acids may be cDNAs or genomic DNAs from organisms other than the one from which the nucleic acid of the invention was first isolated. In such procedures, a nucleic acid sample is contacted with the probe under conditions which permit the probe to specifically hybridize to related sequences. Hybridization of the probe to nucleic acids from the related organism is then detected using any of the methods described above.

In nucleic acid hybridization reactions, the conditions used to achieve a particular level of stringency can vary, depending on the nature of the nucleic acids being hybridized. For example, the length, degree of complementarity, nucleotide sequence composition (e.g., GC v. AT content), and nucleic acid type (e.g., RNA v. DNA) of the hybridizing regions of the nucleic acids can be considered in selecting hybridization conditions. An additional consideration is whether one of the nucleic acids is

immobilized, for example, on a filter. Hybridization can be carried out under conditions of low stringency, moderate stringency or high stringency. As an example of nucleic acid hybridization, a polymer membrane containing immobilized denatured nucleic acids is first prehybridized for 30 minutes at 45°C in a solution consisting of 0.9 M NaCl, 50 mM NaH₂PO₄, pH 7.0, 5.0 mM Na₂EDTA, 0.5% SDS, 10X Denhardt's, and 0.5 mg/ml polyriboadenylic acid. Approximately 2×10^7 cpm (specific activity $4-9 \times 10^8$ cpm/ug) of ³²P end-labeled oligonucleotide probe can then added to the solution. After 12-16 hours of incubation, the membrane is washed for 30 minutes at room temperature (RT) in 1X SET (150 mM NaCl, 20 mM Tris hydrochloride, pH 7.8, 1 mM Na₂EDTA) containing 0.5% SDS, followed by a 30 minute wash in fresh 1X SET at T_m-10°C for the oligonucleotide probe. The membrane is then exposed to auto-radiographic film for detection of hybridization signals.

By varying the stringency of the hybridization conditions used to identify nucleic acids, such as cDNAs or genomic DNAs, which hybridize to the detectable probe, nucleic acids having different levels of homology to the probe can be identified and isolated. Stringency may be varied by conducting the hybridization at varying temperatures below the melting temperatures of the probes. The melting temperature, T_m, is the temperature (under defined ionic strength and pH) at which 50% of the target sequence hybridizes to a perfectly complementary probe. Very stringent conditions are selected to be equal to or about 5°C lower than the T_m for a particular probe. The melting temperature of the probe may be calculated using the following exemplary formulas. For probes between 14 and 70 nucleotides in length the melting temperature (T_m) is calculated using the formula: $T_m = 81.5 + 16.6(\log [Na^+]) + 0.41(\text{fraction G+C}) - (600/N)$ where N is the length of the probe. If the hybridization is carried out in a solution containing formamide, the melting temperature may be calculated using the equation: $T_m = 81.5 + 16.6(\log [Na^+]) + 0.41(\text{fraction G+C}) - (0.63\% \text{ formamide}) - (600/N)$ where N is the length of the probe. Prehybridization may be carried out in 6X SSC, 5X Denhardt's reagent, 0.5% SDS, 100µg denatured fragmented salmon sperm DNA or 6X SSC, 5X Denhardt's reagent, 0.5% SDS, 100µg denatured fragmented salmon sperm DNA, 50% formamide. Formulas for SSC and Denhardt's and other solutions are listed, e.g., in Sambrook.

Hybridization is conducted by adding the detectable probe to the prehybridization solutions listed above. Where the probe comprises double stranded

DNA, it is denatured before addition to the hybridization solution. The filter is contacted with the hybridization solution for a sufficient period of time to allow the probe to hybridize to cDNAs or genomic DNAs containing sequences complementary thereto or homologous thereto. For probes over 200 nucleotides in length, the hybridization may be carried out at 15-25°C below the T_m . For shorter probes, such as oligonucleotide probes, the hybridization may be conducted at 5-10°C below the T_m . In one aspect, hybridizations in 6X SSC are conducted at approximately 68°C. In one aspect, hybridizations in 50% formamide containing solutions are conducted at approximately 42°C. All of the foregoing hybridizations would be considered to be under conditions of high stringency.

Following hybridization, the filter is washed to remove any non-specifically bound detectable probe. The stringency used to wash the filters can also be varied depending on the nature of the nucleic acids being hybridized, the length of the nucleic acids being hybridized, the degree of complementarity, the nucleotide sequence composition (e.g., GC v. AT content), and the nucleic acid type (e.g., RNA v. DNA). Examples of progressively higher stringency condition washes are as follows: 2X SSC, 0.1% SDS at room temperature for 15 minutes (low stringency); 0.1X SSC, 0.5% SDS at room temperature for 30 minutes to 1 hour (moderate stringency); 0.1X SSC, 0.5% SDS for 15 to 30 minutes at between the hybridization temperature and 68°C (high stringency); and 0.15M NaCl for 15 minutes at 72°C (very high stringency). A final low stringency wash can be conducted in 0.1X SSC at room temperature. The examples above are merely illustrative of one set of conditions that can be used to wash filters. One of skill in the art would know that there are numerous recipes for different stringency washes.

Nucleic acids which have hybridized to the probe can be identified by autoradiography or other conventional techniques. The above procedure may be modified to identify nucleic acids having decreasing levels of homology to the probe sequence. For example, to obtain nucleic acids of decreasing homology to the detectable probe, less stringent conditions may be used. For example, the hybridization temperature may be decreased in increments of 5°C from 68°C to 42°C in a hybridization buffer having a Na^+ concentration of approximately 1M. Following hybridization, the filter may be washed with 2X SSC, 0.5% SDS at the temperature of hybridization. These conditions are considered to be "moderate" conditions above 50°C and "low" conditions below 50°C.

An example of "moderate" hybridization conditions is when the above hybridization is conducted at 55°C. An example of "low stringency" hybridization conditions is when the above hybridization is conducted at 45°C.

Alternatively, the hybridization may be carried out in buffers, such as 6X SSC, containing formamide at a temperature of 42°C. In this case, the concentration of formamide in the hybridization buffer may be reduced in 5% increments from 50% to 0% to identify clones having decreasing levels of homology to the probe. Following hybridization, the filter may be washed with 6X SSC, 0.5% SDS at 50°C. These conditions are considered to be "moderate" conditions above 25% formamide and "low" conditions below 25% formamide. A specific example of "moderate" hybridization conditions is when the above hybridization is conducted at 30% formamide. A specific example of "low stringency" hybridization conditions is when the above hybridization is conducted at 10% formamide.

These probes and methods of the invention can be used to isolate nucleic acids having a sequence with at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% sequence identity ("homology") to a nucleic acid sequence of the invention comprising at least about 10, 15, 20, 25, 30, 35, 40, 50, 75, 100, 150, 200, 250, 300, 350, 400, 500, 550, 600, 650, 700, 750, 800, 850, 900, 950, 1000, or more consecutive bases thereof, and the sequences complementary thereto. Homology may be measured using an alignment algorithm, as discussed herein. For example, the homologous polynucleotides may have a coding sequence which is a naturally occurring allelic variant of one of the coding sequences described herein. Such allelic variants may have a substitution, deletion or addition of one or more nucleotides when compared to a nucleic acid of the invention.

Additionally, the probes and methods of the invention can be used to isolate nucleic acids which encode polypeptides having at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99%, sequence identity (homology) to a polypeptide of the invention comprising at least 5, 10, 15, 20, 25, 30, 35, 40, 50, 75, 100, or 150 consecutive amino acids, as

determined using a sequence alignment algorithm (e.g., such as the FASTA version 3.0t78 algorithm with the default parameters, or a BLAST 2.2.2 program with exemplary settings as set forth herein).

Inhibiting Expression of Amylase

5 The invention provides nucleic acids complementary to (e.g., antisense sequences to) the nucleic acid sequences of the invention. Antisense sequences are capable of inhibiting the transport, splicing or transcription of amylase-encoding genes. The inhibition can be effected through the targeting of genomic DNA or messenger RNA. The transcription or function of targeted nucleic acid can be inhibited, for example, by
10 hybridization and/or cleavage. One particularly useful set of inhibitors provided by the present invention includes oligonucleotides which are able to either bind amylase gene or message, in either case preventing or inhibiting the production or function of amylase. The association can be through sequence specific hybridization. Another useful class of inhibitors includes oligonucleotides which cause inactivation or cleavage of amylase
15 message. The oligonucleotide can have enzyme activity which causes such cleavage, such as ribozymes. The oligonucleotide can be chemically modified or conjugated to an enzyme or composition capable of cleaving the complementary nucleic acid. A pool of many different such oligonucleotides can be screened for those with the desired activity.

Antisense Oligonucleotides

20 The invention provides antisense oligonucleotides capable of binding amylase message which can inhibit proteolytic activity by targeting mRNA. Strategies for designing antisense oligonucleotides are well described in the scientific and patent literature, and the skilled artisan can design such amylase oligonucleotides using the novel reagents of the invention. For example, gene walking/ RNA mapping protocols to
25 screen for effective antisense oligonucleotides are well known in the art, see, e.g., Ho (2000) Methods Enzymol. 314:168-183, describing an RNA mapping assay, which is based on standard molecular techniques to provide an easy and reliable method for potent antisense sequence selection. See also Smith (2000) Eur. J. Pharm. Sci. 11:191-198.

Naturally occurring nucleic acids are used as antisense oligonucleotides.

30 The antisense oligonucleotides can be of any length; for example, in alternative aspects, the antisense oligonucleotides are between about 5 to 100, about 10 to 80, about 15 to 60, about 18 to 40. The optimal length can be determined by routine screening. The

antisense oligonucleotides can be present at any concentration. The optimal concentration can be determined by routine screening. A wide variety of synthetic, non-naturally occurring nucleotide and nucleic acid analogues are known which can address this potential problem. For example, peptide nucleic acids (PNAs) containing non-ionic backbones, such as N-(2-aminoethyl) glycine units can be used. Antisense oligonucleotides having phosphorothioate linkages can also be used, as described in WO 97/03211; WO 96/39154; Mata (1997) Toxicol Appl Pharmacol 144:189-197; Antisense Therapeutics, ed. Agrawal (Humana Press, Totowa, N.J., 1996). Antisense oligonucleotides having synthetic DNA backbone analogues provided by the invention can also include phosphoro-dithioate, methylphosphonate, phosphoramidate, alkyl phosphotriester, sulfamate, 3'-thioacetal, methylene(methylimino), 3'-N-carbamate, and morpholino carbamate nucleic acids, as described above.

Combinatorial chemistry methodology can be used to create vast numbers of oligonucleotides that can be rapidly screened for specific oligonucleotides that have appropriate binding affinities and specificities toward any target, such as the sense and antisense amylase sequences of the invention (see, e.g., Gold (1995) J. of Biol. Chem. 270:13581-13584).

Inhibitory Ribozymes

The invention provides ribozymes capable of binding amylase message. These ribozymes can inhibit amylase activity by, e.g., targeting mRNA. Strategies for designing ribozymes and selecting the amylase-specific antisense sequence for targeting are well described in the scientific and patent literature, and the skilled artisan can design such ribozymes using the novel reagents of the invention. Ribozymes act by binding to a target RNA through the target RNA binding portion of a ribozyme which is held in close proximity to an enzymatic portion of the RNA that cleaves the target RNA. Thus, the ribozyme recognizes and binds a target RNA through complementary base-pairing, and once bound to the correct site, acts enzymatically to cleave and inactivate the target RNA. Cleavage of a target RNA in such a manner will destroy its ability to direct synthesis of an encoded protein if the cleavage occurs in the coding sequence. After a ribozyme has bound and cleaved its RNA target, it can be released from that RNA to bind and cleave new targets repeatedly.

In some circumstances, the enzymatic nature of a ribozyme can be advantageous over other technologies, such as antisense technology (where a nucleic acid molecule simply binds to a nucleic acid target to block its transcription, translation or association with another molecule) as the effective concentration of ribozyme necessary to effect a therapeutic treatment can be lower than that of an antisense oligonucleotide. This potential advantage reflects the ability of the ribozyme to act enzymatically. Thus, a single ribozyme molecule is able to cleave many molecules of target RNA. In addition, a ribozyme is typically a highly specific inhibitor, with the specificity of inhibition depending not only on the base pairing mechanism of binding, but also on the mechanism by which the molecule inhibits the expression of the RNA to which it binds. That is, the inhibition is caused by cleavage of the RNA target and so specificity is defined as the ratio of the rate of cleavage of the targeted RNA over the rate of cleavage of non-targeted RNA. This cleavage mechanism is dependent upon factors additional to those involved in base pairing. Thus, the specificity of action of a ribozyme can be greater than that of antisense oligonucleotide binding the same RNA site.

The ribozyme of the invention, e.g., an enzymatic ribozyme RNA molecule, can be formed in a hammerhead motif, a hairpin motif, as a hepatitis delta virus motif, a group I intron motif and/or an RNaseP-like RNA in association with an RNA guide sequence. Examples of hammerhead motifs are described by, e.g., Rossi (1992) *Aids Research and Human Retroviruses* 8:183; hairpin motifs by Hampel (1989) *Biochemistry* 28:4929, and Hampel (1990) *Nuc. Acids Res.* 18:299; the hepatitis delta virus motif by Perrotta (1992) *Biochemistry* 31:16; the RNaseP motif by Guerrier-Takada (1983) *Cell* 35:849; and the group I intron by Cech U.S. Pat. No. 4,987,071. The recitation of these specific motifs is not intended to be limiting. Those skilled in the art will recognize that a ribozyme of the invention, e.g., an enzymatic RNA molecule of this invention, can have a specific substrate binding site complementary to one or more of the target gene RNA regions. A ribozyme of the invention can have a nucleotide sequence within or surrounding that substrate binding site which imparts an RNA cleaving activity to the molecule.

RNA interference (RNAi)

In one aspect, the invention provides an RNA inhibitory molecule, a so-called "RNAi" molecule, comprising an amylase sequence of the invention. The RNAi

molecule comprises a double-stranded RNA (dsRNA) molecule. The RNAi can inhibit expression of an amylase gene. In one aspect, the RNAi is about 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25 or more duplex nucleotides in length. While the invention is not limited by any particular mechanism of action, the RNAi can enter a cell and cause the degradation of a single-stranded RNA (ssRNA) of similar or identical sequences, including endogenous mRNAs. When a cell is exposed to double-stranded RNA (dsRNA), mRNA from the homologous gene is selectively degraded by a process called RNA interference (RNAi). A possible basic mechanism behind RNAi is the breaking of a double-stranded RNA (dsRNA) matching a specific gene sequence into short pieces called short interfering RNA, which trigger the degradation of mRNA that matches its sequence. In one aspect, the RNAi's of the invention are used in gene-silencing therapeutics, see, e.g., Shuey (2002) Drug Discov. Today 7:1040-1046. In one aspect, the invention provides methods to selectively degrade RNA using the RNAi's of the invention. The process may be practiced *in vitro*, *ex vivo* or *in vivo*. In one aspect, the RNAi molecules of the invention can be used to generate a loss-of-function mutation in a cell, an organ or an animal. Methods for making and using RNAi molecules for selectively degrade RNA are well known in the art, see, e.g., U.S. Patent No. 6,506,559; 6,511,824; 6,515,109; 6,489,127.

Modification of Nucleic Acids

The invention provides methods of generating variants of the nucleic acids of the invention, e.g., those encoding an amylase. These methods can be repeated or used in various combinations to generate amylases having an altered or different activity or an altered or different stability from that of an amylase encoded by the template nucleic acid. These methods also can be repeated or used in various combinations, e.g., to generate variations in gene/ message expression, message translation or message stability. In another aspect, the genetic composition of a cell is altered by, e.g., modification of a homologous gene *ex vivo*, followed by its reinsertion into the cell.

A nucleic acid of the invention can be altered by any means. For example, random or stochastic methods, or, non-stochastic, or "directed evolution," methods, see, e.g., U.S. Patent No. 6,361,974. Methods for random mutation of genes are well known in the art, see, e.g., U.S. Patent No. 5,830,696. For example, mutagens can be used to randomly mutate a gene. Mutagens include, e.g., ultraviolet light or gamma irradiation,

or a chemical mutagen, e.g., mitomycin, nitrous acid, photoactivated psoralens, alone or in combination, to induce DNA breaks amenable to repair by recombination. Other chemical mutagens include, for example, sodium bisulfite, nitrous acid, hydroxylamine, hydrazine or formic acid. Other mutagens are analogues of nucleotide precursors, e.g., nitrosoguanidine, 5-bromouracil, 2-aminopurine, or acridine. These agents can be added to a PCR reaction in place of the nucleotide precursor thereby mutating the sequence. Intercalating agents such as proflavine, acriflavine, quinacrine and the like can also be used.

Any technique in molecular biology can be used, e.g., random PCR mutagenesis, see, e.g., Rice (1992) *Proc. Natl. Acad. Sci. USA* 89:5467-5471; or, combinatorial multiple cassette mutagenesis, see, e.g., Cramer (1995) *Biotechniques* 18:194-196. Alternatively, nucleic acids, e.g., genes, can be reassembled after random, or "stochastic," fragmentation, see, e.g., U.S. Patent Nos. 6,291,242; 6,287,862; 6,287,861; 5,955,358; 5,830,721; 5,824,514; 5,811,238; 5,605,793. In alternative aspects, modifications, additions or deletions are introduced by error-prone PCR, shuffling, oligonucleotide-directed mutagenesis, assembly PCR, sexual PCR mutagenesis, in vivo mutagenesis, cassette mutagenesis, recursive ensemble mutagenesis, exponential ensemble mutagenesis, site-specific mutagenesis, gene reassembly, gene site saturated mutagenesis (GSSM), synthetic ligation reassembly (SLR), recombination, recursive sequence recombination, phosphothioate-modified DNA mutagenesis, uracil-containing template mutagenesis, gapped duplex mutagenesis, point mismatch repair mutagenesis, repair-deficient host strain mutagenesis, chemical mutagenesis, radiogenic mutagenesis, deletion mutagenesis, restriction-selection mutagenesis, restriction-purification mutagenesis, artificial gene synthesis, ensemble mutagenesis, chimeric nucleic acid multimer creation, and/or a combination of these and other methods.

The following publications describe a variety of recursive recombination procedures and/or methods which can be incorporated into the methods of the invention: Stemmer (1999) "Molecular breeding of viruses for targeting and other clinical properties" *Tumor Targeting* 4:1-4; Ness (1999) *Nature Biotechnology* 17:893-896; Chang (1999) "Evolution of a cytokine using DNA family shuffling" *Nature Biotechnology* 17:793-797; Minshull (1999) "Protein evolution by molecular breeding" *Current Opinion in Chemical Biology* 3:284-290; Christians (1999) "Directed evolution of thymidine kinase for AZT phosphorylation using DNA family shuffling" *Nature*

Biotechnology 17:259-264; Cramer (1998) "DNA shuffling of a family of genes from diverse species accelerates directed evolution" *Nature* 391:288-291; Cramer (1997) "Molecular evolution of an arsenate detoxification pathway by DNA shuffling," *Nature Biotechnology* 15:436-438; Zhang (1997) "Directed evolution of an effective fucosidase from a galactosidase by DNA shuffling and screening" *Proc. Natl. Acad. Sci. USA* 94:4504-4509; Patten et al. (1997) "Applications of DNA Shuffling to Pharmaceuticals and Vaccines" *Current Opinion in Biotechnology* 8:724-733; Cramer et al. (1996) "Construction and evolution of antibody-phage libraries by DNA shuffling" *Nature Medicine* 2:100-103; Gates et al. (1996) "Affinity selective isolation of ligands from peptide libraries through display on a lac repressor 'headpiece dimer'" *Journal of Molecular Biology* 255:373-386; Stemmer (1996) "Sexual PCR and Assembly PCR" In: *The Encyclopedia of Molecular Biology*. VCH Publishers, New York. pp.447-457; Cramer and Stemmer (1995) "Combinatorial multiple cassette mutagenesis creates all the permutations of mutant and wildtype cassettes" *BioTechniques* 18:194-195; Stemmer et al. (1995) "Single-step assembly of a gene and entire plasmid from large numbers of oligodeoxyribonucleotides" *Gene*, 164:49-53; Stemmer (1995) "The Evolution of Molecular Computation" *Science* 270: 1510; Stemmer (1995) "Searching Sequence Space" *Bio/Technology* 13:549-553; Stemmer (1994) "Rapid evolution of a protein in vitro by DNA shuffling" *Nature* 370:389-391; and Stemmer (1994) "DNA shuffling by random fragmentation and reassembly: In vitro recombination for molecular evolution." *Proc. Natl. Acad. Sci. USA* 91:10747-10751.

Mutational methods of generating diversity include, for example, site-directed mutagenesis (Ling et al. (1997) "Approaches to DNA mutagenesis: an overview" *Anal Biochem.* 254(2): 157-178; Dale et al. (1996) "Oligonucleotide-directed random mutagenesis using the phosphorothioate method" *Methods Mol. Biol.* 57:369-374; Smith (1985) "In vitro mutagenesis" *Ann. Rev. Genet.* 19:423-462; Botstein & Shortle (1985) "Strategies and applications of in vitro mutagenesis" *Science* 229:1193-1201; Carter (1986) "Site-directed mutagenesis" *Biochem. J.* 237:1-7; and Kunkel (1987) "The efficiency of oligonucleotide directed mutagenesis" in *Nucleic Acids & Molecular Biology* (Eckstein, F. and Lilley, D. M. J. eds., Springer Verlag, Berlin)); mutagenesis using uracil containing templates (Kunkel (1985) "Rapid and efficient site-specific mutagenesis without phenotypic selection" *Proc. Natl. Acad. Sci. USA* 82:488-492; Kunkel et al. (1987) "Rapid and efficient site-specific mutagenesis without phenotypic

selection" *Methods in Enzymol.* 154, 367-382; and Bass et al. (1988) "Mutant Trp repressors with new DNA-binding specificities" *Science* 242:240-245); oligonucleotide-directed mutagenesis (*Methods in Enzymol.* 100: 468-500 (1983); *Methods in Enzymol.* 154: 329-350 (1987); Zoller & Smith (1982) "Oligonucleotide-directed mutagenesis using M13-derived vectors: an efficient and general procedure for the production of point mutations in any DNA fragment" *Nucleic Acids Res.* 10:6487-6500; Zoller & Smith (1983) "Oligonucleotide-directed mutagenesis of DNA fragments cloned into M13 vectors" *Methods in Enzymol.* 100:468-500; and Zoller & Smith (1987) "Oligonucleotide-directed mutagenesis: a simple method using two oligonucleotide primers and a single-stranded DNA template" *Methods in Enzymol.* 154:329-350); phosphorothioate-modified DNA mutagenesis (Taylor et al. (1985) "The use of phosphorothioate-modified DNA in restriction enzyme reactions to prepare nicked DNA" *Nucl. Acids Res.* 13: 8749-8764; Taylor et al. (1985) "The rapid generation of oligonucleotide-directed mutations at high frequency using phosphorothioate-modified DNA" *Nucl. Acids Res.* 13: 8765-8787 (1985); Nakamaye (1986) "Inhibition of restriction endonuclease Nci I cleavage by phosphorothioate groups and its application to oligonucleotide-directed mutagenesis" *Nucl. Acids Res.* 14: 9679-9698; Sayers et al. (1988) "Y-T Exonucleases in phosphorothioate-based oligonucleotide-directed mutagenesis" *Nucl. Acids Res.* 16:791-802; and Sayers et al. (1988) "Strand specific cleavage of phosphorothioate-containing DNA by reaction with restriction endonucleases in the presence of ethidium bromide" *Nucl. Acids Res.* 16: 803-814); mutagenesis using gapped duplex DNA (Kramer et al. (1984) "The gapped duplex DNA approach to oligonucleotide-directed mutation construction" *Nucl. Acids Res.* 12: 9441-9456; Kramer & Fritz (1987) *Methods in Enzymol.* "Oligonucleotide-directed construction of mutations via gapped duplex DNA" 154:350-367; Kramer et al. (1988) "Improved enzymatic *in vitro* reactions in the gapped duplex DNA approach to oligonucleotide-directed construction of mutations" *Nucl. Acids Res.* 16: 7207; and Fritz et al. (1988) "Oligonucleotide-directed construction of mutations: a gapped duplex DNA procedure without enzymatic reactions *in vitro*" *Nucl. Acids Res.* 16: 6987-6999).

Additional protocols that can be used to practice the invention include point mismatch repair (Kramer (1984) "Point Mismatch Repair" *Cell* 38:879-887), mutagenesis using repair-deficient host strains (Carter et al. (1985) "Improved oligonucleotide site-directed mutagenesis using M13 vectors" *Nucl. Acids Res.* 13: 4431-

4443; and Carter (1987) "Improved oligonucleotide-directed mutagenesis using M13 vectors" *Methods in Enzymol.* 154: 382-403), deletion mutagenesis (Eghtedarzadeh (1986) "Use of oligonucleotides to generate large deletions" *Nucl. Acids Res.* 14: 5115), restriction-selection and restriction-selection and restriction-purification (Wells et al. 5 (1986) "Importance of hydrogen-bond formation in stabilizing the transition state of subtilisin" *Phil. Trans. R. Soc. Lond. A* 317: 415-423), mutagenesis by total gene synthesis (Nambiar et al. (1984) "Total synthesis and cloning of a gene coding for the ribonuclease S protein" *Science* 223: 1299-1301; Sakamar and Khorana (1988) "Total synthesis and expression of a gene for the α -subunit of bovine rod outer segment guanine 10 nucleotide-binding protein (transducin)" *Nucl. Acids Res.* 14: 6361-6372; Wells et al. (1985) "Cassette mutagenesis: an efficient method for generation of multiple mutations at defined sites" *Gene* 34:315-323; and Grundstrom et al. (1985) "Oligonucleotide-directed mutagenesis by microscale 'shot-gun' gene synthesis" *Nucl. Acids Res.* 13: 3305-3316), double-strand break repair (Mandecki (1986); Arnold (1993) "Protein engineering for 15 unusual environments" *Current Opinion in Biotechnology* 4:450-455. "Oligonucleotide-directed double-strand break repair in plasmids of *Escherichia coli*: a method for site-specific mutagenesis" *Proc. Natl. Acad. Sci. USA*, 83:7177-7181). Additional details on many of the above methods can be found in *Methods in Enzymology* Volume 154, which also describes useful controls for trouble-shooting problems with various mutagenesis 20 methods.

Protocols that can be used to practice the invention are described, e.g., in U.S. Patent Nos. 5,605,793 to Stemmer (Feb. 25, 1997), "Methods for In Vitro Recombination;" U.S. Pat. No. 5,811,238 to Stemmer et al. (Sep. 22, 1998) "Methods for 25 Generating Polynucleotides having Desired Characteristics by Iterative Selection and Recombination;" U.S. Pat. No. 5,830,721 to Stemmer et al. (Nov. 3, 1998), "DNA Mutagenesis by Random Fragmentation and Reassembly;" U.S. Pat. No. 5,834,252 to Stemmer, et al. (Nov. 10, 1998) "End-Complementary Polymerase Reaction;" U.S. Pat. No. 5,837,458 to Minshull, et al. (Nov. 17, 1998), "Methods and Compositions for Cellular and Metabolic Engineering;" WO 95/22625, Stemmer and Crameri, 30 "Mutagenesis by Random Fragmentation and Reassembly;" WO 96/33207 by Stemmer and Lipschutz "End Complementary Polymerase Chain Reaction;" WO 97/20078 by Stemmer and Crameri "Methods for Generating Polynucleotides having Desired Characteristics by Iterative Selection and Recombination;" WO 97/35966 by Minshull

and Stemmer, "Methods and Compositions for Cellular and Metabolic Engineering;" WO 99/41402 by Punnonen et al. "Targeting of Genetic Vaccine Vectors;" WO 99/41383 by Punnonen et al. "Antigen Library Immunization;" WO 99/41369 by Punnonen et al. "Genetic Vaccine Vector Engineering;" WO 99/41368 by Punnonen et al. "Optimization
5 of Immunomodulatory Properties of Genetic Vaccines;" EP 752008 by Stemmer and Crameri, "DNA Mutagenesis by Random Fragmentation and Reassembly;" EP 0932670 by Stemmer "Evolving Cellular DNA Uptake by Recursive Sequence Recombination;" WO 99/23107 by Stemmer et al., "Modification of Virus Tropism and Host Range by Viral Genome Shuffling;" WO 99/21979 by Apt et al., "Human Papillomavirus Vectors;"
10 WO 98/31837 by del Cardayre et al. "Evolution of Whole Cells and Organisms by Recursive Sequence Recombination;" WO 98/27230 by Patten and Stemmer, "Methods and Compositions for Polypeptide Engineering;" WO 98/27230 by Stemmer et al., "Methods for Optimization of Gene Therapy by Recursive Sequence Shuffling and Selection," WO 00/00632, "Methods for Generating Highly Diverse Libraries," WO
15 00/09679, "Methods for Obtaining in Vitro Recombined Polynucleotide Sequence Banks and Resulting Sequences," WO 98/42832 by Arnold et al., "Recombination of Polynucleotide Sequences Using Random or Defined Primers," WO 99/29902 by Arnold et al., "Method for Creating Polynucleotide and Polypeptide Sequences," WO 98/41653 by Vind, "An in Vitro Method for Construction of a DNA Library," WO 98/41622 by
20 Borchert et al., "Method for Constructing a Library Using DNA Shuffling," and WO 98/42727 by Pati and Zarling, "Sequence Alterations using Homologous Recombination."

Protocols that can be used to practice the invention (providing details regarding various diversity generating methods) are described, e.g., in U.S. Patent application serial no. (USSN) 09/407,800, "SHUFFLING OF CODON ALTERED
25 GENES" by Patten et al. filed Sep. 28, 1999; "EVOLUTION OF WHOLE CELLS AND ORGANISMS BY RECURSIVE SEQUENCE RECOMBINATION" by del Cardayre et al., United States Patent No. 6,379,964; "OLIGONUCLEOTIDE MEDIATED NUCLEIC ACID RECOMBINATION" by Crameri et al., United States Patent Nos. 6,319,714; 6,368,861; 6,376,246; 6,423,542; 6,426,224 and PCT/US00/01203; "USE OF CODON-
30 VARIED OLIGONUCLEOTIDE SYNTHESIS FOR SYNTHETIC SHUFFLING" by Welch et al., United States Patent No. 6,436,675; "METHODS FOR MAKING CHARACTER STRINGS, POLYNUCLEOTIDES & POLYPEPTIDES HAVING DESIRED CHARACTERISTICS" by Selifonov et al., filed Jan. 18, 2000,

(PCT/US00/01202) and, e.g. "METHODS FOR MAKING CHARACTER STRINGS, POLYNUCLEOTIDES & POLYPEPTIDES HAVING DESIRED CHARACTERISTICS" by Selifonov et al., filed Jul. 18, 2000 (U.S. Ser. No. 09/618,579); "METHODS OF POPULATING DATA STRUCTURES FOR USE IN EVOLUTIONARY SIMULATIONS" by Selifonov and Stemmer, filed Jan. 18, 2000 (PCT/US00/01138); and "SINGLE-STRANDED NUCLEIC ACID TEMPLATE-MEDIATED RECOMBINATION AND NUCLEIC ACID FRAGMENT ISOLATION" by Affholter, filed Sep. 6, 2000 (U.S. Ser. No. 09/656,549); and United States Patent Nos. 6,177,263; 6,153,410.

Non-stochastic, or "directed evolution," methods include, e.g., gene site saturation mutagenesis (GSSMTM), synthetic ligation reassembly (SLR), or a combination thereof are used to modify the nucleic acids of the invention to generate amylases with new or altered properties (e.g., activity under highly acidic or alkaline conditions, high temperatures, and the like). Polypeptides encoded by the modified nucleic acids can be screened for an activity before testing for proteolytic or other activity. Any testing modality or protocol can be used, e.g., using a capillary array platform. See, e.g., U.S. Patent Nos. 6,361,974; 6,280,926; 5,939,250.

Saturation mutagenesis, or, GSSMTM

In one aspect, codon primers containing a degenerate N,N,G/T sequence are used to introduce point mutations into a polynucleotide, e.g., an amylase or an antibody of the invention, so as to generate a set of progeny polypeptides in which a full range of single amino acid substitutions is represented at each amino acid position, e.g., an amino acid residue in an enzyme active site or ligand binding site targeted to be modified. These oligonucleotides can comprise a contiguous first homologous sequence, a degenerate N,N,G/T sequence, and, optionally, a second homologous sequence. The downstream progeny translational products from the use of such oligonucleotides include all possible amino acid changes at each amino acid site along the polypeptide, because the degeneracy of the N,N,G/T sequence includes codons for all 20 amino acids. In one aspect, one such degenerate oligonucleotide (comprised of, e.g., one degenerate N,N,G/T cassette) is used for subjecting each original codon in a parental polynucleotide template to a full range of codon substitutions. In another aspect, at least two degenerate cassettes are used – either in the same oligonucleotide or not, for subjecting at least two original

codons in a parental polynucleotide template to a full range of codon substitutions. For example, more than one N,N,G/T sequence can be contained in one oligonucleotide to introduce amino acid mutations at more than one site. This plurality of N,N,G/T sequences can be directly contiguous, or separated by one or more additional nucleotide sequence(s). In another aspect, oligonucleotides serviceable for introducing additions and deletions can be used either alone or in combination with the codons containing an N,N,G/T sequence, to introduce any combination or permutation of amino acid additions, deletions, and/or substitutions.

In one aspect, simultaneous mutagenesis of two or more contiguous amino acid positions is done using an oligonucleotide that contains contiguous N,N,G/T triplets, i.e. a degenerate (N,N,G/T)_n sequence. In another aspect, degenerate cassettes having less degeneracy than the N,N,G/T sequence are used. For example, it may be desirable in some instances to use (e.g. in an oligonucleotide) a degenerate triplet sequence comprised of only one N, where said N can be in the first second or third position of the triplet. Any other bases including any combinations and permutations thereof can be used in the remaining two positions of the triplet. Alternatively, it may be desirable in some instances to use (e.g. in an oligo) a degenerate N,N,N triplet sequence.

In one aspect, use of degenerate triplets (e.g., N,N,G/T triplets) allows for systematic and easy generation of a full range of possible natural amino acids (for a total of 20 amino acids) into each and every amino acid position in a polypeptide (in alternative aspects, the methods also include generation of less than all possible substitutions per amino acid residue, or codon, position). For example, for a 100 amino acid polypeptide, 2000 distinct species (i.e. 20 possible amino acids per position X 100 amino acid positions) can be generated. Through the use of an oligonucleotide or set of oligonucleotides containing a degenerate N,N,G/T triplet, 32 individual sequences can code for all 20 possible natural amino acids. Thus, in a reaction vessel in which a parental polynucleotide sequence is subjected to saturation mutagenesis using at least one such oligonucleotide, there are generated 32 distinct progeny polynucleotides encoding 20 distinct polypeptides. In contrast, the use of a non-degenerate oligonucleotide in site-directed mutagenesis leads to only one progeny polypeptide product per reaction vessel. Nondegenerate oligonucleotides can optionally be used in combination with degenerate primers disclosed; for example, nondegenerate oligonucleotides can be used to generate specific point mutations in a working polynucleotide. This provides one means to

generate specific silent point mutations, point mutations leading to corresponding amino acid changes, and point mutations that cause the generation of stop codons and the corresponding expression of polypeptide fragments.

In one aspect, each saturation mutagenesis reaction vessel contains
 5 polynucleotides encoding at least 20 progeny polypeptide (e.g., amylases) molecules such that all 20 natural amino acids are represented at the one specific amino acid position corresponding to the codon position mutagenized in the parental polynucleotide (other aspects use less than all 20 natural combinations). The 32-fold degenerate progeny polypeptides generated from each saturation mutagenesis reaction vessel can be subjected
 10 to clonal amplification (e.g. cloned into a suitable host, e.g., E. coli host, using, e.g., an expression vector) and subjected to expression screening. When an individual progeny polypeptide is identified by screening to display a favorable change in property (when compared to the parental polypeptide, such as increased proteolytic activity under alkaline or acidic conditions), it can be sequenced to identify the correspondingly favorable amino
 15 acid substitution contained therein.

In one aspect, upon mutagenizing each and every amino acid position in a parental polypeptide using saturation mutagenesis as disclosed herein, favorable amino acid changes may be identified at more than one amino acid position. One or more new progeny molecules can be generated that contain a combination of all or part of these
 20 favorable amino acid substitutions. For example, if 2 specific favorable amino acid changes are identified in each of 3 amino acid positions in a polypeptide, the permutations include 3 possibilities at each position (no change from the original amino acid, and each of two favorable changes) and 3 positions. Thus, there are $3 \times 3 \times 3$ or 27 total possibilities, including 7 that were previously examined - 6 single point mutations
 25 (i.e. 2 at each of three positions) and no change at any position.

In another aspect, site-saturation mutagenesis can be used together with another stochastic or non-stochastic means to vary sequence, e.g., synthetic ligation reassembly (see below), shuffling, chimerization, recombination and other mutagenizing processes and mutagenizing agents. This invention provides for the use of any
 30 mutagenizing process(es), including saturation mutagenesis, in an iterative manner.

Synthetic Ligation Reassembly (SLR)

The invention provides a non-stochastic gene modification system termed

“synthetic ligation reassembly,” or simply “SLR,” a “directed evolution process,” to generate polypeptides, e.g., amylases or antibodies of the invention, with new or altered properties. SLR is a method of ligating oligonucleotide fragments together non-stochastically. This method differs from stochastic oligonucleotide shuffling in that the nucleic acid building blocks are not shuffled, concatenated or chimerized randomly, but rather are assembled non-stochastically. See, e.g., U.S. Patent Application Serial No. (USSN) 09/332,835 entitled “Synthetic Ligation Reassembly in Directed Evolution” and filed on June 14, 1999 (“USSN 09/332,835”). In one aspect, SLR comprises the following steps: (a) providing a template polynucleotide, wherein the template polynucleotide comprises sequence encoding a homologous gene; (b) providing a plurality of building block polynucleotides, wherein the building block polynucleotides are designed to cross-over reassemble with the template polynucleotide at a predetermined sequence, and a building block polynucleotide comprises a sequence that is a variant of the homologous gene and a sequence homologous to the template polynucleotide flanking the variant sequence; (c) combining a building block polynucleotide with a template polynucleotide such that the building block polynucleotide cross-over reassembles with the template polynucleotide to generate polynucleotides comprising homologous gene sequence variations.

SLR does not depend on the presence of high levels of homology between polynucleotides to be rearranged. Thus, this method can be used to non-stochastically generate libraries (or sets) of progeny molecules comprised of over 10^{100} different chimeras. SLR can be used to generate libraries comprised of over 10^{1000} different progeny chimeras. Thus, aspects of the present invention include non-stochastic methods of producing a set of finalized chimeric nucleic acid molecule having an overall assembly order that is chosen by design. This method includes the steps of generating by design a plurality of specific nucleic acid building blocks having serviceable mutually compatible ligatable ends, and assembling these nucleic acid building blocks, such that a designed overall assembly order is achieved.

The mutually compatible ligatable ends of the nucleic acid building blocks to be assembled are considered to be “serviceable” for this type of ordered assembly if they enable the building blocks to be coupled in predetermined orders. Thus, the overall assembly order in which the nucleic acid building blocks can be coupled is specified by the design of the ligatable ends. If more than one assembly step is to be used, then the

overall assembly order in which the nucleic acid building blocks can be coupled is also specified by the sequential order of the assembly step(s). In one aspect, the annealed building pieces are treated with an enzyme, such as a ligase (e.g. T4 DNA ligase), to achieve covalent bonding of the building pieces.

5 In one aspect, the design of the oligonucleotide building blocks is obtained by analyzing a set of progenitor nucleic acid sequence templates that serve as a basis for producing a progeny set of finalized chimeric polynucleotides. These parental oligonucleotide templates thus serve as a source of sequence information that aids in the design of the nucleic acid building blocks that are to be mutagenized, e.g., chimerized or shuffled. In one aspect of this method, the sequences of a plurality of parental nucleic acid templates are aligned in order to select one or more demarcation points. The demarcation points can be located at an area of homology, and are comprised of one or more nucleotides. These demarcation points are preferably shared by at least two of the progenitor templates. The demarcation points can thereby be used to delineate the boundaries of oligonucleotide building blocks to be generated in order to rearrange the parental polynucleotides. The demarcation points identified and selected in the progenitor molecules serve as potential chimerization points in the assembly of the final chimeric progeny molecules. A demarcation point can be an area of homology (comprised of at least one homologous nucleotide base) shared by at least two parental polynucleotide sequences. Alternatively, a demarcation point can be an area of homology that is shared by at least half of the parental polynucleotide sequences, or, it can be an area of homology that is shared by at least two thirds of the parental polynucleotide sequences. Even more preferably a serviceable demarcation points is an area of homology that is shared by at least three fourths of the parental polynucleotide sequences, or, it can be shared by at almost all of the parental polynucleotide sequences. In one aspect, a demarcation point is an area of homology that is shared by all of the parental polynucleotide sequences.

15 20 25 30 In one aspect, a ligation reassembly process is performed exhaustively in order to generate an exhaustive library of progeny chimeric polynucleotides. In other words, all possible ordered combinations of the nucleic acid building blocks are represented in the set of finalized chimeric nucleic acid molecules. At the same time, in another aspect, the assembly order (i.e. the order of assembly of each building block in the 5' to 3' sequence of each finalized chimeric nucleic acid) in each combination is by

design (or non-stochastic) as described above. Because of the non-stochastic nature of this invention, the possibility of unwanted side products is greatly reduced.

In another aspect, the ligation reassembly method is performed systematically. For example, the method is performed in order to generate a systematically compartmentalized library of progeny molecules, with compartments that can be screened systematically, e.g. one by one. In other words this invention provides that, through the selective and judicious use of specific nucleic acid building blocks, coupled with the selective and judicious use of sequentially stepped assembly reactions, a design can be achieved where specific sets of progeny products are made in each of several reaction vessels. This allows a systematic examination and screening procedure to be performed. Thus, these methods allow a potentially very large number of progeny molecules to be examined systematically in smaller groups. Because of its ability to perform chimerizations in a manner that is highly flexible yet exhaustive and systematic as well, particularly when there is a low level of homology among the progenitor molecules, these methods provide for the generation of a library (or set) comprised of a large number of progeny molecules. Because of the non-stochastic nature of the instant ligation reassembly invention, the progeny molecules generated preferably comprise a library of finalized chimeric nucleic acid molecules having an overall assembly order that is chosen by design. The saturation mutagenesis and optimized directed evolution methods also can be used to generate different progeny molecular species. It is appreciated that the invention provides freedom of choice and control regarding the selection of demarcation points, the size and number of the nucleic acid building blocks, and the size and design of the couplings. It is appreciated, furthermore, that the requirement for intermolecular homology is highly relaxed for the operability of this invention. In fact, demarcation points can even be chosen in areas of little or no intermolecular homology. For example, because of codon wobble, i.e. the degeneracy of codons, nucleotide substitutions can be introduced into nucleic acid building blocks without altering the amino acid originally encoded in the corresponding progenitor template. Alternatively, a codon can be altered such that the coding for an originally amino acid is altered. This invention provides that such substitutions can be introduced into the nucleic acid building block in order to increase the incidence of intermolecular homologous demarcation points and thus to allow an increased number of couplings to be

achieved among the building blocks, which in turn allows a greater number of progeny chimeric molecules to be generated.

In another aspect, the synthetic nature of the step in which the building blocks are generated allows the design and introduction of nucleotides (e.g., one or more nucleotides, which may be, for example, codons or introns or regulatory sequences) that can later be optionally removed in an in vitro process (e.g. by mutagenesis) or in an in vivo process (e.g. by utilizing the gene splicing ability of a host organism). It is appreciated that in many instances the introduction of these nucleotides may also be desirable for many other reasons in addition to the potential benefit of creating a serviceable demarcation point.

In one aspect, a nucleic acid building block is used to introduce an intron. Thus, functional introns are introduced into a man-made gene manufactured according to the methods described herein. The artificially introduced intron(s) can be functional in a host cells for gene splicing much in the way that naturally-occurring introns serve functionally in gene splicing.

Optimized Directed Evolution System

The invention provides a non-stochastic gene modification system termed "optimized directed evolution system" to generate polypeptides, e.g., amylases or antibodies of the invention, with new or altered properties. Optimized directed evolution is directed to the use of repeated cycles of reductive reassortment, recombination and selection that allow for the directed molecular evolution of nucleic acids through recombination. Optimized directed evolution allows generation of a large population of evolved chimeric sequences, wherein the generated population is significantly enriched for sequences that have a predetermined number of crossover events.

A crossover event is a point in a chimeric sequence where a shift in sequence occurs from one parental variant to another parental variant. Such a point is normally at the juncture of where oligonucleotides from two parents are ligated together to form a single sequence. This method allows calculation of the correct concentrations of oligonucleotide sequences so that the final chimeric population of sequences is enriched for the chosen number of crossover events. This provides more control over choosing chimeric variants having a predetermined number of crossover events.

In addition, this method provides a convenient means for exploring a tremendous amount of the possible protein variant space in comparison to other systems. Previously, if one generated, for example, 10^{13} chimeric molecules during a reaction, it would be extremely difficult to test such a high number of chimeric variants for a particular activity. Moreover, a significant portion of the progeny population would have a very high number of crossover events which resulted in proteins that were less likely to have increased levels of a particular activity. By using these methods, the population of chimeric molecules can be enriched for those variants that have a particular number of crossover events. Thus, although one can still generate 10^{13} chimeric molecules during a reaction, each of the molecules chosen for further analysis most likely has, for example, only three crossover events. Because the resulting progeny population can be skewed to have a predetermined number of crossover events, the boundaries on the functional variety between the chimeric molecules is reduced. This provides a more manageable number of variables when calculating which oligonucleotide from the original parental polynucleotides might be responsible for affecting a particular trait.

One method for creating a chimeric progeny polynucleotide sequence is to create oligonucleotides corresponding to fragments or portions of each parental sequence. Each oligonucleotide preferably includes a unique region of overlap so that mixing the oligonucleotides together results in a new variant that has each oligonucleotide fragment assembled in the correct order. Additional information can also be found, e.g., in USSN 09/332,835; U.S. Patent No. 6,361,974.

The number of oligonucleotides generated for each parental variant bears a relationship to the total number of resulting crossovers in the chimeric molecule that is ultimately created. For example, three parental nucleotide sequence variants might be provided to undergo a ligation reaction in order to find a chimeric variant having, for example, greater activity at high temperature. As one example, a set of 50 oligonucleotide sequences can be generated corresponding to each portions of each parental variant. Accordingly, during the ligation reassembly process there could be up to 50 crossover events within each of the chimeric sequences. The probability that each of the generated chimeric polynucleotides will contain oligonucleotides from each parental variant in alternating order is very low. If each oligonucleotide fragment is present in the ligation reaction in the same molar quantity it is likely that in some positions oligonucleotides from the same parental polynucleotide will ligate next to one another

and thus not result in a crossover event. If the concentration of each oligonucleotide from each parent is kept constant during any ligation step in this example, there is a 1/3 chance (assuming 3 parents) that an oligonucleotide from the same parental variant will ligate within the chimeric sequence and produce no crossover.

Accordingly, a probability density function (PDF) can be determined to predict the population of crossover events that are likely to occur during each step in a ligation reaction given a set number of parental variants, a number of oligonucleotides corresponding to each variant, and the concentrations of each variant during each step in the ligation reaction. The statistics and mathematics behind determining the PDF is described below. By utilizing these methods, one can calculate such a probability density function, and thus enrich the chimeric progeny population for a predetermined number of crossover events resulting from a particular ligation reaction. Moreover, a target number of crossover events can be predetermined, and the system then programmed to calculate the starting quantities of each parental oligonucleotide during each step in the ligation reaction to result in a probability density function that centers on the predetermined number of crossover events. These methods are directed to the use of repeated cycles of reductive reassortment, recombination and selection that allow for the directed molecular evolution of a nucleic acid encoding a polypeptide through recombination. This system allows generation of a large population of evolved chimeric sequences, wherein the generated population is significantly enriched for sequences that have a predetermined number of crossover events. A crossover event is a point in a chimeric sequence where a shift in sequence occurs from one parental variant to another parental variant. Such a point is normally at the juncture of where oligonucleotides from two parents are ligated together to form a single sequence. The method allows calculation of the correct concentrations of oligonucleotide sequences so that the final chimeric population of sequences is enriched for the chosen number of crossover events. This provides more control over choosing chimeric variants having a predetermined number of crossover events.

In addition, these methods provide a convenient means for exploring a tremendous amount of the possible protein variant space in comparison to other systems. By using the methods described herein, the population of chimerics molecules can be enriched for those variants that have a particular number of crossover events. Thus, although one can still generate 10^{13} chimeric molecules during a reaction, each of the

molecules chosen for further analysis most likely has, for example, only three crossover events. Because the resulting progeny population can be skewed to have a predetermined number of crossover events, the boundaries on the functional variety between the chimeric molecules is reduced. This provides a more manageable number of variables when calculating which oligonucleotide from the original parental polynucleotides might be responsible for affecting a particular trait.

In one aspect, the method creates a chimeric progeny polynucleotide sequence by creating oligonucleotides corresponding to fragments or portions of each parental sequence. Each oligonucleotide preferably includes a unique region of overlap so that mixing the oligonucleotides together results in a new variant that has each oligonucleotide fragment assembled in the correct order. See also USSN 09/332,835.

Determining Crossover Events

Aspects of the invention include a system and software that receive a desired crossover probability density function (PDF), the number of parent genes to be reassembled, and the number of fragments in the reassembly as inputs. The output of this program is a "fragment PDF" that can be used to determine a recipe for producing reassembled genes, and the estimated crossover PDF of those genes. The processing described herein is preferably performed in MATLAB™ (The Mathworks, Natick, Massachusetts) a programming language and development environment for technical computing.

Iterative Processes

In practicing the invention, these processes can be iteratively repeated. For example, a nucleic acid (or, the nucleic acid) responsible for an altered or new amylase phenotype is identified, re-isolated, again modified, re-tested for activity. This process can be iteratively repeated until a desired phenotype is engineered. For example, an entire biochemical anabolic or catabolic pathway can be engineered into a cell, including, e.g., starch hydrolysis activity.

Similarly, if it is determined that a particular oligonucleotide has no affect at all on the desired trait (e.g., a new amylase phenotype), it can be removed as a variable by synthesizing larger parental oligonucleotides that include the sequence to be removed. Since incorporating the sequence within a larger sequence prevents any crossover events, there will no longer be any variation of this sequence in the progeny polynucleotides.

This iterative practice of determining which oligonucleotides are most related to the desired trait, and which are unrelated, allows more efficient exploration all of the possible protein variants that might be provide a particular trait or activity.

In vivo shuffling

5 *In vivo* shuffling of molecules is use in methods of the invention that provide variants of polypeptides of the invention, e.g., antibodies, amylases, and the like. *In vivo* shuffling can be performed utilizing the natural property of cells to recombine multimers. While recombination *in vivo* has provided the major natural route to molecular diversity, genetic recombination remains a relatively complex process that
10 involves 1) the recognition of homologies; 2) strand cleavage, strand invasion, and metabolic steps leading to the production of recombinant chiasma; and finally 3) the resolution of chiasma into discrete recombined molecules. The formation of the chiasma requires the recognition of homologous sequences.

In one aspect, the invention provides a method for producing a hybrid
15 polynucleotide from at least a first polynucleotide (e.g., an amylase of the invention) and a second polynucleotide (e.g., an enzyme, such as an amylase of the invention or any other amylase, or, a tag or an epitope). The invention can be used to produce a hybrid polynucleotide by introducing at least a first polynucleotide and a second polynucleotide which share at least one region of partial sequence homology into a suitable host cell.
20 The regions of partial sequence homology promote processes which result in sequence reorganization producing a hybrid polynucleotide. The term "hybrid polynucleotide", as used herein, is any nucleotide sequence which results from the method of the present invention and contains sequence from at least two original polynucleotide sequences. Such hybrid polynucleotides can result from intermolecular recombination events which
25 promote sequence integration between DNA molecules. In addition, such hybrid polynucleotides can result from intramolecular reductive reassortment processes which utilize repeated sequences to alter a nucleotide sequence within a DNA molecule.

Producing sequence variants

The invention also provides additional methods for making sequence
30 variants of the nucleic acid (e.g., amylase) sequences of the invention. The invention also provides additional methods for isolating amylases using the nucleic acids and polypeptides of the invention. In one aspect, the invention provides for variants of an

amylase coding sequence (e.g., a gene, cDNA or message) of the invention, which can be altered by any means, including, e.g., random or stochastic methods, or, non-stochastic, or "directed evolution," methods, as described above.

The isolated variants may be naturally occurring. Variant can also be created *in vitro*. Variants may be created using genetic engineering techniques such as site directed mutagenesis, random chemical mutagenesis, Exonuclease III deletion procedures, and standard cloning techniques. Alternatively, such variants, fragments, analogs, or derivatives may be created using chemical synthesis or modification procedures. Other methods of making variants are also familiar to those skilled in the art. These include procedures in which nucleic acid sequences obtained from natural isolates are modified to generate nucleic acids which encode polypeptides having characteristics which enhance their value in industrial or laboratory applications. In such procedures, a large number of variant sequences having one or more nucleotide differences with respect to the sequence obtained from the natural isolate are generated and characterized. These nucleotide differences can result in amino acid changes with respect to the polypeptides encoded by the nucleic acids from the natural isolates.

For example, variants may be created using error prone PCR. In error prone PCR, PCR is performed under conditions where the copying fidelity of the DNA polymerase is low, such that a high rate of point mutations is obtained along the entire length of the PCR product. Error prone PCR is described, e.g., in Leung, D.W., et al., Technique, 1:11-15, 1989) and Caldwell, R. C. & Joyce G.F., PCR Methods Applic., 2:28-33, 1992. Briefly, in such procedures, nucleic acids to be mutagenized are mixed with PCR primers, reaction buffer, $MgCl_2$, $MnCl_2$, Taq polymerase and an appropriate concentration of dNTPs for achieving a high rate of point mutation along the entire length of the PCR product. For example, the reaction may be performed using 20 fmoles of nucleic acid to be mutagenized, 30 pmole of each PCR primer, a reaction buffer comprising 50mM KCl, 10mM Tris HCl (pH 8.3) and 0.01% gelatin, 7mM $MgCl_2$, 0.5mM $MnCl_2$, 5 units of Taq polymerase, 0.2mM dGTP, 0.2mM dATP, 1mM dCTP, and 1mM dTTP. PCR may be performed for 30 cycles of 94°C for 1 min, 45°C for 1 min, and 72°C for 1 min. However, it will be appreciated that these parameters may be varied as appropriate. The mutagenized nucleic acids are cloned into an appropriate vector and the activities of the polypeptides encoded by the mutagenized nucleic acids is evaluated.

Variants may also be created using oligonucleotide directed mutagenesis to generate site-specific mutations in any cloned DNA of interest. Oligonucleotide mutagenesis is described, e.g., in Reidhaar-Olson (1988) *Science* 241:53-57. Briefly, in such procedures a plurality of double stranded oligonucleotides bearing one or more mutations to be introduced into the cloned DNA are synthesized and inserted into the cloned DNA to be mutagenized. Clones containing the mutagenized DNA are recovered and the activities of the polypeptides they encode are assessed.

Another method for generating variants is assembly PCR. Assembly PCR involves the assembly of a PCR product from a mixture of small DNA fragments. A large number of different PCR reactions occur in parallel in the same vial, with the products of one reaction priming the products of another reaction. Assembly PCR is described in, e.g., U.S. Patent No. 5,965,408.

Still another method of generating variants is sexual PCR mutagenesis. In sexual PCR mutagenesis, forced homologous recombination occurs between DNA molecules of different but highly related DNA sequence in vitro, as a result of random fragmentation of the DNA molecule based on sequence homology, followed by fixation of the crossover by primer extension in a PCR reaction. Sexual PCR mutagenesis is described, e.g., in Stemmer (1994) *Proc. Natl. Acad. Sci. USA* 91:10747-10751. Briefly, in such procedures a plurality of nucleic acids to be recombined are digested with DNase to generate fragments having an average size of 50-200 nucleotides. Fragments of the desired average size are purified and resuspended in a PCR mixture. PCR is conducted under conditions which facilitate recombination between the nucleic acid fragments. For example, PCR may be performed by resuspending the purified fragments at a concentration of 10-30ng/:l in a solution of 0.2mM of each dNTP, 2.2mM MgCl₂, 50mM KCL, 10mM Tris HCl, pH 9.0, and 0.1% Triton X-100. 2.5 units of Taq polymerase per 100:1 of reaction mixture is added and PCR is performed using the following regime: 94°C for 60 seconds, 94°C for 30 seconds, 50-55°C for 30 seconds, 72°C for 30 seconds (30-45 times) and 72°C for 5 minutes. However, it will be appreciated that these parameters may be varied as appropriate. In some aspects, oligonucleotides may be included in the PCR reactions. In other aspects, the Klenow fragment of DNA polymerase I may be used in a first set of PCR reactions and Taq polymerase may be used in a subsequent set of PCR reactions. Recombinant sequences are isolated and the activities of the polypeptides they encode are assessed.

Variants may also be created by *in vivo* mutagenesis. In some aspects, random mutations in a sequence of interest are generated by propagating the sequence of interest in a bacterial strain, such as an *E. coli* strain, which carries mutations in one or more of the DNA repair pathways. Such "mutator" strains have a higher random
5 mutation rate than that of a wild-type parent. Propagating the DNA in one of these strains will eventually generate random mutations within the DNA. Mutator strains suitable for use for *in vivo* mutagenesis are described, e.g., in PCT Publication No. WO 91/16427.

Variants may also be generated using cassette mutagenesis. In cassette mutagenesis a small region of a double stranded DNA molecule is replaced with a
10 synthetic oligonucleotide "cassette" that differs from the native sequence. The oligonucleotide often contains completely and/or partially randomized native sequence.

Recursive ensemble mutagenesis may also be used to generate variants. Recursive ensemble mutagenesis is an algorithm for protein engineering (protein
mutagenesis) developed to produce diverse populations of phenotypically related mutants
15 whose members differ in amino acid sequence. This method uses a feedback mechanism to control successive rounds of combinatorial cassette mutagenesis. Recursive ensemble mutagenesis is described, e.g., in Arkin (1992) Proc. Natl. Acad. Sci. USA 89:7811-7815.

In some aspects, variants are created using exponential ensemble mutagenesis. Exponential ensemble mutagenesis is a process for generating
20 combinatorial libraries with a high percentage of unique and functional mutants, wherein small groups of residues are randomized in parallel to identify, at each altered position, amino acids which lead to functional proteins. Exponential ensemble mutagenesis is described, e.g., in Delegrave (1993) Biotechnology Res. 11:1548-1552. Random and site-directed mutagenesis are described, e.g., in Arnold (1993) Current Opinion in
25 Biotechnology 4:450-455.

In some aspects, the variants are created using shuffling procedures wherein portions of a plurality of nucleic acids which encode distinct polypeptides are fused together to create chimeric nucleic acid sequences which encode chimeric polypeptides as described in, e.g., U.S. Patent Nos. 5,965,408; 5,939,250 (see also
30 discussion, above).

The invention also provides variants of polypeptides of the invention (e.g., amylases) comprising sequences in which one or more of the amino acid residues (e.g., of an exemplary polypeptide of the invention) are substituted with a conserved or non-

conserved amino acid residue (e.g., a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code. Conservative substitutions are those that substitute a given amino acid in a polypeptide by another amino acid of like characteristics. Thus, polypeptides of the invention include those with conservative substitutions of sequences of the invention, e.g., the exemplary polypeptides of the invention, including but not limited to the following replacements: replacements of an aliphatic amino acid such as Alanine, Valine, Leucine and Isoleucine with another aliphatic amino acid; replacement of a Serine with a Threonine or vice versa; replacement of an acidic residue such as Aspartic acid and Glutamic acid with another acidic residue; replacement of a residue bearing an amide group, such as Asparagine and Glutamine, with another residue bearing an amide group; exchange of a basic residue such as Lysine and Arginine with another basic residue; and replacement of an aromatic residue such as Phenylalanine, Tyrosine with another aromatic residue. Other variants are those in which one or more of the amino acid residues of the polypeptides of the invention includes a substituent group.

Other variants within the scope of the invention are those in which the polypeptide is associated with another compound, such as a compound to increase the half-life of the polypeptide, for example, polyethylene glycol.

Additional variants within the scope of the invention are those in which additional amino acids are fused to the polypeptide, such as a leader sequence, a secretory sequence, a proprotein sequence or a sequence which facilitates purification, enrichment, or stabilization of the polypeptide.

In some aspects, the variants, fragments, derivatives and analogs of the polypeptides of the invention retain the same biological function or activity as the exemplary polypeptides, e.g., amylase activity, as described herein. In other aspects, the variant, fragment, derivative, or analog includes a proprotein, such that the variant, fragment, derivative, or analog can be activated by cleavage of the proprotein portion to produce an active polypeptide.

Optimizing codons to achieve high levels of protein expression in host cells

The invention provides methods for modifying amylase-encoding nucleic acids to modify codon usage. In one aspect, the invention provides methods for modifying codons in a nucleic acid encoding an amylase to increase or decrease its

expression in a host cell. The invention also provides nucleic acids encoding an amylase modified to increase its expression in a host cell, amylase so modified, and methods of making the modified amylases. The method comprises identifying a "non-preferred" or a "less preferred" codon in amylase-encoding nucleic acid and replacing one or more of these non-preferred or less preferred codons with a "preferred codon" encoding the same amino acid as the replaced codon and at least one non-preferred or less preferred codon in the nucleic acid has been replaced by a preferred codon encoding the same amino acid. A preferred codon is a codon over-represented in coding sequences in genes in the host cell and a non-preferred or less preferred codon is a codon under-represented in coding sequences in genes in the host cell.

Host cells for expressing the nucleic acids, expression cassettes and vectors of the invention include bacteria, yeast, fungi, plant cells, insect cells and mammalian cells. Thus, the invention provides methods for optimizing codon usage in all of these cells, codon-altered nucleic acids and polypeptides made by the codon-altered nucleic acids. Exemplary host cells include gram negative bacteria, such as *Escherichia coli*; gram positive bacteria, such as *Bacillus cereus*, *Streptomyces*, *Lactobacillus gasseri*, *Lactococcus lactis*, *Lactococcus cremoris*, *Bacillus subtilis*. Exemplary host cells also include eukaryotic organisms, e.g., various yeast, such as *Saccharomyces* sp., including *Saccharomyces cerevisiae*, *Schizosaccharomyces pombe*, *Pichia pastoris*, and *Kluyveromyces lactis*, *Hansenula polymorpha*, *Aspergillus niger*, and mammalian cells and cell lines and insect cells and cell lines. Thus, the invention also includes nucleic acids and polypeptides optimized for expression in these organisms and species.

For example, the codons of a nucleic acid encoding an amylase isolated from a bacterial cell are modified such that the nucleic acid is optimally expressed in a bacterial cell different from the bacteria from which the amylase was derived, a yeast, a fungi, a plant cell, an insect cell or a mammalian cell. Methods for optimizing codons are well known in the art, see, e.g., U.S. Patent No. 5,795,737; Baca (2000) Int. J. Parasitol. 30:113-118; Hale (1998) Protein Expr. Purif. 12:185-188; Narum (2001) Infect. Immun. 69:7250-7253. See also Narum (2001) Infect. Immun. 69:7250-7253, describing optimizing codons in mouse systems; Outchkourov (2002) Protein Expr. Purif. 24:18-24, describing optimizing codons in yeast; Feng (2000) Biochemistry 39:15399-15409, describing optimizing codons in *E. coli*; Humphreys (2000) Protein Expr. Purif. 20:252-264, describing optimizing codon usage that affects secretion in *E. coli*.

Transgenic non-human animals

The invention provides transgenic non-human animals comprising a nucleic acid, a polypeptide (e.g., an amylase), an expression cassette or vector or a transfected or transformed cell of the invention. The invention also provides methods of making and using these transgenic non-human animals.

The transgenic non-human animals can be, e.g., goats, rabbits, sheep, pigs, cows, rats and mice, comprising the nucleic acids of the invention. These animals can be used, e.g., as *in vivo* models to study amylase activity, or, as models to screen for agents that change the amylase activity *in vivo*. The coding sequences for the polypeptides to be expressed in the transgenic non-human animals can be designed to be constitutive, or, under the control of tissue-specific, developmental-specific or inducible transcriptional regulatory factors. Transgenic non-human animals can be designed and generated using any method known in the art; see, e.g., U.S. Patent Nos. 6,211,428; 6,187,992; 6,156,952; 6,118,044; 6,111,166; 6,107,541; 5,959,171; 5,922,854; 5,892,070; 5,880,327; 5,891,698; 5,639,940; 5,573,933; 5,387,742; 5,087,571, describing making and using transformed cells and eggs and transgenic mice, rats, rabbits, sheep, pigs and cows. See also, e.g., Pollock (1999) J. Immunol. Methods 231:147-157, describing the production of recombinant proteins in the milk of transgenic dairy animals; Baguisi (1999) Nat. Biotechnol. 17:456-461, demonstrating the production of transgenic goats. U.S. Patent No. 6,211,428, describes making and using transgenic non-human mammals which express in their brains a nucleic acid construct comprising a DNA sequence. U.S. Patent No. 5,387,742, describes injecting cloned recombinant or synthetic DNA sequences into fertilized mouse eggs, implanting the injected eggs in pseudo-pregnant females, and growing to term transgenic mice whose cells express proteins related to the pathology of Alzheimer's disease. U.S. Patent No. 6,187,992, describes making and using a transgenic mouse whose genome comprises a disruption of the gene encoding amyloid precursor protein (APP).

"Knockout animals" can also be used to practice the methods of the invention. For example, in one aspect, the transgenic or modified animals of the invention comprise a "knockout animal," e.g., a "knockout mouse," engineered not to express an endogenous gene, which is replaced with a gene expressing an amylase of the invention, or, a fusion protein comprising an amylase of the invention.

Transgenic Plants and Seeds

The invention provides transgenic plants and seeds comprising a nucleic acid, a polypeptide (e.g., an amylase, such as an alpha amylase), an expression cassette or vector or a transfected or transformed cell of the invention. The invention also provides
5 plant products, e.g., oils, seeds, leaves, extracts and the like, comprising a nucleic acid and/or a polypeptide (e.g., an amylase, such as an alpha amylase) of the invention. The transgenic plant can be dicotyledonous (a dicot) or monocotyledonous (a monocot). The invention also provides methods of making and using these transgenic plants and seeds. The transgenic plant or plant cell expressing a polypeptide of the present invention may
10 be constructed in accordance with any method known in the art. See, for example, U.S. Patent No. 6,309,872.

Nucleic acids and expression constructs of the invention can be introduced into a plant cell by any means. For example, nucleic acids or expression constructs can be introduced into the genome of a desired plant host, or, the nucleic acids or expression
15 constructs can be episomes. Introduction into the genome of a desired plant can be such that the host's a-amylase production is regulated by endogenous transcriptional or translational control elements. The invention also provides "knockout plants" where insertion of gene sequence by, e.g., homologous recombination, has disrupted the expression of the endogenous gene. Means to generate "knockout" plants are well-known
20 in the art, see, e.g., Strepp (1998) Proc Natl. Acad. Sci. USA 95:4368-4373; Miao (1995) Plant J 7:359-365. See discussion on transgenic plants, below.

The nucleic acids of the invention can be used to confer desired traits on essentially any plant, e.g., on starch-producing plants, such as potato, wheat, rice, barley, and the like. Nucleic acids of the invention can be used to manipulate metabolic
25 pathways of a plant in order to optimize or alter host's expression of a-amylase. The can change the ratio of starch/sugar conversion in a plant. This can facilitate industrial processing of a plant. Alternatively, alpha-amylases of the invention can be used in production of a transgenic plant to produce a compound not naturally produced by that plant. This can lower production costs or create a novel product.

30 In one aspect, the first step in production of a transgenic plant involves making an expression construct for expression in a plant cell. These techniques are well known in the art. They can include selecting and cloning a promoter, a coding sequence for facilitating efficient binding of ribosomes to mRNA and selecting the appropriate

gene terminator sequences. One exemplary constitutive promoter is CaMV35S, from the cauliflower mosaic virus, which generally results in a high degree of expression in plants. Other promoters are more specific and respond to cues in the plant's internal or external environment. An exemplary light-inducible promoter is the promoter from the *cab* gene, encoding the major chlorophyll *a/b* binding protein.

In one aspect, the nucleic acid is modified to achieve greater expression in a plant cell. For example, a sequence of the invention is likely to have a higher percentage of A-T nucleotide pairs compared to that seen in a plant, some of which prefer G-C nucleotide pairs. Therefore, A-T nucleotides in the coding sequence can be substituted with G-C nucleotides without significantly changing the amino acid sequence to enhance production of the gene product in plant cells.

Selectable marker gene can be added to the gene construct in order to identify plant cells or tissues that have successfully integrated the transgene. This may be necessary because achieving incorporation and expression of genes in plant cells is a rare event, occurring in just a few percent of the targeted tissues or cells. Selectable marker genes encode proteins that provide resistance to agents that are normally toxic to plants, such as antibiotics or herbicides. Only plant cells that have integrated the selectable marker gene will survive when grown on a medium containing the appropriate antibiotic or herbicide. As for other inserted genes, marker genes also require promoter and termination sequences for proper function.

In one aspect, making transgenic plants or seeds comprises incorporating sequences of the invention and, optionally, marker genes into a target expression construct (e.g., a plasmid), along with positioning of the promoter and the terminator sequences. This can involve transferring the modified gene into the plant through a suitable method. For example, a construct may be introduced directly into the genomic DNA of the plant cell using techniques such as electroporation and microinjection of plant cell protoplasts, or the constructs can be introduced directly to plant tissue using ballistic methods, such as DNA particle bombardment. For example, see, e.g., Christou (1997) *Plant Mol. Biol.* 35:197-203; Pawlowski (1996) *Mol. Biotechnol.* 6:17-30; Klein (1987) *Nature* 327:70-73; Takumi (1997) *Genes Genet. Syst.* 72:63-69, discussing use of particle bombardment to introduce transgenes into wheat; and Adam (1997) *supra*, for use of particle bombardment to introduce YACs into plant cells. For example, Rinehart (1997) *supra*, used particle bombardment to generate transgenic cotton plants. Apparatus

for accelerating particles is described U.S. Pat. No. 5,015,580; and, the commercially available BioRad (Biolistics) PDS-2000 particle acceleration instrument; see also, John, U.S. Patent No. 5,608,148; and Ellis, U.S. Patent No. 5,681,730, describing particle-mediated transformation of gymnosperms.

5 In one aspect, protoplasts can be immobilized and injected with a nucleic acids, e.g., an expression construct. Although plant regeneration from protoplasts is not easy with cereals, plant regeneration is possible in legumes using somatic embryogenesis from protoplast derived callus. Organized tissues can be transformed with naked DNA using gene gun technique, where DNA is coated on tungsten microprojectiles, shot
10 1/100th the size of cells, which carry the DNA deep into cells and organelles. Transformed tissue is then induced to regenerate, usually by somatic embryogenesis. This technique has been successful in several cereal species including maize and rice.

Nucleic acids, e.g., expression constructs, can also be introduced in to plant cells using recombinant viruses. Plant cells can be transformed using viral vectors,
15 such as, e.g., tobacco mosaic virus derived vectors (Rouwendal (1997) Plant Mol. Biol. 33:989-999), see Porta (1996) "Use of viral replicons for the expression of genes in plants," Mol. Biotechnol. 5:209-221.

Alternatively, nucleic acids, e.g., an expression construct, can be combined with suitable T-DNA flanking regions and introduced into a conventional *Agrobacterium*
20 *tumefaciens* host vector. The virulence functions of the *Agrobacterium tumefaciens* host will direct the insertion of the construct and adjacent marker into the plant cell DNA when the cell is infected by the bacteria. *Agrobacterium tumefaciens*-mediated transformation techniques, including disarming and use of binary vectors, are well described in the scientific literature. See, e.g., Horsch (1984) *Science* 233:496-498;
25 Fraley (1983) *Proc. Natl. Acad. Sci. USA* 80:4803 (1983); *Gene Transfer to Plants*, Potrykus, ed. (Springer-Verlag, Berlin 1995). The DNA in an *A. tumefaciens* cell is contained in the bacterial chromosome as well as in another structure known as a Ti (tumor-inducing) plasmid. The Ti plasmid contains a stretch of DNA termed T-DNA (~20 kb long) that is transferred to the plant cell in the infection process and a series of vir
30 (virulence) genes that direct the infection process. *A. tumefaciens* can only infect a plant through wounds: when a plant root or stem is wounded it gives off certain chemical signals, in response to which, the vir genes of *A. tumefaciens* become activated and direct a series of events necessary for the transfer of the T-DNA from the Ti plasmid to the

plant's chromosome. The T-DNA then enters the plant cell through the wound. One speculation is that the T-DNA waits until the plant DNA is being replicated or transcribed, then inserts itself into the exposed plant DNA. In order to use *A. tumefaciens* as a transgene vector, the tumor-inducing section of T-DNA have to be removed, while retaining the T-DNA border regions and the vir genes. The transgene is then inserted between the T-DNA border regions, where it is transferred to the plant cell and becomes integrated into the plant's chromosomes.

The invention provides for the transformation of monocotyledonous plants using the nucleic acids of the invention, including important cereals, see Hiei (1997) Plant Mol. Biol. 35:205-218. See also, e.g., Horsch, Science (1984) 233:496; Fraley (1983) Proc. Natl Acad. Sci USA 80:4803; Thykjaer (1997) supra; Park (1996) Plant Mol. Biol. 32:1135-1148, discussing T-DNA integration into genomic DNA. See also D'Halluin, U.S. Patent No. 5,712,135, describing a process for the stable integration of a DNA comprising a gene that is functional in a cell of a cereal, or other monocotyledonous plant.

In one aspect, the third step can involve selection and regeneration of whole plants capable of transmitting the incorporated target gene to the next generation. Such regeneration techniques rely on manipulation of certain phytohormones in a tissue culture growth medium, typically relying on a biocide and/or herbicide marker that has been introduced together with the desired nucleotide sequences. Plant regeneration from cultured protoplasts is described in Evans et al., *Protoplasts Isolation and Culture, Handbook of Plant Cell Culture*, pp. 124-176, MacMillan Publishing Company, New York, 1983; and Binding, *Regeneration of Plants, Plant Protoplasts*, pp. 21-73, CRC Press, Boca Raton, 1985. Regeneration can also be obtained from plant callus, explants, organs, or parts thereof. Such regeneration techniques are described generally in Klee (1987) Ann. Rev. of Plant Phys. 38:467-486. To obtain whole plants from transgenic tissues such as immature embryos, they can be grown under controlled environmental conditions in a series of media containing nutrients and hormones, a process known as tissue culture. Once whole plants are generated and produce seed, evaluation of the progeny begins.

After the expression cassette is stably incorporated in transgenic plants, it can be introduced into other plants by sexual crossing. Any of a number of standard breeding techniques can be used, depending upon the species to be crossed. Since

transgenic expression of the nucleic acids of the invention leads to phenotypic changes, plants comprising the recombinant nucleic acids of the invention can be sexually crossed with a second plant to obtain a final product. Thus, the seed of the invention can be derived from a cross between two transgenic plants of the invention, or a cross between a
 5 plant of the invention and another plant. The desired effects (e.g., expression of the polypeptides of the invention to produce a plant in which flowering behavior is altered) can be enhanced when both parental plants express the polypeptides (e.g., an amylase, such as an alpha amylase) of the invention. The desired effects can be passed to future plant generations by standard propagation means.

10 The nucleic acids and polypeptides of the invention are expressed in or inserted in any plant or seed. Transgenic plants of the invention can be dicotyledonous or monocotyledonous. Examples of monocot transgenic plants of the invention are grasses, such as meadow grass (blue grass, *Poa*), forage grass such as festuca, lolium, temperate grass, such as *Agrostis*, and cereals, e.g., wheat, oats, rye, barley, rice, sorghum, and
 15 maize (corn). Examples of dicot transgenic plants of the invention are tobacco, legumes, such as lupins, potato, sugar beet, pea, bean and soybean, and cruciferous plants (family *Brassicaceae*), such as cauliflower, rape seed, and the closely related model organism *Arabidopsis thaliana*. Thus, the transgenic plants and seeds of the invention include a broad range of plants, including, but not limited to, species from the genera *Anacardium*,
 20 *Arachis*, *Asparagus*, *Atropa*, *Avena*, *Brassica*, *Citrus*, *Citrullus*, *Capsicum*, *Carthamus*, *Cocos*, *Coffea*, *Cucumis*, *Cucurbita*, *Daucus*, *Elaeis*, *Fragaria*, *Glycine*, *Gossypium*, *Helianthus*, *Heterocallis*, *Hordeum*, *Hyoscyamus*, *Lactuca*, *Linum*, *Lolium*, *Lupinus*, *Lycopersicon*, *Malus*, *Manihot*, *Majorana*, *Medicago*, *Nicotiana*, *Olea*, *Oryza*, *Panicum*, *Pennisetum*, *Persea*, *Phaseolus*, *Pistachia*, *Pisum*, *Pyrus*, *Prunus*, *Raphanus*, *Ricinus*,
 25 *Secale*, *Senecio*, *Sinapis*, *Solanum*, *Sorghum*, *Theobromus*, *Trigonella*, *Triticum*, *Vicia*, *Vitis*, *Vigna*, and *Zea*.

In alternative embodiments, the nucleic acids of the invention are expressed in plants which contain fiber cells, including, e.g., cotton, silk cotton tree (Kapok, *Ceiba pentandra*), desert willow, creosote bush, winterfat, balsa, ramie, kenaf,
 30 hemp, roselle, jute, sisal abaca and flax. In alternative embodiments, the transgenic plants of the invention can be members of the genus *Gossypium*, including members of any *Gossypium* species, such as *G. arboreum*; *G. herbaceum*, *G. barbadense*, and *G. hirsutum*.

The invention also provides for transgenic plants to be used for producing large amounts of the polypeptides (e.g., an amylase, such as an alpha amylase) of the invention. For example, see Palmgren (1997) Trends Genet. 13:348; Chong (1997) Transgenic Res. 6:289-296 (producing human milk protein beta-casein in transgenic potato plants using an auxin-inducible, bidirectional mannopine synthase (mas1',2') promoter with *Agrobacterium tumefaciens*-mediated leaf disc transformation methods).

Using known procedures, one of skill can screen for plants of the invention by detecting the increase or decrease of transgene mRNA or protein in transgenic plants. Means for detecting and quantitation of mRNAs or proteins are well known in the art.

10 Polypeptides and peptides

In one aspect, the invention provides isolated or recombinant polypeptides having a sequence identity (e.g., at least about 50%, 51%, 52%, 53%, 54%, 55%, 56%, 57%, 58%, 59%, 60%, 61%, 62%, 63%, 64%, 65%, 66%, 67%, 68%, 69%, 70%, 71%, 72%, 73%, 74%, 75%, 76%, 77%, 78%, 79%, 80%, 81%, 82%, 83%, 84%, 85%, 86%, 15 87%, 88%, 89%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or more, or complete (100%) sequence identity) to an exemplary sequence of the invention, e.g., SEQ ID NO:2, SEQ ID NO:4, SEQ ID NO:6, SEQ ID NO:10, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:16, SEQ ID NO:18, SEQ ID NO:20, SEQ ID NO:22, SEQ ID NO:24, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:30, SEQ ID NO:32, SEQ ID NO:34, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:40, SEQ ID NO:42, SEQ ID NO:44, SEQ ID NO:46, SEQ ID NO:48, SEQ ID NO:50, SEQ ID NO:52, SEQ ID NO:54, SEQ ID NO:56, SEQ ID NO:58, SEQ ID NO:60, SEQ ID NO:62, SEQ ID NO:64, SEQ ID NO:66, SEQ ID NO:68, SEQ ID NO:70, SEQ ID NO:72, SEQ ID NO:74, SEQ ID NO:76, SEQ ID NO:78, SEQ ID NO:80, SEQ ID NO:82, SEQ ID NO:84, SEQ ID NO:86, SEQ ID NO:88, SEQ ID NO:90, SEQ ID NO:92, SEQ ID NO:94, SEQ ID NO:96, SEQ ID NO:98, SEQ ID NO:100, SEQ ID NO:102, SEQ ID NO:104, SEQ ID NO:106, SEQ ID NO:108, SEQ ID NO:110, SEQ ID NO:112, SEQ ID NO:114, SEQ ID NO:116, SEQ ID NO:118, SEQ ID NO:120, SEQ ID NO:122, SEQ ID NO:124, SEQ ID NO:126, SEQ ID NO:128, SEQ ID NO:130, SEQ ID NO:132, SEQ ID NO:134, SEQ ID NO:136, SEQ ID NO:138, SEQ ID NO:140, SEQ ID NO:142, SEQ ID NO:144, SEQ ID NO:146, SEQ ID NO:148, SEQ ID NO:150, SEQ ID NO:152, SEQ ID NO:154, SEQ ID NO:156, SEQ ID NO:158, SEQ ID NO:160, SEQ ID NO:162, SEQ ID NO:164, SEQ ID

NO:166, SEQ ID NO:168, SEQ ID NO:190, SEQ ID NO:192, SEQ ID NO:194, SEQ ID
 NO:204, SEQ ID NO:206, SEQ ID NO:208, SEQ ID NO:210, SEQ ID NO:212, SEQ ID
 NO:323, SEQ ID NO:325, SEQ ID NO:327, SEQ ID NO:329, SEQ ID NO:331, SEQ ID
 NO:333, SEQ ID NO:335, SEQ ID NO:337, SEQ ID NO:339, SEQ ID NO:341, SEQ ID
 5 NO:343, SEQ ID NO:345, SEQ ID NO:347, SEQ ID NO:349, SEQ ID NO:351, SEQ ID
 NO:353, SEQ ID NO:355, SEQ ID NO:357, SEQ ID NO:359, SEQ ID NO:361, SEQ ID
 NO:363, SEQ ID NO:365, SEQ ID NO:367, SEQ ID NO:369, SEQ ID NO:371, SEQ ID
 NO:373, SEQ ID NO:375, SEQ ID NO:377, SEQ ID NO:379, SEQ ID NO:381, SEQ ID
 NO:383, SEQ ID NO:385, SEQ ID NO:387, SEQ ID NO:389, SEQ ID NO:391, SEQ ID
 10 NO:393, SEQ ID NO:395, SEQ ID NO:397, SEQ ID NO:399, SEQ ID NO:401, SEQ ID
 NO:403, SEQ ID NO:405, SEQ ID NO:407, SEQ ID NO:409, SEQ ID NO:411, SEQ ID
 NO:413, SEQ ID NO:415, SEQ ID NO:417, SEQ ID NO:419, SEQ ID NO:421, SEQ ID
 NO:423, SEQ ID NO:425, SEQ ID NO:427, SEQ ID NO:429, SEQ ID NO:431, SEQ ID
 NO:433, SEQ ID NO:435, SEQ ID NO:437, SEQ ID NO:439, SEQ ID NO:441, SEQ ID
 15 NO:443, SEQ ID NO:445, SEQ ID NO:447, SEQ ID NO:449, SEQ ID NO:451, SEQ ID
 NO:453, SEQ ID NO:455, SEQ ID NO:457, SEQ ID NO:459, SEQ ID NO:461, SEQ ID
 NO:461, SEQ ID NO:463, SEQ ID NO:464, SEQ ID NO:466, SEQ ID NO:468, SEQ ID
 NO:469, SEQ ID NO:470, SEQ ID NO:471, SEQ ID NO:472, SEQ ID NO:474, SEQ ID
 NO:476, SEQ ID NO:477, SEQ ID NO:479, SEQ ID NO:481, SEQ ID NO:482, SEQ ID
 20 NO:483, SEQ ID NO:485, SEQ ID NO:487, SEQ ID NO:488, SEQ ID NO:489, SEQ ID
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 NO:510, SEQ ID NO:512, SEQ ID NO:513, SEQ ID NO:514, SEQ ID NO:516, SEQ ID
 25 NO:518, SEQ ID NO:518, SEQ ID NO:520, SEQ ID NO:521, SEQ ID NO:523, SEQ ID
 NO:525, SEQ ID NO:526, SEQ ID NO:528, SEQ ID NO:530, SEQ ID NO:531, SEQ ID
 NO:533, SEQ ID NO:535, SEQ ID NO:536, SEQ ID NO:537, SEQ ID NO:538, SEQ ID
 NO:540, SEQ ID NO:542, SEQ ID NO:543, SEQ ID NO:545, SEQ ID NO:547, SEQ ID
 NO:548, SEQ ID NO:549, SEQ ID NO:550, SEQ ID NO:551, SEQ ID NO:553, SEQ ID
 30 NO:555, SEQ ID NO:556, SEQ ID NO:557, SEQ ID NO:559, SEQ ID NO:561, SEQ ID
 NO:562, SEQ ID NO:563, SEQ ID NO:564, SEQ ID NO:566, SEQ ID NO:568, SEQ ID
 NO:570, SEQ ID NO:572, SEQ ID NO:574, SEQ ID NO:576, SEQ ID NO:578, SEQ ID
 NO:580, SEQ ID NO:582, SEQ ID NO:584, SEQ ID NO:586, SEQ ID NO:588, SEQ ID

NO:589, SEQ ID NO:590, SEQ ID NO:591, SEQ ID NO:592, SEQ ID NO:594, SEQ ID NO:604, SEQ ID NO:606, SEQ ID NO:608, SEQ ID NO:610, SEQ ID NO:612, SEQ ID NO:614, SEQ ID NO:616, SEQ ID NO:618, SEQ ID NO:620 or SEQ ID NO:622, and subsequences thereof and variants thereof. In one aspect, the polypeptide has an amylase activity, e.g., an alpha amylase activity or a glucoamylase activity.

The identity can be over the full length of the polypeptide, or, the identity can be over a region of at least about 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 90, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700 or more residues. Polypeptides of the invention can also be shorter than the full length of exemplary polypeptides. In alternative aspects, the invention provides polypeptides (peptides, fragments) ranging in size between about 5 and the full length of a polypeptide, e.g., an enzyme, such as an amylase; exemplary sizes being of about 5, 10, 15, 20, 25, 30, 35, 40, 45, 50, 55, 60, 65, 70, 75, 80, 85, 90, 100, 125, 150, 175, 200, 250, 300, 350, 400, 450, 500, 550, 600, 650, 700, or more residues, e.g., contiguous residues of an exemplary amylase of the invention. Peptides of the invention can be useful as, e.g., labeling probes, antigens, toleragens, motifs, amylase active sites.

For example, the following table summarizes characteristics (e.g., activity, initial source, signal sequence location and exemplary signal sequence) of exemplary polypeptides of the invention. For example, the polypeptide having a sequence as set forth in SEQ ID NO:437, encoded by SEQ ID NO:436, was artificially generated; the polypeptide having a sequence as set forth in SEQ ID NO:439, encoded by SEQ ID NO:438, has amylase activity under alkaline conditions and was initially derived (isolated) from an unknown source; the polypeptide having a sequence as set forth in SEQ ID NO:441, encoded by SEQ ID NO:440, has amylase activity under alkaline conditions and was initially derived (isolated) from an unknown source, and has a signal sequence consisting of amino acid residues 1 to 32 of SEQ ID NO:441 ("AA 1-32"); see also discussion below regarding signal sequences of the invention, etc.:

SEQ ID NO:P	NOTES	Source	Signal location	Signal Sequence
436, 437	Reassembled amylase ALKALINE	Artificial		
438, 439	AMYLASE	Unknown		

440, 441	ALKALINE AMYLASE	Unknown	AA1-32	MNQIVNFKSHFYRKIALLSITFI WAAGSLSA
442, 443	ALKALINE AMYLASE	Unknown	AA1-27	MNRYLRLAALTALAPLAYPWG NLVRA
444, 445	ALKALINE AMYLASE	Unknown	AA1-24	MTPFGQPMMPGARMMAANMA PVRA
446, 447	ALKALINE AMYLASE	Unknown		
448, 449	ALKALINE AMYLASE	Unknown	AA1-23	MRLIMKKMIILITLAWVFTGCES
450, 451	ALKALINE AMYLASE	Unknown	AA1-49	MNDSINLYNFFPYNRPMSSINKTN TMKQMINWLGSLALLMLLLSCG EATE
452, 453	ALKALINE AMYLASE	Unknown	AA1-34	MMQLNPWFSTTLKAAGLATALA AVSACQPASESA
454, 455	ALKALINE AMYLASE	Unknown	AA1-37	MDLLEYKNTIQRRQTMTRKLL FIVATVILAVLVSFSS
456, 457	ALKALINE AMYLASE	Unknown	AA1-26	MMQLNPWFSSASLKAAGLATALA AVSA
458, 459	ALKALINE AMYLASE	Unknown	AA1-29	MFKVSLRSKDMKKLSLIVTILVLA LTLSA
460, 461	fungal	Cochliobolus heterostrophus ATCC 48331		
462-466	fungal	Fungal	AA1-22	MSRSSTILFVLAAANLASLVDA
467-474	fungal	Cochliobolus heterostrophus ATCC 48331		
475-479	fungal	Fungal		NOTE: AA1-122 may be removed and the remaining DNA/protein sequences still encode for an amylase
480-485	fungal	Fungal	AA1-19	MKFSLLATIVASISPLARA
486-493	fungal	Fungal	AA1-54	MRRKSTDKYKKVSIRAHLAACE QLAISKMLFSRTATILSLLCVQAT AISPRGSA
494-499	fungal	Fungal	AA1-22	MGFSKMLLGALIGIASLNGVQS
500-510	fungal	Fungal		
511-516	fungal	Fungal	AA1-21	MKYSIIPFVPLFAGLSRAASS MNMNIFLLIISLAFFSTVNCYTMS NA
517, 518	fungal	Fungal	AA1-26	
519-523	fungal	Fungal		
524-528	fungal	Fungal		

529-533	fungal	Cochliobolus heterostrophus ATCC 48331		
534-540	fungal	Cochliobolus heterostrophus ATCC 48331	AA1-20	MLLLNIFTTLFFYITCIVSA
541-545	fungal	Fungal		
546-553	fungal	Fungal	AA1-23	MASSLLSSLSSISTFNSTQILQA
554-559	fungal	Cochliobolus heterostrophus ATCC 48331	AA1-19	MTTALSSGQVAPTPHTAAA
560-566	fungal	Fungal	AA1-33	MLTTSEKRTSTAFVTWSMLWVV LLTSFVKDVHA
567, 568	ALKALINE AMYLASE	Unknown		
569, 570		Thermococcus alcaliphilus AEDII12RA		
571, 572		Unknown	AA1-28	MQSNGNVKGRSAVLALALLLT AVAATA
573, 574		Bacteria	AA1-27	MKKTFKLILVLMLSLTLVFGLTAP IQA
575, 576		Unknown		
577, 578		Unknown	AA1-34	MKPFLKKSIIITLLASTCLFTAWLI PSIAVPTVSA
579, 580		Unknown	AA1-29	MFKRRALGFLLAFLLVFTAVFGS MPMEFA
581, 582		Unknown	AA1-27	MKKFYKLTTALALSLSLALSLLG PAHA
583, 584		Unknown		
585, 586		Bacteria	AA1-28	MSLFKKSFPWILSLLLLFLFIAPF SIQT
587-594	GLUCOAMYLASE	Thermomyces lanuginosus ATCC 200065	AA1-23	MLFQPTLCAALGLAALIVQGGEA
603, 604		Unknown	AA1-31	MQNTAKNSIWQVRHSAIALSA LSLSFGLQA
605, 606		Unknown	AA1-34	MVNHLKKWIAGMALTLALLTGT VVPGLPVQVASA
607, 608		Unknown		

609, 610	Unknown	AA1-31	MQNTAKNSIWQRVRHSAIALSA
611, 612	Unknown		LSLSFGLQA
613, 614	Unknown	AA1-31	MQNTAKNSIWQRVRHSAIALSA
			LSLSFGLQA
615, 616	Unknown	AA1-34	MSERGVRRRAVRTALVGLAAAAT
			AAVTLGAPTAQA
617, 618	Unknown	AA1-27	MNRYLRLAALTALAPLAYPWG
			NLARA
619, 620	Bacteria	AA1-29	MARKSVAAALALVAGAAAVAVT
			GNTAAQA
621, 622	Unknown	AA1-31	MQNTAKNSIWQRVRHSAIALSA
			LSLSFGLQA

Polypeptides and peptides of the invention can be isolated from natural sources, be synthetic, or be recombinantly generated polypeptides. Peptides and proteins can be recombinantly expressed *in vitro* or *in vivo*. The peptides and polypeptides of the invention can be made and isolated using any method known in the art. Polypeptide and peptides of the invention can also be synthesized, whole or in part, using chemical methods well known in the art. See e.g., Caruthers (1980) Nucleic Acids Res. Symp. Ser. 215-223; Horn (1980) Nucleic Acids Res. Symp. Ser. 225-232; Banga, A.K., Therapeutic Peptides and Proteins, Formulation, Processing and Delivery Systems (1995) Technomic Publishing Co., Lancaster, PA. For example, peptide synthesis can be performed using various solid-phase techniques (see e.g., Roberge (1995) Science 269:202; Merrifield (1997) Methods Enzymol. 289:3-13) and automated synthesis may be achieved, e.g., using the ABI 431A Peptide Synthesizer (Perkin Elmer) in accordance with the instructions provided by the manufacturer.

The peptides and polypeptides of the invention can also be glycosylated. The glycosylation can be added post-translationally either chemically or by cellular biosynthetic mechanisms, wherein the later incorporates the use of known glycosylation motifs, which can be native to the sequence or can be added as a peptide or added in the nucleic acid coding sequence. The glycosylation can be O-linked or N-linked. Glycosylation can be added to any polypeptide of the invention to generate an enzyme that is more thermotolerant or thermostable than the "parent" enzyme (to which the

glycosylation was added). The glycosylation can be added by either chemical or by cellular biosynthetic mechanisms.

5 The invention provides amylases having a broad range of specific activity over a broad range of temperatures, e.g., at about 37°C in the range from about 10 to 10,000, or, 100 to about 1000 units per milligram of protein. Amylases of the invention can also have activity at temperatures as high as 120°C. In alternative aspects, the amylase used in these methods is active at these temperatures, e.g., active at temperatures in a range of between about 80°C to about 115°C, between about 100°C to about 110°C, and from about 105°C to about 108°C. However, amylases of the invention can also have 10 activity at low temperatures, e.g., as low as 4°C to 5°C.

The T_m of an enzyme of the invention can be shifted (for example, can be shifted between about 10°C to 90°C) by heat activation. For example, the T_m of SEQ ID NO:336/337 can be shifted about 17°C to 87°C by heat activation: for example, 80°C preincubation for 5 minutes.

15 The peptides and polypeptides of the invention, as defined above, include all "mimetic" and "peptidomimetic" forms. The terms "mimetic" and "peptidomimetic" refer to a synthetic chemical compound which has substantially the same structural and/or functional characteristics of the polypeptides of the invention. The mimetic can be either entirely composed of synthetic, non-natural analogues of amino acids, or, is a chimeric 20 molecule of partly natural peptide amino acids and partly non-natural analogs of amino acids. The mimetic can also incorporate any amount of natural amino acid conservative substitutions as long as such substitutions also do not substantially alter the mimetic's structure and/or activity. As with polypeptides of the invention which are conservative variants, routine experimentation will determine whether a mimetic is within the scope of 25 the invention, i.e., that its structure and/or function is not substantially altered. Thus, in one aspect, a mimetic composition is within the scope of the invention if it has an amylase activity.

Polypeptide mimetic compositions of the invention can contain any combination of non-natural structural components. In alternative aspect, mimetic 30 compositions of the invention include one or all of the following three structural groups: a) residue linkage groups other than the natural amide bond ("peptide bond") linkages; b) non-natural residues in place of naturally occurring amino acid residues; or c) residues which induce secondary structural mimicry, i.e., to induce or stabilize a secondary

structure, e.g., a beta turn, gamma turn, beta sheet, alpha helix conformation, and the like. For example, a polypeptide of the invention can be characterized as a mimetic when all or some of its residues are joined by chemical means other than natural peptide bonds. Individual peptidomimetic residues can be joined by peptide bonds, other chemical bonds
 5 or coupling means, such as, e.g., glutaraldehyde, N-hydroxysuccinimide esters, bifunctional maleimides, N,N'-dicyclohexylcarbodiimide (DCC) or N,N'-diisopropylcarbodiimide (DIC). Linking groups that can be an alternative to the traditional amide bond ("peptide bond") linkages include, e.g., ketomethylene (e.g., -C(=O)-CH₂- for -C(=O)-NH-), aminomethylene (CH₂-NH), ethylene, olefin (CH=CH),
 10 ether (CH₂-O), thioether (CH₂-S), tetrazole (CN₄-), thiazole, retroamide, thioamide, or ester (see, e.g., Spatola (1983) in Chemistry and Biochemistry of Amino Acids, Peptides and Proteins, Vol. 7, pp 267-357, "Peptide Backbone Modifications," Marcell Dekker, NY).

A polypeptide of the invention can also be characterized as a mimetic by
 15 containing all or some non-natural residues in place of naturally occurring amino acid residues. Non-natural residues are well described in the scientific and patent literature; a few exemplary non-natural compositions useful as mimetics of natural amino acid residues and guidelines are described below. Mimetics of aromatic amino acids can be generated by replacing by, e.g., D- or L- naphylalanine; D- or L- phenylglycine; D- or L-
 20 2 thieneylalanine; D- or L-1, -2, 3-, or 4- pyreneylalanine; D- or L-3 thieneylalanine; D- or L-(2-pyridinyl)-alanine; D- or L-(3-pyridinyl)-alanine; D- or L-(2-pyrazinyl)-alanine; D- or L-(4-isopropyl)-phenylglycine; D-(trifluoromethyl)-phenylglycine; D-(trifluoromethyl)-phenylalanine; D-p-fluoro-phenylalanine; D- or L-p-biphenylphenylalanine; D- or L-p-methoxy-biphenylphenylalanine; D- or L-2-
 25 indole(alkyl)alanines; and, D- or L-alkylainines, where alkyl can be substituted or unsubstituted methyl, ethyl, propyl, hexyl, butyl, pentyl, isopropyl, iso-butyl, sec-isotyl, iso-pentyl, or a non-acidic amino acids. Aromatic rings of a non-natural amino acid include, e.g., thiazolyl, thiophenyl, pyrazolyl, benzimidazolyl, naphthyl, furanyl, pyrrolyl, and pyridyl aromatic rings.

30 Mimetics of acidic amino acids can be generated by substitution by, e.g., non-carboxylate amino acids while maintaining a negative charge; (phosphono)alanine; sulfated threonine. Carboxyl side groups (e.g., aspartyl or glutamyl) can also be selectively modified by reaction with carbodiimides (R'-N-C-N-R') such as, e.g., 1-

cyclohexyl-3(2-morpholinyl-(4-ethyl) carbodiimide or 1-ethyl-3(4-azonia- 4,4-dimetholpentyl) carbodiimide. Aspartyl or glutamyl can also be converted to asparaginy and glutaminy residues by reaction with ammonium ions. Mimetics of basic amino acids can be generated by substitution with, e.g., (in addition to lysine and arginine) the amino acids ornithine, citrulline, or (guanidino)-acetic acid, or (guanidino)alkyl-acetic acid, where alkyl is defined above. Nitrile derivative (e.g., containing the CN-moiety in place of COOH) can be substituted for asparagine or glutamine. Asparaginy and glutaminy residues can be deaminated to the corresponding aspartyl or glutamyl residues. Arginine residue mimetics can be generated by reacting arginyl with, e.g., one or more conventional reagents, including, e.g., phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, or ninhydrin, preferably under alkaline conditions. Tyrosine residue mimetics can be generated by reacting tyrosyl with, e.g., aromatic diazonium compounds or tetranitromethane. N-acetylimidizol and tetranitromethane can be used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively. Cysteine residue mimetics can be generated by reacting cysteinyl residues with, e.g., alpha-haloacetates such as 2-chloroacetic acid or chloroacetamide and corresponding amines; to give carboxymethyl or carboxyamidomethyl derivatives. Cysteine residue mimetics can also be generated by reacting cysteinyl residues with, e.g., bromo-trifluoroacetone, alpha-bromo-beta-(5-imidozoyl) propionic acid; chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide; methyl 2-pyridyl disulfide; p-chloromercuribenzoate; 2-chloromercuri-4-nitrophenol; or, chloro-7-nitrobenzo-oxa-1,3-diazole. Lysine mimetics can be generated (and amino terminal residues can be altered) by reacting lysinyl with, e.g., succinic or other carboxylic acid anhydrides. Lysine and other alpha-amino-containing residue mimetics can also be generated by reaction with imidoesters, such as methyl picolinimide, pyridoxal phosphate, pyridoxal, chloroborohydride, trinitrobenzenesulfonic acid, O-methylisourea, 2,4, pentanedione, and transamidase-catalyzed reactions with glyoxylate. Mimetics of methionine can be generated by reaction with, e.g., methionine sulfoxide. Mimetics of proline include, e.g., pipecolic acid, thiazolidine carboxylic acid, 3- or 4- hydroxy proline, dehydropyrolidine, 3- or 4-methylproline, or 3,3,-dimethylproline. Histidine residue mimetics can be generated by reacting histidyl with, e.g., diethylprocarbonate or para-bromophenacyl bromide. Other mimetics include, e.g., those generated by hydroxylation of proline and lysine; phosphorylation of the hydroxyl groups of seryl or threonyl residues; methylation of the alpha-amino groups of lysine,

arginine and histidine; acetylation of the N-terminal amine; methylation of main chain amide residues or substitution with N-methyl amino acids; or amidation of C-terminal carboxyl groups.

5 A residue, e.g., an amino acid, of a polypeptide of the invention can also be replaced by an amino acid (or peptidomimetic residue) of the opposite chirality. Thus, any amino acid naturally occurring in the L-configuration (which can also be referred to as the R or S, depending upon the structure of the chemical entity) can be replaced with the amino acid of the same chemical structural type or a peptidomimetic, but of the opposite chirality, referred to as the D- amino acid, but also can be referred to as the R- or
10 S- form.

The invention also provides methods for modifying the polypeptides of the invention by either natural processes, such as post-translational processing (e.g., phosphorylation, acylation, etc), or by chemical modification techniques, and the resulting modified polypeptides. Modifications can occur anywhere in the polypeptide,
15 including the peptide backbone, the amino acid side-chains and the amino or carboxyl termini. It will be appreciated that the same type of modification may be present in the same or varying degrees at several sites in a given polypeptide. Also a given polypeptide may have many types of modifications. Modifications include acetylation, acylation, ADP-ribosylation, amidation, covalent attachment of flavin, covalent attachment of a
20 heme moiety, covalent attachment of a nucleotide or nucleotide derivative, covalent attachment of a lipid or lipid derivative, covalent attachment of a phosphatidylinositol, cross-linking cyclization, disulfide bond formation, demethylation, formation of covalent cross-links, formation of cysteine, formation of pyroglutamate, formylation, gamma-carboxylation, glycosylation, GPI anchor formation, hydroxylation, iodination,
25 methylation, myristoylation, oxidation, pegylation, proteolytic processing, phosphorylation, prenylation, racemization, selenoylation, sulfation, and transfer-RNA mediated addition of amino acids to protein such as arginylation. See, e.g., Creighton, T.E., *Proteins – Structure and Molecular Properties* 2nd Ed., W.H. Freeman and Company, New York (1993); *Posttranslational Covalent Modification of Proteins*, B.C.
30 Johnson, Ed., Academic Press, New York, pp. 1-12 (1983).

Solid-phase chemical peptide synthesis methods can also be used to synthesize the polypeptide or fragments of the invention. Such method have been known in the art since the early 1960's (Merrifield, R. B., *J. Am. Chem. Soc.*, 85:2149-2154,

1963) (See also Stewart, J. M. and Young, J. D., Solid Phase Peptide Synthesis, 2nd Ed.,
Pierce Chemical Co., Rockford, Ill., pp. 11-12)) and have recently been employed in
commercially available laboratory peptide design and synthesis kits (Cambridge Research
Biochemicals). Such commercially available laboratory kits have generally utilized the
5 teachings of H. M. Geysen et al, Proc. Natl. Acad. Sci., USA, 81:3998 (1984) and provide
for synthesizing peptides upon the tips of a multitude of "rods" or "pins" all of which are
connected to a single plate. When such a system is utilized, a plate of rods or pins is
inverted and inserted into a second plate of corresponding wells or reservoirs, which
contain solutions for attaching or anchoring an appropriate amino acid to the pin's or rod's
10 tips. By repeating such a process step, i.e., inverting and inserting the rod's and pin's tips
into appropriate solutions, amino acids are built into desired peptides. In addition, a
number of available Fmoc peptide synthesis systems are available. For example,
assembly of a polypeptide or fragment can be carried out on a solid support using an
Applied Biosystems, Inc. Model 431A™ automated peptide synthesizer. Such equipment
15 provides ready access to the peptides of the invention, either by direct synthesis or by
synthesis of a series of fragments that can be coupled using other known techniques.

The invention provides novel amylases (e.g., alpha amylases), including
the exemplary enzymes of the invention, nucleic acids encoding them, antibodies that
bind them, and methods for making and using them. In one aspect, the polypeptides of
20 the invention have an amylase activity, as described herein, including, e.g., the ability to
hydrolyze starches into sugars. In one aspect, the polypeptides of the invention have an
alpha amylase activity. In alternative aspects, the amylases of the invention have
activities that have been modified from those of the exemplary amylases described herein.

The invention includes amylases of the invention with and without signal
25 sequences (including signal sequences of the invention, see e.g., Table 3, below, or other
signal sequences) and the signal sequences themselves (e.g., Table 3, below). The
invention also include polypeptides (e.g., fusion proteins) comprising a signal sequence of
the invention, see, e.g., Table 3, below. The polypeptide comprising a signal sequence of
the invention can be an amylase of the invention or another amylase or another enzyme or
30 other polypeptide.

The invention includes immobilized amylases, anti-amylase antibodies and
fragments thereof. The invention provides methods for inhibiting amylase activity, e.g.,
using dominant negative mutants or anti-amylase antibodies of the invention. The

invention includes heterocomplexes, e.g., fusion proteins, heterodimers, etc., comprising the amylases of the invention.

In one aspect, amylases (e.g., alpha amylases) of the invention hydrolyze internal polysaccharide bonds, e.g., α -1,4- and 1,6-glucosidic bonds in starch to produce smaller molecular weight maltodextrines. In one aspect, this hydrolysis is largely at random. Thus, the invention provides methods for producing smaller molecular weight maltodextrines.

Amylases of the invention can be used in laboratory and industrial settings to hydrolyze starch or any maltodextrine-comprising compound for a variety of purposes. These amylases can be used alone to provide specific hydrolysis or can be combined with other amylases to provide a "cocktail" with a broad spectrum of activity. Exemplary uses include the removal or partial or complete hydrolysis of starch or any maltodextrine-comprising compound from biological, food, animal feed, pharmaceutical or industrial samples.

For example, the amylases of the present invention can be formulated in laundry detergents to aid in the removal of starch-containing stains. In one aspect, the invention provides detergents comprising amylases of the invention, including amylases active under alkaline conditions, and methods of making and using them. These detergent compositions include laundry and dishwashing (e.g., autodishwashing) solutions and application. Amylases of the invention can be used as cleaning agents in any detergent matrices (see industrial applications below). The amylases of the present invention can be used in the initial stages (liquefaction) of starch processing, in wet corn milling, in alcohol production, in the textile industry for starch desizing, in baking applications, in the beverage industry, in oilfields in drilling processes; in inking of recycled paper; and in animal feed.

Amylases of the invention can have an amylase activity under various conditions, e.g., extremes in pH and/or temperature, oxidizing agents, and the like. The invention provides methods leading to alternative amylase preparations with different catalytic efficiencies and stabilities, e.g., towards temperature, oxidizing agents and changing wash conditions. In one aspect, amylase variants can be produced using techniques of site-directed mutagenesis and/or random mutagenesis. In one aspect, directed evolution can be used to produce a great variety of amylase variants with alternative specificities and stability.

The proteins of the invention are also useful as research reagents to identify amylase modulators, e.g., activators or inhibitors of amylase activity. Briefly, test samples (compounds, broths, extracts, and the like) are added to amylase assays to determine their ability to inhibit substrate cleavage. Inhibitors identified in this way can be used in industry and research to reduce or prevent undesired proteolysis. As with amylases, inhibitors can be combined to increase the spectrum of activity.

The invention also provides methods of discovering new amylases using the nucleic acids, polypeptides and antibodies of the invention. In one aspect, lambda phage libraries are screened for expression-based discovery of amylases. In one aspect, the invention uses lambda phage libraries in screening to allow detection of toxic clones; improved access to substrate; reduced need for engineering a host, by-passing the potential for any bias resulting from mass excision of the library; and, faster growth at low clone densities. Screening of lambda phage libraries can be in liquid phase or in solid phase. In one aspect, the invention provides screening in liquid phase. This gives a greater flexibility in assay conditions; additional substrate flexibility; higher sensitivity for weak clones; and ease of automation over solid phase screening.

The invention provides screening methods using the proteins and nucleic acids of the invention and robotic automation to enable the execution of many thousands of biocatalytic reactions and screening assays in a short period of time, e.g., per day, as well as ensuring a high level of accuracy and reproducibility (see discussion of arrays, below). As a result, a library of derivative compounds can be produced in a matter of weeks. For further teachings on modification of molecules, including small molecules, see PCT/US94/09174.

The present invention includes amylase enzymes which are non-naturally occurring carbonyl hydrolase variants (e.g., amylase variants) having a different proteolytic activity, stability, substrate specificity, pH profile and/or performance characteristic as compared to the precursor carbonyl hydrolase from which the amino acid sequence of the variant is derived. Specifically, such amylase variants have an amino acid sequence not found in nature, which is derived by substitution of a plurality of amino acid residues of a precursor amylase with different amino acids. The precursor amylase may be a naturally-occurring amylase or a recombinant amylase. The useful amylase variants encompass the substitution of any of the naturally occurring L-amino acids at the designated amino acid residue positions.

Amylase Signal Sequences

The invention provides signal sequences consisting of or comprising a peptide having a sequence comprising residues 1 to 12, 1 to 13, 1 to 14, 1 to 15, 1 to 16, 1 to 17, 1 to 18, 1 to 19, 1 to 20, 1 to 21, 1 to 22, 1 to 23, 1 to 24, 1 to 25, 1 to 26, 1 to 27, 1 to 28, 1 to 28, 1 to 30 or 1 to 31, 1 to 32, 1 to 33, 1 to 34, 1 to 35, 1 to 36, 1 to 37, 1 to 38, or 1 to 39, or longer, of a polypeptide of the invention. For example, the invention provides amylase (e.g., alpha amylase or glucoamylase) signal sequences and nucleic acids encoding these signal sequences, e.g., exemplary peptides of the invention having sequences as set forth in Table 3, SEQ ID NO:7, SEQ ID NO:8, and SEQ ID NO:213 through 257, and polypeptides comprising (or consisting of) sequences as set forth in Table 3, SEQ ID NO:7, SEQ ID NO:8, and SEQ ID NO:213 through 257. The invention also provides amylase signal sequences and nucleic acids encoding these signal sequences, e.g., peptides comprising or consisting of residues 1 to 27 of SEQ ID NO:323 (encoded by SEQ ID NO:322), peptides comprising or consisting of residues 1 to 22 of SEQ ID NO:333 (encoded by SEQ ID NO:332), peptides comprising or consisting of residues 1 to 20 of SEQ ID NO:335 (encoded by SEQ ID NO:334), peptides comprising or consisting of residues 1 to 35 of SEQ ID NO:337 (encoded by SEQ ID NO:336), etc., see Table 3 for, in addition to these signal sequences, additional amylase signal sequences and nucleic acids encoding these signal sequences.

The invention also provides amylase signal sequences and nucleic acids encoding these signal sequences comprising or consisting of residues 1 to 32 or 1 to 33 of SEQ ID NO:441; residues 1 to 27 or 1 to 28 of SEQ ID NO:443; residues 1 to 24 or 1 to 25 of SEQ ID NO:445; residues 1 to 23 or 1 to 24 of SEQ ID NO:449; residues 1 to 49 or 1 to 50 of SEQ ID NO:451; residues 1 to 34 or 1 to 35 of SEQ ID NO:453; residues 1 to 37 or 1 to 38 of SEQ ID NO:455; residues 1 to 26 or 1 to 27 of SEQ ID NO:457; residues 1 to 29 or 1 to 30 of SEQ ID NO:459; residues 1 to 22 or 1 to 23 of SEQ ID NO:466; residues 1 to 19 or 1 to 20 of SEQ ID NO:485; residues 1 to 54 or 1 to 55 of SEQ ID NO:493; residues 1 to 22 or 1 to 23 of SEQ ID NO:499; residues 21 or 1 to 22 of SEQ ID NO:516; residues 1 to 26 or 1 to 27 of SEQ ID NO:518; residues 1 to 20 or 1 to 21 of SEQ ID NO:540; residues 1 to 23 or 1 to 24 of SEQ ID NO:553; residues 1 to 19 or 1 to 20 of SEQ ID NO:559; residues 1 to 33 or 1 to 34 of SEQ ID NO:566.

For example, regarding Table 3, the invention provides peptides comprising or consisting of amino acid residues 1 to 23 (SEQ ID NO:213) of SEQ ID NO:87, etc.

Table 3

<u>SEQ ID NO.</u>	<u>Signal Sequence</u>
SEQ ID NO: 87	AA1-23 (SEQ ID NO:213)
SEQ ID NO: 91	AA1-23 (SEQ ID NO: 214)
SEQ ID NO: 93	AA1-33 (SEQ ID NO: 215)
SEQ ID NO: 97	AA1-31 (SEQ ID NO: 216)
SEQ ID NO: 99	AA1-30 (SEQ ID NO: 217)
SEQ ID NO: 103	AA1-22 (SEQ ID NO: 218)
SEQ ID NO: 105	AA1-33 (SEQ ID NO: 219)
SEQ ID NO: 109	AA1-25 (SEQ ID NO: 220)
SEQ ID NO: 111	AA1-35 (SEQ ID NO: 221)
SEQ ID NO: 113	AA1-28 (SEQ ID NO: 222)
SEQ ID NO: 117	AA1-21 (SEQ ID NO: 223)
SEQ ID NO: 119	AA1-30 (SEQ ID NO: 224)
SEQ ID NO: 123	AA1-35 (SEQ ID NO: 225)
SEQ ID NO: 125	AA1-28 (SEQ ID NO: 226)
SEQ ID NO: 127	AA1-30 (SEQ ID NO: 227)
SEQ ID NO: 131	AA1-30 (SEQ ID NO: 228)
SEQ ID NO: 133	AA1-30 (SEQ ID NO: 229)
SEQ ID NO: 137	AA1-28 (SEQ ID NO: 230)
SEQ ID NO: 139	AA1-23 (SEQ ID NO: 231)
SEQ ID NO: 141	AA1-23 (SEQ ID NO: 232)
SEQ ID NO: 143	AA1-30 (SEQ ID NO: 233)
SEQ ID NO: 145	AA1-27 (SEQ ID NO: 234)
SEQ ID NO: 147	AA1-29 (SEQ ID NO: 235)
SEQ ID NO: 149	AA1-28 (SEQ ID NO: 236)
SEQ ID NO: 69	AA1-27 (SEQ ID NO: 237)
SEQ ID NO: 153	AA1-26 (SEQ ID NO: 238)
SEQ ID NO: 155	AA1-33 (SEQ ID NO: 239)
SEQ ID NO: 157	AA1-25 (SEQ ID NO: 240)
SEQ ID NO: 159	AA1-25 (SEQ ID NO: 241)
SEQ ID NO: 161	AA1-36 (SEQ ID NO: 242)
SEQ ID NO: 167	AA1-36 (SEQ ID NO: 243)
SEQ ID NO: 169	AA1-23 (SEQ ID NO: 244)
SEQ ID NO: 173	AA1-25 (SEQ ID NO: 245)

SEQ ID NO: 175	AA1-22 (SEQ ID NO: 246)
SEQ ID NO: 177	AA1-23 (SEQ ID NO: 247)
SEQ ID NO: 179	AA1-23 (SEQ ID NO: 248)
SEQ ID NO: 185	AA1-25 (SEQ ID NO: 249)
SEQ ID NO: 189	AA1-36 (SEQ ID NO: 250)
SEQ ID NO: 191	AA1-25 (SEQ ID NO: 251)
SEQ ID NO: 193	AA1-25 (SEQ ID NO: 252)
SEQ ID NO: 197	AA1-23 (SEQ ID NO: 253)
SEQ ID NO: 199	AA1-23 (SEQ ID NO: 254)
SEQ ID NO: 201	AA1-30 (SEQ ID NO: 255)
SEQ ID NO: 203	AA1-25 (SEQ ID NO: 256)
SEQ ID NO: 205	AA1-16 (SEQ ID NO: 257)
SEQ ID NO: 73	AA1-16 (SEQ ID NO: 7)
SEQ ID NO: 79	AA1-26 (SEQ ID NO: 8)
SEQ ID NO: 322, 323	Residues 1 through 27
SEQ ID NO: 332, 333	Residues 1 through 22
SEQ ID NO: 334, 335	Residues 1 through 20
SEQ ID NO: 336, 337	Residues 1 through 35
SEQ ID NO: 338, 339	Residues 1 through 50
SEQ ID NO: 342, 343	Residues 1 through 23
SEQ ID NO: 344, 345	Residues 1 through 22
SEQ ID NO: 346, 347	Residues 1 through 21
SEQ ID NO: 350, 351	Residues 1 through 21
SEQ ID NO: 352, 353	Residues 1 through 27
SEQ ID NO: 354, 355	Residues 1 through 24
SEQ ID NO: 358, 359	Residues 1 through 29
SEQ ID NO: 362, 363	Residues 1 through 20
SEQ ID NO: 364, 365	Residues 1 through 29
SEQ ID NO: 366, 367	Residues 1 through 24
SEQ ID NO: 370, 371	Residues 1 through 22
SEQ ID NO: 372, 373	Residues 1 through 25
SEQ ID NO: 374, 375	Residues 1 through 21
SEQ ID NO: 376, 377	Residues 1 through 37
SEQ ID NO: 378, 379	Residues 1 through 27
SEQ ID NO: 380, 381	Residues 1 through 29
SEQ ID NO: 382, 383	Residues 1 through 35
SEQ ID NO: 384, 385	Residues 1 through 37
SEQ ID NO: 386, 387	Residues 1 through 25
SEQ ID NO: 388, 389	Residues 1 through 21

SEQ ID NO:390, 391	Residues 1 through 58
SEQ ID NO:394, 395	Residues 1 through 57
SEQ ID NO:396, 397	Residues 1 through 19
SEQ ID NO:400, 401	Residues 1 through 19
SEQ ID NO:402, 403	Residues 1 through 19
SEQ ID NO:404, 405	Residues 1 through 26
SEQ ID NO:406, 407	Residues 1 through 21
SEQ ID NO:408, 409	Residues 1 through 51
SEQ ID NO:410, 411	Residues 1 through 21
SEQ ID NO:416, 417	Residues 1 through 24
SEQ ID NO:418, 419	Residues 1 through 44
SEQ ID NO:420, 421	Residues 1 through 44
SEQ ID NO:422, 423	Residues 1 through 27
SEQ ID NO:424, 425	Residues 1 through 37
SEQ ID NO:428, 429	Residues 1 through 30
SEQ ID NO:430, 431	Residues 1 through 33
SEQ ID NO:432, 433	Residues 1 through 34
SEQ ID NO:434, 435	Residues 1 through 27

The amylase signal sequences of the invention can be isolated peptides, or, sequences joined to another amylase or a non-amylase polypeptide, e.g., as a fusion protein. In one aspect, the invention provides polypeptides comprising amylase signal sequences of the invention. In one aspect, polypeptides comprising amylase signal sequences of the invention comprise sequences heterologous to an amylase of the invention (e.g., a fusion protein comprising an amylase signal sequence of the invention and sequences from another amylase or a non-amylase protein). In one aspect, the invention provides amylases of the invention with heterologous signal sequences, e.g., sequences with a yeast signal sequence. For example, an amylase of the invention comprising a heterologous signal sequence in a vectors, e.g., a pPIC series vector (Invitrogen, Carlsbad, CA).

In one aspect, the signal sequences of the invention are identified following identification of novel amylase polypeptides. The pathways by which proteins are sorted and transported to their proper cellular location are often referred to as protein targeting pathways. One of the most important elements in all of these targeting systems is a short amino acid sequence at the amino terminus of a newly synthesized polypeptide called the signal sequence. This signal sequence directs a protein to its appropriate

location in the cell and is removed during transport or when the protein reaches its final destination. Most lysosomal, membrane, or secreted proteins have an amino-terminal signal sequence that marks them for translocation into the lumen of the endoplasmic reticulum. More than 100 signal sequences for proteins in this group have been
5 determined. The signal sequences can vary in length from 13 to 36 amino acid residues. Various methods of recognition of signal sequences are known to those of skill in the art. For example, in one aspect, novel amylase signal peptides are identified by a method referred to as SignalP. SignalP uses a combined neural network which recognizes both signal peptides and their cleavage sites. (Nielsen, et al., "Identification of prokaryotic and
10 eukaryotic signal peptides and prediction of their cleavage sites." Protein Engineering, vol. 10, no. 1, p. 1-6 (1997).

It should be understood that in some aspects amylases of the invention may not have signal sequences. In one aspect, the invention provides the amylases of the invention lacking all or part of a signal sequence, e.g. the signal sequences of the
15 invention (see Table 3, below). In one aspect, the invention provides a nucleic acid sequence encoding a signal sequence from one amylase operably linked to a nucleic acid sequence of a different amylase or, optionally, a signal sequence from a non-amylase protein may be desired. Table 3 shows exemplary signal sequences of the invention.

20

Amylase prepro and signal sequences and catalytic domains

In addition to signal sequences (e.g., signal peptides (SPs)), as discussed above, the invention provides prepro domains and catalytic domains (CDs). The SPs, prepro domains and/or CDs of the invention can be isolated or recombinant peptides or
25 can be part of a fusion protein, e.g., as a heterologous domain in a chimeric protein. The invention provides nucleic acids encoding these catalytic domains (CDs) (e.g., "active sites"), prepro domains and signal sequences (SPs, e.g., a peptide having a sequence comprising/ consisting of amino terminal residues of a polypeptide of the invention).

The amylase signal sequences (SPs), catalytic domains (CDs) and/or
30 prepro sequences of the invention can be isolated peptides, or, sequences joined to another amylase or a non- amylase polypeptide, e.g., as a fusion (chimeric) protein. In one aspect, polypeptides comprising amylase signal sequences SPs and/or prepro of the

invention comprise sequences heterologous to amylases of the invention (e.g., a fusion protein comprising an SP and/or prepro of the invention and sequences from another amylase or a non- amylase protein). In one aspect, the invention provides amylases of the invention with heterologous CDs, SPs and/or prepro sequences, e.g., sequences with a yeast signal sequence. An amylase of the invention can comprise a heterologous CD, SP and/or prepro in a vector, e.g., a pPIC series vector (Invitrogen, Carlsbad, CA).

In one aspect, SPs, CDs, and/or prepro sequences of the invention are identified following identification of novel amylase polypeptides. The pathways by which proteins are sorted and transported to their proper cellular location are often referred to as protein targeting pathways. One of the most important elements in all of these targeting systems is a short amino acid sequence at the amino terminus of a newly synthesized polypeptide called the signal sequence. This signal sequence directs a protein to its appropriate location in the cell and is removed during transport or when the protein reaches its final destination. Most lysosomal, membrane, or secreted proteins have an amino-terminal signal sequence that marks them for translocation into the lumen of the endoplasmic reticulum. The signal sequences can vary in length from 13 to 45 or more amino acid residues. Various methods of recognition of signal sequences are known to those of skill in the art. For example, in one aspect, novel hydrolase signal peptides are identified by a method referred to as SignalP. SignalP uses a combined neural network which recognizes both signal peptides and their cleavage sites. (Nielsen, et al., "Identification of prokaryotic and eukaryotic signal peptides and prediction of their cleavage sites." Protein Engineering, vol. 10, no. 1, p. 1-6 (1997)).

In some aspects, an amylase of the invention may not have SPs and/or prepro sequences, and/or catalytic domains (CDs). In one aspect, the invention provides amylases lacking all or part of an SP, a CD and/or a prepro domain. In one aspect, the invention provides a nucleic acid sequence encoding a signal sequence (SP), a CD and/or prepro from one amylase operably linked to a nucleic acid sequence of a different amylase or, optionally, a signal sequence (SPs), a CD and/or prepro domain from a non-amylase protein may be desired.

The invention also provides isolated or recombinant polypeptides comprising signal sequences (SPs), prepro domain and/or catalytic domains (CDs) of the invention and heterologous sequences. The heterologous sequences are sequences not naturally associated (e.g., to an amylase) with an SP, prepro domain and/or CD. The

sequence to which the SP, prepro domain and/or CD are not naturally associated can be on the SP's, prepro domain and/or CD's amino terminal end, carboxy terminal end, and/or on both ends of the SP and/or CD. In one aspect, the invention provides an isolated or recombinant polypeptide comprising (or consisting of) a polypeptide comprising a signal sequence (SP), prepro domain and/or catalytic domain (CD) of the invention with the proviso that it is not associated with any sequence to which it is naturally associated (e.g., amylase sequence). Similarly in one aspect, the invention provides isolated or recombinant nucleic acids encoding these polypeptides. Thus, in one aspect, the isolated or recombinant nucleic acid of the invention comprises coding sequence for a signal sequence (SP), prepro domain and/or catalytic domain (CD) of the invention and a heterologous sequence (i.e., a sequence not naturally associated with the a signal sequence (SP), prepro domain and/or catalytic domain (CD) of the invention). The heterologous sequence can be on the 3' terminal end, 5' terminal end, and/or on both ends of the SP, prepro domain and/or CD coding sequence.

The polypeptides of the invention include amylases in an active or inactive form. For example, the polypeptides of the invention include proproteins before "maturation" or processing of prepro sequences, e.g., by a proprotein-processing enzyme, such as a proprotein convertase to generate an "active" mature protein. The polypeptides of the invention include amylases inactive for other reasons, e.g., before "activation" by a post-translational processing event, e.g., an endo- or exo-peptidase or proteinase action, a phosphorylation event, an amidation, a glycosylation or a sulfation, a dimerization event, and the like. Methods for identifying "prepro" domain sequences, CDs, and signal sequences are well known in the art, see, e.g., Van de Ven (1993) Crit. Rev. Oncog. 4(2):115-136. For example, to identify a prepro sequence, the protein is purified from the extracellular space and the N-terminal protein sequence is determined and compared to the unprocessed form.

The polypeptides of the invention include all active forms, including active subsequences, e.g., catalytic domains (CDs) or active sites, of an enzyme of the invention. In one aspect, the invention provides catalytic domains or active sites as set forth below.

In one aspect, the invention provides a peptide or polypeptide comprising or consisting of an active site domain as predicted through use of a database such as Pfam (which is a large collection of multiple sequence alignments and hidden Markov models covering many common protein families, The Pfam protein families database, A. Bateman, E.

Birney, L. Cerruti, R. Durbin, L. Etwiller, S.R. Eddy, S. Griffiths-Jones, K.L. Howe, M. Marshall, and E.L.L. Sonnhammer, Nucleic Acids Research, 30(1):276-280, 2002) or equivalent.

Hybrid amylases and peptide libraries

5 In one aspect, the invention provides hybrid amylases and fusion proteins, including peptide libraries, comprising sequences of the invention. The peptide libraries of the invention can be used to isolate peptide modulators (e.g., activators or inhibitors) of targets, such as amylase substrates, receptors, enzymes. The peptide libraries of the invention can be used to identify formal binding partners of targets, such as ligands, e.g.,
10 cytokines, hormones and the like.

In one aspect, the fusion proteins of the invention (e.g., the peptide moiety) are conformationally stabilized (relative to linear peptides) to allow a higher binding affinity for targets. The invention provides fusions of amylases of the invention and other peptides, including known and random peptides. They can be fused in such a manner that
15 the structure of the amylases is not significantly perturbed and the peptide is metabolically or structurally conformationally stabilized. This allows the creation of a peptide library that is easily monitored both for its presence within cells and its quantity.

Amino acid sequence variants of the invention can be characterized by a predetermined nature of the variation, a feature that sets them apart from a naturally
20 occurring form, e.g., an allelic or interspecies variation of an amylase sequence. In one aspect, the variants of the invention exhibit the same qualitative biological activity as the naturally occurring analogue. Alternatively, the variants can be selected for having modified characteristics. In one aspect, while the site or region for introducing an amino acid sequence variation is predetermined, the mutation per se need not be predetermined.

25 For example, in order to optimize the performance of a mutation at a given site, random mutagenesis may be conducted at the target codon or region and the expressed amylase variants screened for the optimal combination of desired activity. Techniques for making substitution mutations at predetermined sites in DNA having a known sequence are well known, as discussed herein for example, M13 primer mutagenesis and PCR mutagenesis.
30 Screening of the mutants can be done using assays of proteolytic activities. In alternative aspects, amino acid substitutions can be single residues; insertions can be on the order of from about 1 to 20 amino acids, although considerably larger insertions can be done.

Deletions can range from about 1 to about 20, 30, 40, 50, 60, 70 residues or more. To obtain a final derivative with the optimal properties, substitutions, deletions, insertions or any combination thereof may be used. Generally, these changes are done on a few amino acids to minimize the alteration of the molecule. However, larger changes may be
5 tolerated in certain circumstances.

The invention provides amylases where the structure of the polypeptide backbone, the secondary or the tertiary structure, e.g., an alpha-helical or beta-sheet structure, has been modified. In one aspect, the charge or hydrophobicity has been modified. In one aspect, the bulk of a side chain has been modified. Substantial changes
10 in function or immunological identity are made by selecting substitutions that are less conservative. For example, substitutions can be made which more significantly affect: the structure of the polypeptide backbone in the area of the alteration, for example a alpha-helical or a beta-sheet structure; a charge or a hydrophobic site of the molecule, which can be at an active site; or a side chain. The invention provides substitutions in
15 polypeptide of the invention where (a) a hydrophilic residues, e.g. seryl or threonyl, is substituted for (or by) a hydrophobic residue, e.g. leucyl, isoleucyl, phenylalanyl, valyl or alanyl; (b) a cysteine or proline is substituted for (or by) any other residue; (c) a residue having an electropositive side chain, e.g. lysyl, arginyl, or histidyl, is substituted for (or by) an electronegative residue, e.g. glutamyl or aspartyl; or (d) a residue having a bulky
20 side chain, e.g. phenylalanine, is substituted for (or by) one not having a side chain, e.g. glycine. The variants can exhibit the same qualitative biological activity (i.e. amylase activity) although variants can be selected to modify the characteristics of the amylases as needed.

In one aspect, amylases of the invention comprise epitopes or purification
25 tags, signal sequences or other fusion sequences, etc. In one aspect, the amylases of the invention can be fused to a random peptide to form a fusion polypeptide. By "fused" or "operably linked" herein is meant that the random peptide and the amylase are linked together, in such a manner as to minimize the disruption to the stability of the amylase structure, e.g., it retains amylase activity. The fusion polypeptide (or fusion
30 polynucleotide encoding the fusion polypeptide) can comprise further components as well, including multiple peptides at multiple loops.

In one aspect, the peptides and nucleic acids encoding them are randomized, either fully randomized or they are biased in their randomization, e.g. in

nucleotide/residue frequency generally or per position. "Randomized" means that each nucleic acid and peptide consists of essentially random nucleotides and amino acids, respectively. In one aspect, the nucleic acids which give rise to the peptides can be chemically synthesized, and thus may incorporate any nucleotide at any position. Thus, when the nucleic acids are expressed to form peptides, any amino acid residue may be incorporated at any position. The synthetic process can be designed to generate randomized nucleic acids, to allow the formation of all or most of the possible combinations over the length of the nucleic acid, thus forming a library of randomized nucleic acids. The library can provide a sufficiently structurally diverse population of randomized expression products to affect a probabilistically sufficient range of cellular responses to provide one or more cells exhibiting a desired response. Thus, the invention provides an interaction library large enough so that at least one of its members will have a structure that gives it affinity for some molecule, protein, or other factor.

Screening Methodologies and "On-line" Monitoring Devices

In practicing the methods of the invention, a variety of apparatus and methodologies can be used to in conjunction with the polypeptides and nucleic acids of the invention, e.g., to screen polypeptides for amylase activity, to screen compounds as potential modulators, e.g., activators or inhibitors, of an amylase activity, for antibodies that bind to a polypeptide of the invention, for nucleic acids that hybridize to a nucleic acid of the invention, to screen for cells expressing a polypeptide of the invention and the like.

Capillary Arrays

Capillary arrays, such as the GIGAMATRIX™, Diversa Corporation, San Diego, CA, can be used to in the methods of the invention. Nucleic acids or polypeptides of the invention can be immobilized to or applied to an array, including capillary arrays. Arrays can be used to screen for or monitor libraries of compositions (e.g., small molecules, antibodies, nucleic acids, etc.) for their ability to bind to or modulate the activity of a nucleic acid or a polypeptide of the invention. Capillary arrays provide another system for holding and screening samples. For example, a sample screening apparatus can include a plurality of capillaries formed into an array of adjacent capillaries, wherein each capillary comprises at least one wall defining a lumen for retaining a sample. The apparatus can further include interstitial material disposed

between adjacent capillaries in the array, and one or more reference indicia formed within of the interstitial material. A capillary for screening a sample, wherein the capillary is adapted for being bound in an array of capillaries, can include a first wall defining a lumen for retaining the sample, and a second wall formed of a filtering material, for
5 filtering excitation energy provided to the lumen to excite the sample.

A polypeptide or nucleic acid, e.g., a ligand, can be introduced into a first component into at least a portion of a capillary of a capillary array. Each capillary of the capillary array can comprise at least one wall defining a lumen for retaining the first component. An air bubble can be introduced into the capillary behind the first component. A second
10 component can be introduced into the capillary, wherein the second component is separated from the first component by the air bubble. A sample of interest can be introduced as a first liquid labeled with a detectable particle into a capillary of a capillary array, wherein each capillary of the capillary array comprises at least one wall defining a lumen for retaining the first liquid and the detectable particle, and wherein the at least one
15 wall is coated with a binding material for binding the detectable particle to the at least one wall. The method can further include removing the first liquid from the capillary tube, wherein the bound detectable particle is maintained within the capillary, and introducing a second liquid into the capillary tube.

The capillary array can include a plurality of individual capillaries comprising at least one
20 outer wall defining a lumen. The outer wall of the capillary can be one or more walls fused together. Similarly, the wall can define a lumen that is cylindrical, square, hexagonal or any other geometric shape so long as the walls form a lumen for retention of a liquid or sample. The capillaries of the capillary array can be held together in close proximity to form a planar structure. The capillaries can be bound together, by being
25 fused (e.g., where the capillaries are made of glass), glued, bonded, or clamped side-by-side. The capillary array can be formed of any number of individual capillaries, for example, a range from 100 to 4,000,000 capillaries. A capillary array can form a micro titer plate having about 100,000 or more individual capillaries bound together.

Arrays, or "Biochips"

30 Nucleic acids or polypeptides of the invention can be immobilized to or applied to an array. Arrays can be used to screen for or monitor libraries of compositions (e.g., small molecules, antibodies, nucleic acids, etc.) for their ability to bind to or modulate the activity of a nucleic acid or a polypeptide of the invention. For example, in

one aspect of the invention, a monitored parameter is transcript expression of an amylase gene. One or more, or, all the transcripts of a cell can be measured by hybridization of a sample comprising transcripts of the cell, or, nucleic acids representative of or complementary to transcripts of a cell, by hybridization to immobilized nucleic acids on an array, or "biochip." By using an "array" of nucleic acids on a microchip, some or all of the transcripts of a cell can be simultaneously quantified. Alternatively, arrays comprising genomic nucleic acid can also be used to determine the genotype of a newly engineered strain made by the methods of the invention. Polypeptide arrays" can also be used to simultaneously quantify a plurality of proteins. The present invention can be practiced with any known "array," also referred to as a "microarray" or "nucleic acid array" or "polypeptide array" or "antibody array" or "biochip," or variation thereof. Arrays are generically a plurality of "spots" or "target elements," each target element comprising a defined amount of one or more biological molecules, e.g., oligonucleotides, immobilized onto a defined area of a substrate surface for specific binding to a sample molecule, e.g., mRNA transcripts.

In practicing the methods of the invention, any known array and/or method of making and using arrays can be incorporated in whole or in part, or variations thereof, as described, for example, in U.S. Patent Nos. 6,277,628; 6,277,489; 6,261,776; 6,258,606; 6,054,270; 6,048,695; 6,045,996; 6,022,963; 6,013,440; 5,965,452; 5,959,098; 5,856,174; 5,830,645; 5,770,456; 5,632,957; 5,556,752; 5,143,854; 5,807,522; 5,800,992; 5,744,305; 5,700,637; 5,556,752; 5,434,049; see also, e.g., WO 99/51773; WO 99/09217; WO 97/46313; WO 96/17958; see also, e.g., Johnston (1998) Curr. Biol. 8:R171-R174; Schummer (1997) Biotechniques 23:1087-1092; Kern (1997) Biotechniques 23:120-124; Solinas-Toldo (1997) Genes, Chromosomes & Cancer 20:399-407; Bowtell (1999) Nature Genetics Supp. 21:25-32. See also published U.S. patent applications Nos. 20010018642; 20010019827; 20010016322; 20010014449; 20010014448; 20010012537; 20010008765.

Antibodies and Antibody-based screening methods

The invention provides isolated or recombinant antibodies that specifically bind to an amylase of the invention. These antibodies can be used to isolate, identify or quantify the amylases of the invention or related polypeptides. These antibodies can be used to isolate other polypeptides within the scope the invention or other related

amylases. The antibodies can be designed to bind to an active site of an amylase. Thus, the invention provides methods of inhibiting amylases using the antibodies of the invention.

The antibodies can be used in immunoprecipitation, staining, immunoaffinity columns, and the like. If desired, nucleic acid sequences encoding for specific antigens can be generated by immunization followed by isolation of polypeptide or nucleic acid, amplification or cloning and immobilization of polypeptide onto an array of the invention. Alternatively, the methods of the invention can be used to modify the structure of an antibody produced by a cell to be modified, e.g., an antibody's affinity can be increased or decreased. Furthermore, the ability to make or modify antibodies can be a phenotype engineered into a cell by the methods of the invention.

Methods of immunization, producing and isolating antibodies (polyclonal and monoclonal) are known to those of skill in the art and described in the scientific and patent literature, see, e.g., Coligan, CURRENT PROTOCOLS IN IMMUNOLOGY, Wiley/Greene, NY (1991); Stites (eds.) BASIC AND CLINICAL IMMUNOLOGY (7th ed.) Lange Medical Publications, Los Altos, CA ("Stites"); Goding, MONOCLONAL ANTIBODIES: PRINCIPLES AND PRACTICE (2d ed.) Academic Press, New York, NY (1986); Kohler (1975) Nature 256:495; Harlow (1988) ANTIBODIES, A LABORATORY MANUAL, Cold Spring Harbor Publications, New York. Antibodies also can be generated in vitro, e.g., using recombinant antibody binding site expressing phage display libraries, in addition to the traditional in vivo methods using animals. See, e.g., Hoogenboom (1997) Trends Biotechnol. 15:62-70; Katz (1997) Annu. Rev. Biophys. Biomol. Struct. 26:27-45.

Polypeptides or peptides can be used to generate antibodies which bind specifically to the polypeptides, e.g., the amylases, of the invention. The resulting antibodies may be used in immunoaffinity chromatography procedures to isolate or purify the polypeptide or to determine whether the polypeptide is present in a biological sample. In such procedures, a protein preparation, such as an extract, or a biological sample is contacted with an antibody capable of specifically binding to one of the polypeptides of the invention.

In immunoaffinity procedures, the antibody is attached to a solid support, such as a bead or other column matrix. The protein preparation is placed in contact with the antibody under conditions in which the antibody specifically binds to one of the

polypeptides of the invention. After a wash to remove non-specifically bound proteins, the specifically bound polypeptides are eluted.

The ability of proteins in a biological sample to bind to the antibody may be determined using any of a variety of procedures familiar to those skilled in the art. For example, binding may be determined by labeling the antibody with a detectable label such as a fluorescent agent, an enzymatic label, or a radioisotope. Alternatively, binding of the antibody to the sample may be detected using a secondary antibody having such a detectable label thereon. Particular assays include ELISA assays, sandwich assays, radioimmunoassays, and Western Blots.

Polyclonal antibodies generated against the polypeptides of the invention can be obtained by direct injection of the polypeptides into an animal or by administering the polypeptides to a non-human animal. The antibody so obtained will then bind the polypeptide itself. In this manner, even a sequence encoding only a fragment of the polypeptide can be used to generate antibodies which may bind to the whole native polypeptide. Such antibodies can then be used to isolate the polypeptide from cells expressing that polypeptide.

For preparation of monoclonal antibodies, any technique which provides antibodies produced by continuous cell line cultures can be used. Examples include the hybridoma technique, the trioma technique, the human B-cell hybridoma technique, and the EBV-hybridoma technique (see, e.g., Cole (1985) in *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc., pp. 77-96).

Techniques described for the production of single chain antibodies (see, e.g., U.S. Patent No. 4,946,778) can be adapted to produce single chain antibodies to the polypeptides of the invention. Alternatively, transgenic mice may be used to express humanized antibodies to these polypeptides or fragments thereof.

Antibodies generated against the polypeptides of the invention may be used in screening for similar polypeptides (e.g., amylases) from other organisms and samples. In such techniques, polypeptides from the organism are contacted with the antibody and those polypeptides which specifically bind the antibody are detected. Any of the procedures described above may be used to detect antibody binding.

Kits

The invention provides kits comprising the compositions, e.g., nucleic acids, expression cassettes, vectors, cells, transgenic seeds or plants or plant parts, polypeptides (e.g., amylases) and/or antibodies of the invention. The kits also can contain instructional material teaching the methodologies and industrial uses of the invention, as
 5 described herein.

Measuring Metabolic Parameters

The methods of the invention provide whole cell evolution, or whole cell engineering, of a cell to develop a new cell strain having a new phenotype, e.g., a new or modified amylase activity, by modifying the genetic composition of the cell. The genetic
 10 composition can be modified by addition to the cell of a nucleic acid of the invention. To detect the new phenotype, at least one metabolic parameter of a modified cell is monitored in the cell in a "real time" or "on-line" time frame. In one aspect, a plurality of cells, such as a cell culture, is monitored in "real time" or "on-line." In one aspect, a plurality of metabolic parameters is monitored in "real time" or "on-line." Metabolic
 15 parameters can be monitored using the amylases of the invention.

Metabolic flux analysis (MFA) is based on a known biochemistry framework. A linearly independent metabolic matrix is constructed based on the law of mass conservation and on the pseudo-steady state hypothesis (PSSH) on the intracellular metabolites. In practicing the methods of the invention, metabolic networks are
 20 established, including the:

- identity of all pathway substrates, products and intermediary metabolites
- identity of all the chemical reactions interconverting the pathway metabolites, the stoichiometry of the pathway reactions,
- identity of all the enzymes catalyzing the reactions, the enzyme reaction kinetics,
- 25 • the regulatory interactions between pathway components, e.g. allosteric interactions, enzyme-enzyme interactions etc,
- intracellular compartmentalization of enzymes or any other supramolecular organization of the enzymes, and,
- the presence of any concentration gradients of metabolites, enzymes or effector
 30 molecules or diffusion barriers to their movement.

Once the metabolic network for a given strain is built, mathematic presentation by matrix notion can be introduced to estimate the intracellular metabolic

fluxes if the on-line metabolome data is available. Metabolic phenotype relies on the changes of the whole metabolic network within a cell. Metabolic phenotype relies on the change of pathway utilization with respect to environmental conditions, genetic regulation, developmental state and the genotype, etc. In one aspect of the methods of the invention, after the on-line MFA calculation, the dynamic behavior of the cells, their phenotype and other properties are analyzed by investigating the pathway utilization. For example, if the glucose supply is increased and the oxygen decreased during the yeast fermentation, the utilization of respiratory pathways will be reduced and/or stopped, and the utilization of the fermentative pathways will dominate. Control of physiological state of cell cultures will become possible after the pathway analysis. The methods of the invention can help determine how to manipulate the fermentation by determining how to change the substrate supply, temperature, use of inducers, etc. to control the physiological state of cells to move along desirable direction. In practicing the methods of the invention, the MFA results can also be compared with transcriptome and proteome data to design experiments and protocols for metabolic engineering or gene shuffling, etc.

In practicing the methods of the invention, any modified or new phenotype can be conferred and detected, including new or improved characteristics in the cell. Any aspect of metabolism or growth can be monitored.

Monitoring expression of an mRNA transcript

In one aspect of the invention, the engineered phenotype comprises increasing or decreasing the expression of an mRNA transcript (e.g., an amylase message) or generating new (e.g., amylase) transcripts in a cell. This increased or decreased expression can be traced by testing for the presence of an amylase of the invention or by amylase activity assays. mRNA transcripts, or messages, also can be detected and quantified by any method known in the art, including, e.g., Northern blots, quantitative amplification reactions, hybridization to arrays, and the like. Quantitative amplification reactions include, e.g., quantitative PCR, including, e.g., quantitative reverse transcription polymerase chain reaction, or RT-PCR; quantitative real time RT-PCR, or "real-time kinetic RT-PCR" (see, e.g., Kreuzer (2001) Br. J. Haematol. 114:313-318; Xia (2001) Transplantation 72:907-914).

In one aspect of the invention, the engineered phenotype is generated by knocking out expression of a homologous gene. The gene's coding sequence or one or more

transcriptional control elements can be knocked out, e.g., promoters or enhancers. Thus, the expression of a transcript can be completely ablated or only decreased.

In one aspect of the invention, the engineered phenotype comprises increasing the expression of a homologous gene. This can be effected by knocking out of
5 a negative control element, including a transcriptional regulatory element acting in cis- or trans- , or, mutagenizing a positive control element. One or more, or, all the transcripts of a cell can be measured by hybridization of a sample comprising transcripts of the cell, or, nucleic acids representative of or complementary to transcripts of a cell, by hybridization to immobilized nucleic acids on an array.

10 *Monitoring expression of a polypeptides, peptides and amino acids*

In one aspect of the invention, the engineered phenotype comprises increasing or decreasing the expression of a polypeptide (e.g., an amylase) or generating new polypeptides in a cell. This increased or decreased expression can be traced by determining the amount of amylase present or by amylase activity assays. Polypeptides,
15 peptides and amino acids also can be detected and quantified by any method known in the art, including, e.g., nuclear magnetic resonance (NMR), spectrophotometry, radiography (protein radiolabeling), electrophoresis, capillary electrophoresis, high performance liquid chromatography (HPLC), thin layer chromatography (TLC), hyperdiffusion chromatography, various immunological methods, e.g. immunoprecipitation,
20 immunodiffusion, immuno-electrophoresis, radioimmunoassays (RIAs), enzyme-linked immunosorbent assays (ELISAs), immuno-fluorescent assays, gel electrophoresis (e.g., SDS-PAGE), staining with antibodies, fluorescent activated cell sorter (FACS), pyrolysis mass spectrometry, Fourier-Transform Infrared Spectrometry, Raman spectrometry, GC-MS, and LC-Electrospray and cap-LC-tandem-electrospray mass spectrometries, and the
25 like. Novel bioactivities can also be screened using methods, or variations thereof, described in U.S. Patent No. 6,057,103. Furthermore, as discussed below in detail, one or more, or, all the polypeptides of a cell can be measured using a protein array.

Industrial Applications

Detergent Compositions

30 The invention provides detergent compositions comprising one or more polypeptides of the invention, for example, amylases of the invention, such as alpha amylases, glucoamylases, etc., and methods of making and using these compositions.

The invention incorporates all methods of making and using detergent compositions, see, e.g., U.S. Patent No. 6,413,928; 6,399,561; 6,365,561; 6,380,147. The detergent compositions can be a one and two part aqueous composition, a non-aqueous liquid composition, a cast solid, a granular form, a particulate form, a compressed tablet, a gel and/or a paste and a slurry form. The invention also provides methods capable of a rapid removal of gross food soils, films of food residue and other minor food compositions using these detergent compositions. Amylases of the invention can facilitate the removal of starchy stains by means of catalytic hydrolysis of the starch polysaccharide. Amylases of the invention can be used in dishwashing detergents in textile laundering detergents.

10 The actual active enzyme content depends upon the method of manufacture of a detergent composition and is not critical, assuming the detergent solution has the desired enzymatic activity. In one aspect, the amount of amylase present in the final solution ranges from about 0.001 mg to 0.5 mg per gram of the detergent composition. The particular enzyme chosen for use in the process and products of this invention depends upon the conditions of final utility, including the physical product form, use pH, use temperature, and soil types to be degraded or altered. The enzyme can be chosen to provide optimum activity and stability for any given set of utility conditions. In one aspect, the polypeptides of the present invention are active in the pH ranges of from about 4 to about 12 and in the temperature range of from about 20°C to about 95°C.

20 The detergents of the invention can comprise cationic, semi-polar nonionic or zwitterionic surfactants; or, mixtures thereof.

Amylases of the present invention can be formulated into powdered and liquid detergents having pH between 4.0 and 12.0 at levels of about 0.01 to about 5% (preferably 0.1% to 0.5%) by weight. These detergent compositions can also include other enzymes such as known proteases, cellulases, lipases or endoglycosidases, as well as builders and stabilizers. The addition of amylases of the invention to conventional cleaning compositions does not create any special use limitation. In other words, any temperature and pH suitable for the detergent is also suitable for the present compositions as long as the pH is within the above range, and the temperature is below the described enzyme's denaturing temperature. In addition, the polypeptides of the invention can be used in a cleaning composition without detergents, again either alone or in combination with builders and stabilizers.

The present invention provides cleaning compositions including detergent compositions for cleaning hard surfaces, detergent compositions for cleaning fabrics, dishwashing compositions, oral cleaning compositions, denture cleaning compositions, and contact lens cleaning solutions.

5 In one aspect, the invention provides a method for washing an object comprising contacting the object with a polypeptide of the invention under conditions sufficient for washing. In one aspect, a polypeptide of the invention (e.g., an alkaline-active amylase) is used in a detergent, i.e., as a detergent additive. The detergent composition of the invention may, for example, be formulated as a hand or machine
10 laundry detergent composition comprising a polypeptide of the invention. Detergent compositions of the invention include laundry and dishwashing (e.g., autodishwashing) solutions and application. A laundry additive suitable for pre-treatment of stained fabrics can comprise a polypeptide of the invention. A fabric softener composition can comprise a polypeptide of the invention. Alternatively, a polypeptide of the invention can be
15 formulated as a detergent composition for use in general household hard surface cleaning operations. In alternative aspects, detergent additives and detergent compositions of the invention may comprise one or more other enzymes such as a protease, a lipase, a cutinase, another amylase, a carbohydrase, a cellulase, a pectinase, a mannanase, an arabinase, a galactanase, a xylanase, an oxidase, e.g., a lactase, and/or a peroxidase. The
20 properties of the enzyme(s) of the invention are chosen to be compatible with the selected detergent (i.e. pH-optimum, compatibility with other enzymatic and non-enzymatic ingredients, etc.) and the enzyme(s) is present in effective amounts. In one aspect, amylase enzymes of the invention are used to remove malodorous materials from fabrics. Various detergent compositions and methods for making them that can be used in
25 practicing the invention are described in, e.g., U.S. Patent Nos. 6,333,301; 6,329,333; 6,326,341; 6,297,038; 6,309,871; 6,204,232; 6,197,070; 5,856,164.

Treating fabrics

The invention provides methods of treating fabrics using one or more polypeptides of the invention. The polypeptides of the invention can be used in any
30 fabric-treating method, which are well known in the art, see, e.g., U.S. Patent No. 6,077,316. For example, in one aspect, the feel and appearance of a fabric is improved by

a method comprising contacting the fabric with an amylase of the invention in a solution. In one aspect, the fabric is treated with the solution under pressure.

In one aspect, the enzymes of the invention are applied during or after the weaving of textiles, or during the desizing stage, or one or more additional fabric processing steps. During the weaving of textiles, the threads are exposed to considerable mechanical strain. Prior to weaving on mechanical looms, warp yarns are often coated with sizing starch or starch derivatives in order to increase their tensile strength and to prevent breaking. The enzymes of the invention can be applied to remove these sizing starch or starch derivatives. After the textiles have been woven, a fabric can proceed to a desizing stage. This can be followed by one or more additional fabric processing steps. Desizing is the act of removing size from textiles. After weaving, the size coating must be removed before further processing the fabric in order to ensure a homogeneous and wash-proof result. The invention provides a method of desizing comprising enzymatic hydrolysis of the size by the action of an enzyme of the invention.

The enzymes of the invention can be used to desize fabrics, including cotton-containing fabrics, as detergent additives, e.g., in aqueous compositions. The invention provides methods for producing a stonewashed look on indigo-dyed denim fabric and garments. For the manufacture of clothes, the fabric can be cut and sewn into clothes or garments, which is afterwards finished. In particular, for the manufacture of denim jeans, different enzymatic finishing methods have been developed. The finishing of denim garment normally is initiated with an enzymatic desizing step, during which garments are subjected to the action of amylolytic enzymes in order to provide softness to the fabric and make the cotton more accessible to the subsequent enzymatic finishing steps. The invention provides methods of finishing denim garments (e.g., a "bio-stoning process"), enzymatic desizing and providing softness to fabrics using the amylases of the invention. The invention provides methods for quickly softening denim garments in a desizing and/or finishing process.

Foods and food processing: liquification of starch

The enzymes of the invention have numerous applications in food processing industry. The amylases of the invention are used in starch to fructose processing. Starch to fructose processing can consist of four steps: liquefaction of

granular starch, saccharification of the liquefied starch into dextrose, purification, and isomerization to fructose.

The invention provides methods of starch liquefaction using the enzymes of the invention. Concentrated suspensions of starch polymer granules are converted into a solution of soluble shorter chain length dextrans of low viscosity. This step is useful for convenient handling with standard equipment and for efficient conversion to glucose or 10^3 other sugars. In one aspect, the granular starch is liquefied by gelatinizing the granules by raising the temperature of the granular starch to over about 72°C. The heating process instantaneously disrupts the insoluble starch granules to produce a water soluble starch solution. The solubilized starch solution can then be liquefied by an amylase of the invention. Thus, the invention provides enzymatic starch liquefaction processes using an amylase of the invention.

Figure 26, Figure 27 and Figure 28 illustrate alternative exemplary starch processes, including starch liquefaction processes, of the invention (using at least one enzyme of the invention). For example, Figure 26 illustrates an exemplary starch liquefaction process of the invention comprising treating a starch slurry (e.g., having about 30% to 35% solids) with steam for primary liquefaction (e.g., at about 105°C for about 5 minutes), input into a flash tank, followed by secondary liquefaction (e.g., at about 90°C to 95°C for about 90 minutes), each or one of these steps involving use of an enzyme of the invention. Figure 27 illustrates another exemplary starch liquefaction process of the invention comprising treating a starch slurry at about between pH 4 to pH 5, e.g., pH 4.5, adjusting the pH, calcium addition, liquefaction at about pH 5 to pH 6, e.g., pH 5.4, at about 95°C using an alpha amylase of the invention, followed by another pH and temperature adjustment for saccharification at about between pH 4 to pH 5, e.g., pH 4.5, at a temperature of between about 60°C to 65°C using a glucoamylase of the invention. Figure 28 illustrates another exemplary starch process of the invention comprising treating a starch slurry at about between pH 4 to pH 5, e.g., pH 4.5, (optional adjusting pH, calcium addition), combined liquefaction-saccharification using an alpha amylase and/or a glucoamylase of the invention at about between pH 4 to pH 5, e.g., pH 4.5, at a temperature of greater than about 90°C, or, greater than about 95°C, followed by another pH and temperature adjustment for saccharification at about between pH 4 to pH 5, e.g., pH 4.5, at a temperature of between about 60°C to 65°C using a glucoamylase of the invention. In one aspect, the combined liquefaction-saccharification of the invention

is a "one-pot" process. In one aspect, the entire process is a "one-pot" process. Any one of these processes, and any one of these steps, can also comprise, or can further comprise, another enzyme of the invention (e.g., a glucosidase such as an α -1,6-glucosidase, a maltase, etc.), or another enzyme such as a pullulanase or an isomerase.

5 An exemplary enzymatic liquefaction process involves adjusting the pH of a granular starch slurry to between 6.0 and 6.5 and the addition of calcium hydroxide, sodium hydroxide or sodium carbonate. In one aspect, calcium hydroxide is added. This provides calcium ions to stabilize the glucoamylase of the invention against inactivation. In one aspect, upon addition of amylase, the suspension is pumped through a steam jet to
10 instantaneously raise the temperature to between 80°-115°C. In one aspect, the starch is immediately gelatinized and, due to the presence of amylase, depolymerized through random hydrolysis of α -1,4-glycosidic bonds by amylase to a fluid mass. The fluid mass can be easily pumped.

 The invention provides various enzymatic starch liquefaction processes
15 using an amylase of the invention. In one aspect of the liquefaction process of the invention, an amylase is added to the starch suspension and the suspension is held at a temperature of between about 80°-100°C to partially hydrolyze the starch granules. In one aspect, the partially hydrolyzed starch suspension is pumped through a jet at temperatures in excess of about 105°C to thoroughly gelatinize any remaining granular
20 structure. In one aspect, after cooling the gelatinized starch, a second addition of amylase is made to further hydrolyze the starch.

 The invention provides enzymes and processes for hydrolyzing liquid (liquefied) and granular starch. Such starch can be derived from any source, e.g., corn, wheat, milo, sorghum, rye or bulgher. The invention applies to any grain starch source
25 which is useful in liquefaction, e.g., any other grain or vegetable source known to produce starch suitable for liquefaction. The methods of the invention comprise liquefying starch from any natural material, such as rice, germinated rice, corn, barley, milo, wheat, legumes and sweet potato. The liquefying process can substantially hydrolyze the starch to produce a syrup. The temperature range of the liquefaction can be any liquefaction
30 temperature which is known to be effective in liquefying starch. For example, the temperature of the starch can be between about 80°C to about 115°C, between about 100°C to about 110°C, and from about 105°C to about 108°C. In alternative aspects, the amylase used in these methods is active at these temperatures, e.g., active at temperatures

in a range of between about 80°C to about 115°C, between about 100°C to about 110°C, and from about 105°C to about 108°C.

The invention provides methods for liquefaction saccharification as illustrated in Figure 17. In one aspect, alpha-amylases of the invention are used in the illustrated liquefaction step (some current industrial methods use *B. licheniformis* a-
5 amylase). In one aspect, the process takes place at about pH 6.0 at a temperature anywhere in the range of between about 95°C to 105°C, for a length of time anywhere between about 0.5 and 5 hours, e.g., 60, 90 or 120 minutes. In one aspect, in a corn steep process, prior to liquefaction cellulases, proteases and/or protein thio reductases are added.

10 In one aspect of a liquefaction process of the invention, an amylase of the invention that has activity at about pH 4.5 (or, anywhere between about pH 5 and pH 5), that may or may not be Ca^{2+} dependent is added. Eliminating the addition of salts in the front end of the process eliminates the need to remove them at the back end of the process. In one aspect of a liquefaction process of the invention, an amylase that is more
15 active is used. This can allow one to decrease the amount of enzyme needed. In one aspect, liquefaction and saccharification are done in the same pot, as a "one-pot process," for example, under conditions comprising about 90°C to 95°C (or, anywhere between about 80°C to 105°C), as about a 3 hour process (or, as a process lasting between about 1 and 5 hours). In this aspect, the enzyme load can be cut in half again.

20 In one aspect of a saccharification process of the invention, a glucoamylase of the invention is used. In one aspect, glucoamylases of the invention are used in the illustrated saccharification step (some current industrial methods use *A. niger* glucoamylase). In one aspect, the process takes place at about pH 4.5, in a temperature range of between about 60°C to 62°C (or, anywhere in the range of between about 50°C
25 to 72°C, or, between about 40°C to 80°C) as a process lasting between about 12 and 96 or more hours. In one aspect of a saccharification process of the invention, a glucoamylase of the invention is used to give a higher level of dextrose in the syrup. In one aspect, other enzymes are added, e.g., pullulanases to increase the amount of glucose.

In one aspect, amylases of the invention are used in the illustrated
30 isomerization step (some current industrial methods use *Streptomyces sp.* glucose isomerase). In one aspect, the isomerization reaction of the invention takes place under conditions comprising anywhere between about pH 5 and pH 10, or anywhere between about pH 6 and pH 9, or anywhere between about pH 7.0 and 8.5. In one aspect, the

isomerization reaction of the invention takes place under conditions comprising between about 40°C to 75°C, or between about 50°C to 65°C, or between about 55°C to 60°C.

In one aspect of an isomerization process of the invention, a xylose isomerase is used. In one aspect, cobalt is used in the reaction (some known thermostable glucose isomerases require cobalt). In one aspect, an enzyme of the invention is used that lacks dependency, or has less dependency, on cobalt. In one aspect, an enzyme of the invention is used that has activity at a lower pH, e.g., pH 7.0, pH 6.5, pH 6, pH 5.5, pH 5, pH 4.5, pH 4, pH 3.5 or less, or, e.g., between a range of about pH 3.5 to 7.0). In one aspect, this allows less color formation (otherwise, excess color may have to be removed).
10 In one aspect, the temperature is increased during isomerization, e.g. to between about 80°C to 110°C, 85°C to 105°C, or 90°C to 100°C. This can increase the amount of fructose produced, e.g. to about 51%. However, in one aspect, for sodas (e.g., soft drinks and the like), the fructose level can be anywhere between about 45% and 65%, or 50% and 60%, e.g., about 55%.

15 In one aspect, one, some or all of the enzymes used in processes of the invention (including the enzymes of the invention) are immobilized, e.g., immobilized on any surface, e.g., a flat surface or an enzyme column, e.g., immobilized on an array, a bead, fiber, pore, capillary and the like. In one aspect, by being immobilized, they can be reused.

20 In one aspect, the invention provides “enzyme cocktails” using at least one enzyme of the invention. In one aspect, “enzyme cocktails” are used in the processes of the invention, e.g., including the liquefaction saccharification methods as illustrated in Figure 17. For example, in one aspect, cell wall degrading enzymes (CWDE) are used, e.g., for textile, pulp and paper, and laundry processes of the invention, including, e.g.,
25 combinations of cellulases, hemicellulases, xylanase, galactomannanases, glucomannanases, arabinofuranosidases, and others. In one aspect, “enzyme cocktails” used in the processes of the invention for bio-bleaching (e.g., pulp and paper, laundry processes), include combinations of laccases, peroxidases, oxidases and the like. In one aspect, cell wall degrading enzymes are combined with bio-bleaching enzymes and
30 enzymes of the invention to degrade plant cell walls to release color agents.

Processes to produce high MW dextrose syrups

The invention provides processes to produce high MW dextrose syrups using enzymes of the invention, including methods for producing oligosaccharides having a MW tightly groups at about 20,000 MW. In one aspect, amylases of the invention of
5 archael origin, including the archael-derived amylases of SEQ ID NO:80 (encoded by SEQ ID NO:79), SEQ ID NO:82 (encoded by SEQ ID NO:81), SEQ ID NO:116 (encoded by SEQ ID NO:115), SEQ ID NO:323 (encoded by SEQ ID NO:322), SEQ NO:570 (encoded by SEQ ID NO:169) and enzymes of the invention having the same activity as these archael amylases, are used to liquefy a starch-comprising composition, e.g., a
10 corn starch, to produce an oligosaccharide pattern that is tightly grouped at about 20,000 MW (*Bacillus* amylases will produce syrups containing much higher MW fragments, and high MW oligosaccharides are not fully converted to glucose by glucoamylases, e.g., *Aspergillus* glucoamylases, during saccharification).

Using the amylases of the invention of archael origin to catalyze the
15 hydrolysis of a starch-comprising composition, e.g., a corn starch, the approximately 20,000 MW fragments are produced. These approximately 20,000 MW fragments can be rapidly and fully converted to glucose. Thus, in one aspect, saccharified syrups resulting from *Bacillus* amylase liquefaction contain less dextrose than saccharified syrups from liquefaction using amylases of the invention.

20 *Processes to produce homogenous maltodextrins*

The invention provides processes to produce homogenous maltodextrins using enzymes of the invention. The homogenous maltodextrins produced by the methods of the invention can be used in a wide variety of food, drug and coating applications. In one aspect, amylases of the invention of archael origin, including the
25 archael amylases of SEQ ID NO:79, SEQ ID NO:80, SEQ ID NO:81, SEQ ID NO:82, SEQ ID NO:115, SEQ ID NO:116, SEQ ID NO:322, SEQ ID NO:323, and enzymes of the invention having the same activity as these archael amylases, can generate an extremely uniform maltodextrin composition (conventional manufacturing processes using either acid or enzymatic hydrolysis of starch result in a broad, typically bimodal
30 MW distribution of oligosaccharides). The homogenous maltodextrins produced by the methods of the invention have a homogenous MW distribution and can be used in a

variety of maltodextrin-comprising products, resulting in lower viscosity, clear (no haze) solutions, better coating properties, better film-forming properties, and the like.

In one aspect, amylases of the invention of archael origin (and enzymes of the invention having the same activity as these archael amylases) are used to liquefy corn starch to produce a uniform maltodextrin-comprising composition. In one aspect, the liquefaction is conducted at a pH of between about pH 4.5 to about pH 6.5, e.g., pH 5.0 or 5.5, at temperatures up to about 105°C. The uniform maltodextrin composition can be produced at DE's ranging from about 5 to as high as about 20. The syrups produced by these archael-derived amylases of the invention can be filtered, treated with charcoal and/or spray-dried to yield the maltodextrin-comprising product.

Enzymatic dry milling processes

The invention provides enzymatic dry milling processes using an amylase of the invention. In dry milling, whole grain is ground and combined with water. The germ is optionally removed by flotation separation or equivalent techniques. The resulting mixture, which contains starch, fiber, protein and other components of the grain, is liquefied using amylase. In one aspect, enzymatic liquefaction is done at lower temperatures than the starch liquification processes discussed above. In one aspect, after gelatinization the starch solution is held at an elevated temperature in the presence of amylase until a DE of 10-20 is achieved. In one aspect, this is a period of about 1-3 hours. Dextrose equivalent (DE) is the industry standard for measuring the concentration of total reducing sugars, calculated as D-glucose on a dry weight basis. Unhydrolyzed granular starch has a DE of virtually zero, whereas the DE of D-glucose is defined as 100.

Enzymatic wet milling processes

The invention provides wet milling processes, e.g., corn wet milling, using an enzyme, e.g., an amylase, of the invention. Corn wet milling is a process which produces corn oil, gluten meal, gluten feed and starch. Thus, the invention provides methods of making corn oil, gluten meal, gluten feed and starch using an enzyme of the invention. In one aspect, an alkaline-amylase of the invention is used in the liquefaction of starch. In one aspect, a glucoamylase of the invention is used in saccharification to produce glucose. An exemplary corn wet milling process of the invention (using at least one enzyme of the invention) is illustrated in Figure 25. Figure 25 illustrates an exemplary corn oil process of the invention comprising steeping, de-germing, de-fibering

and gluten separation, followed by liquefaction using an enzyme of the invention (e.g., an alpha amylase), and saccharification using an enzyme of the invention (e.g., glucoamylase).

In one aspect, corn (a kernel that consists of a outer seed coat (fiber), starch, a combination of starch and glucose and the inner germ), is subjected to a four step process, which results in the production of starch. In one aspect, the corn is steeped, de-germed, de-fibered, and the gluten is separated. In a steeping process the solubles are taken out. The product remaining after removal of the solubles is de-germed, resulting in production of corn oil and production of an oil cake, which is added to the solubles from the steeping step. The remaining product is de-fibered and the fiber solids are added to the oil cake/solubles mixture. This mixture of fiber solids, oil cake and solubles forms a gluten feed. After de-fibering, the remaining product is subjected to gluten separation. This separation results in a gluten meal and starch. The starch is then subjected to liquefaction and saccharification using polypeptides of the invention to produce glucose.

Figure 25 illustrates an exemplary corn wet milling process of the invention (using at least one enzyme of the invention). Figure 26, Figure 27 and Figure 28 illustrate alternative exemplary starch processes, including starch liquefaction processes, of the invention (using at least one enzyme of the invention).

Anti-staling processes

The invention provides anti-staling processes (e.g., of baked products such as bread) using an amylase of the invention. The invention provides methods to slow the increase of the firmness of the crumb (of the baked product) and a decrease of the elasticity of the crumb using an amylase of the invention. Staling of baked products (such as bread) is more serious as time passes between the moment of preparation of the bread product and the moment of consumption. The term staling is used to describe changes undesirable to the consumer in the properties of the bread product after leaving the oven, such as an increase of the firmness of the crumb, a decrease of the elasticity of the crumb, and changes in the crust, which becomes tough and leathery. The firmness of the bread crumb increases further during storage up to a level, which is considered as negative. Amylases of the invention are used to retard staling of the bread as described e.g., in U.S. Patent Nos. 6,197,352; 2,615,810 and 3,026,205; Silberstein (1964) Baker's Digest 38:66-72.

In one aspect, an enzyme of the invention is used to retard the staling of baked products while not hydrolyzing starch into the branched dextrins. Branched dextrins are formed by cleaving off the branched chains of the dextrins generated by α -amylase hydrolysis which cannot be degraded further by the α -amylase. This can
5 produce a gummy crumb in the resulting bread. Accordingly, the invention provides a process for retarding the staling of baked products (e.g., leavened baked products) comprising adding an enzyme of the invention comprising exoamylase activity to a flour or a dough used for producing a baked product. Exoamylases of the invention can have glucoamylase, β -amylase (which releases maltose in the beta-configuration) and/or
10 maltogenic amylase activity.

The invention also provides a process for preparing a dough or a baked product prepared from the dough which comprises adding an amylase of the invention to the dough in an amount which is effective to retard the staling of the bread. The invention also provides a dough comprising said amylase and a premix comprising flour
15 together with said amylase. Finally, the invention provides an enzymatic baking additive, which contains said amylase.

The invention also provides a high yield process for producing high quality corn fiber gum by treatment of corn fiber with an enzyme of the invention followed by hydrogen peroxide treatment to obtain an extract of milled corn fiber. See,
20 e.g., U.S. Patent No. 6,147,206.

Animal feeds and additives

The invention provides methods for treating animal feeds and additives using amylase enzymes of the invention. The invention provides animal feeds and additives comprising amylases of the invention. In one aspect, treating animal feeds and
25 additives using amylase enzymes of the invention can help in the availability of starch in the animal feed or additive. This can result in release of readily digestible and easily absorbed sugars.

Use of an amylase of the invention can increase the digestive capacity of animals and birds. Use of an amylase of the invention can ensure availability of an
30 adequate nutrient supply for better growth and performance. In one aspect, the enzymes of the invention can be added as feed additives for animals. In another aspect, the animal feed can be treated with amylases prior to animal consumption. In another aspect, the

amylases may be supplied by expressing the enzymes directly in transgenic feed crops (as, e.g., transgenic plants, seeds and the like), such as corn. As discussed above, the invention provides transgenic plants, plant parts and plant cells comprising a nucleic acid sequence encoding a polypeptide of the invention. In one aspect, the nucleic acid is
5 expressed such that the amylase is produced in recoverable quantities. The amylase can be recovered from any plant or plant part. Alternatively, the plant or plant part containing the recombinant polypeptide can be used as such for improving the quality of a food or feed, e.g., improving nutritional value, palatability, and rheological properties, or to destroy an antinutritive factor.

10 *Paper or pulp treatment*

The enzymes of the invention can be in paper or pulp treatment or paper deinking. For example, in one aspect, the invention provides a paper treatment process using amylases of the invention. In one aspect, the enzymes of the invention can be used to modify starch in the paper thereby converting it into a liquefied form. In another
15 aspect, paper components of recycled photocopied paper during chemical and enzymatic deinking processes. In one aspect, amylases of the invention can be used in combination with cellulases. The paper can be treated by the following three processes: 1) disintegration in the presence of an enzyme of the invention, 2) disintegration with a deinking chemical and an enzyme of the invention, and/or 3) disintegration after soaking
20 with an enzyme of the invention. The recycled paper treated with amylase can have a higher brightness due to removal of toner particles as compared to the paper treated with just cellulase. While the invention is not limited by any particular mechanism, the effect of an amylase of the invention may be due to its behavior as surface-active agents in pulp suspension.

25 The invention provides methods of treating paper and paper pulp using one or more polypeptides of the invention. The polypeptides of the invention can be used in any paper- or pulp-treating method, which are well known in the art, see, e.g., U.S. Patent No. 6,241,849; 6,066,233; 5,582,681. For example, in one aspect, the invention provides a method for deinking and decolorizing a printed paper containing a dye, comprising
30 pulping a printed paper to obtain a pulp slurry, and dislodging an ink from the pulp slurry in the presence of an enzyme of the invention (other enzymes can also be added). In another aspect, the invention provides a method for enhancing the freeness of pulp, e.g.,

pulp made from secondary fiber, by adding an enzymatic mixture comprising an enzyme of the invention (can also include other enzymes, e.g., pectinase enzymes) to the pulp and treating under conditions to cause a reaction to produce an enzymatically treated pulp.

The freeness of the enzymatically treated pulp is increased from the initial freeness of the
5 secondary fiber pulp without a loss in brightness.

Repulping: treatment of lignocellulosic materials

The invention also provides a method for the treatment of lignocellulosic fibers, wherein the fibers are treated with a polypeptide of the invention, in an amount which is efficient for improving the fiber properties. The amylases of the invention may
10 also be used in the production of lignocellulosic materials such as pulp, paper and cardboard, from starch reinforced waste paper and cardboard, especially where repulping occurs at pH above 7 and where amylases can facilitate the disintegration of the waste material through degradation of the reinforcing starch. The amylases of the invention can be useful in a process for producing a papermaking pulp from starch-coated printed paper.
15 The process may be performed as described in, e.g., WO 95/14807.

An exemplary process comprises disintegrating the paper to produce a pulp, treating with a starch-degrading enzyme before, during or after the disintegrating, and separating ink particles from the pulp after disintegrating and enzyme treatment. See
20 also U.S. Patent No. 6,309,871 and other US patents cited herein. Thus, the invention includes a method for enzymatic deinking of recycled paper pulp, wherein the polypeptide is applied in an amount which is efficient for effective de-inking of the fiber surface.

Waste treatment

25 The enzymes of the invention can be used in a variety of other industrial applications, e.g., in waste treatment. For example, in one aspect, the invention provides a solid waste digestion process using enzymes of the invention. The methods can comprise reducing the mass and volume of substantially untreated solid waste. Solid waste can be treated with an enzymatic digestive process in the presence of an enzymatic
30 solution (including an enzyme of the invention) at a controlled temperature. This results in a reaction without appreciable bacterial fermentation from added microorganisms. The solid waste is converted into a liquefied waste and any residual solid waste. The resulting

liquefied waste can be separated from said any residual solidified waste. See e.g., U.S. Patent No. 5,709,796.

Oral care products

The invention provides oral care product comprising an amylase of the invention. Exemplary oral care products include toothpastes, dental creams, gels or tooth powders, odontics, mouth washes, pre- or post brushing rinse formulations, chewing gums, lozenges, or candy. See, e.g., U.S. Patent No. 6,264,925.

Brewing and fermenting

The invention provides methods of brewing (e.g., fermenting) beer comprising an amylase of the invention. In one exemplary process, starch-containing raw materials are disintegrated and processed to form a malt. An amylase of the invention is used at any point in the fermentation process. For example, amylases of the invention can be used in the processing of barley malt. The major raw material of beer brewing is barley malt. This can be a three stage process. First, the barley grain can be steeped to increase water content, e.g., to around about 40%. Second, the grain can be germinated by incubation at 15-25°C for 3 to 6 days when enzyme synthesis is stimulated under the control of gibberellins. During this time amylase levels rise significantly. In one aspect, amylases of the invention are added at this (or any other) stage of the process. The action of the amylase results in an increase in fermentable reducing sugars. This can be expressed as the diastatic power, DP, which can rise from around 80 to 190 in 5 days at 12°C.

Amylases of the invention can be used in any beer producing process, as described, e.g., in U.S. Patent No. 5,762,991; 5,536,650; 5,405,624; 5,021,246; 4,788,066.

25 *Use in drilling well and mining operations*

The invention also includes methods using enzymes of the invention in well and drilling operations, e.g., gas, oil or other drilling or mining operations. For example, in one aspect, enzymes of the invention are used to increase the flow of production fluids from a subterranean formation, e.g., a well or a mine. In one aspect, the enzymes of the invention are used to remove viscous, starch-containing fluids that can be damaging, e.g., fluids formed during production operations. These starch-containing fluids can be found within a subterranean formation which surrounds a completed well

bore. In one aspect, an amylase of the invention is used in an oil well drilling fluid to aid in the carrying away of drilling mud.

In one aspect, the method comprises allowing production fluids (comprising enzymes of the invention) to flow from the well bore or a mine. The methods can comprise reducing the flow of production fluids from the formation below expected flow rates and formulating an enzyme treatment by blending together an aqueous fluid and a polypeptide of the invention. The methods can comprise pumping the enzyme treatment to a desired location within the well bore or other drilled shaft and allowing the enzyme treatment to degrade the viscous, starch-containing, damaging fluid. The methods can comprise removing the fluid from the subterranean formation to the well or shaft surface. In one aspect, the enzyme treatment is effective to attack the alpha glucosidic linkages in the starch-containing fluid. In one aspect, amylases of the invention are used in mine drilling, well drilling (e.g., gas or oil well drilling), and the like to carry away drilling mud, e.g., while drilling the hole (well bore or shaft).

The enzymes of the invention can be used in any well, shaft or mine drilling operation, many of which are well known in the art. For example, the invention provides methods of introducing an enzyme of the invention, which in one aspect can also comprise an oil or gas field production chemical, into a rock formation comprising oil and/or gas, which comprises passing a microemulsion comprising the enzyme (and, in one aspect, the chemical) down a production well and then into the formation. In one aspect, a production well is subjected to a "shut-in" treatment whereby an aqueous composition comprising an enzyme of the invention is injected into the production well under pressure and "squeezed" into the formation and held there. See, e.g., U.S. Patent No. 6,581,687.

In one aspect, the amylases of the invention used in gas, oil or other drilling or mining operations are active at high or low pH and/or high or low temperatures, e.g., amylases of the invention used in these processes are active under conditions comprising about pH 6.5, pH 6, pH 5.5, pH 5, pH 4.5 or pH 4, or lower, or, under conditions comprising about pH 7, pH 7.5 pH 8.0, pH 8.5, pH 9, pH 9.5, pH 10, pH 10.5 or pH 11 or higher. In one aspect, the amylases of the invention used in these processes are active under conditions comprising a temperature range of anywhere between about 0°C to about 37°C, or, between about 37°C to about 95°C or more, or,

between about 80°C to about 120°C, e.g., 85°C, 90°C, 95°C, 98°C, 100°C, 105°C, 110°C, 115°C, 120°C or more.

Delayed release compositions

The invention provides delayed release or "controlled release"

- 5 compositions comprising a desired composition coated by a latex polymer, e.g., a latex paint, or equivalent. The delayed release/ controlled release compositions of the invention can comprise any desired composition, including enzymes or any active ingredient, including small molecules, drugs, polysaccharides, lipids, nucleic acids, vitamins, antibiotics, insecticides, and the like. In one aspect, the coating will not readily
10 dissolve at a relatively low temperature but will decompose to release the desired composition (e.g., enzyme) at a relatively higher temperature.

The invention provides methods for the delayed release/ controlled release of compositions wherein the composition is coated by a latex polymer, e.g., a latex paint, or equivalent.

- 15 The delayed release/ controlled release compositions and methods of the invention can be used for a variety of medical and industrial applications, for example, in one aspect, delayed release/ controlled release enzyme compositions of the invention comprise enzymes involved in guar fracturing fluids in enhanced oil recovery operations. The oilfield guar degrading application of the invention is facilitated by a coating that will
20 not readily dissolve at low temperature but will decompose to release the enzyme at higher temperatures.

- In another aspect, the delayed release/ controlled release enzyme compositions of the invention comprise animal feeds or nutritional supplements comprising, e.g., enzymes, vitamins, antibiotics and/or other food, drug or nutritional
25 supplements. These active compounds in the animal feeds or nutritional supplements are protected from pelleting conditions or gastric digestion by the coating on a delayed release/ controlled release composition of the invention.

- In one aspect, the release is a temperature activated release, e.g., the desired composition (e.g., enzyme) is released at an elevated temperature, e.g., between
30 about 37°C to about 95°C or more, e.g., 85°C, 90°C, 95°C, 98°C, 100°C or more. The rate of release can be controlled by the thickness or amount of "barrier" or latex polymer, applied to the desired composition, e.g., a pellet or matrix comprising the desired

composition. Thus, the invention provides pellets or matrices having a range of thicknesses of latex polymer or equivalent and methods of using them.

The invention provides delayed release/ controlled release enzyme compositions, e.g., in one aspect, comprising an enzyme of the invention. In one aspect, the invention provides an enzyme (e.g., an enzyme of the invention), or a pelleted composition comprising an enzyme (e.g., an enzyme of the invention), coated with a latex polymer, e.g., a latex paint, or equivalent. In one aspect, the invention provides methods of making delayed release enzyme compositions comprising coating an enzyme (e.g., an enzyme of the invention), or a pelleted composition comprising an enzyme (e.g., an enzyme of the invention), with a latex polymer, e.g., a latex paint, or equivalent. In one aspect, the invention provides methods of making delayed release/ controlled release compositions comprising coating a desired compound with a latex polymer, e.g., a latex paint, or equivalent.

Latex polymers that are used in the delayed release/ controlled release compositions (e.g., delayed release/ controlled release enzyme compositions) and methods of the invention include, but are not limited to, various types such as the following: acrylics; alkyds; celluloses; coumarone-indenes; epoxys; esters; hydrocarbons; maleics; melamines; natural resins; oleo resins; phenolics; polyamides; polyesters; rosins; silicones; styrenes; terpenes; ureas; urethanes; vinyls; and the like. Latex polymers that are used in the delayed release compositions and methods of the invention also include, but are not limited to, one or more homo- or copolymers containing one or more of the following monomers: (meth)acrylates; vinyl acetate; styrene; ethylene; vinyl chloride; butadiene; vinylidene chloride; vinyl versatate; vinyl propionate; t-butyl acrylate; acrylonitrile; neoprene; maleates; fumarates; and the like, including plasticized or other derivatives thereof.

The amount of latex polymer used in the latex composition of the invention is not critical, but may be any amount following well established procedures using latex polymers. In alternative aspects, the amount of dry latex polymer is at least about 1, or, from about 2 to about 50, or, from about 3 to about 40 weight percent of the total latex composition. The latex composition of the invention may optionally contain other components such as those generally used in latex compositions. These additional components include, but are not limited to, one or more of the following: solvents such as aliphatic or aromatic hydrocarbons, alcohols, esters, ketones, glycols, glycol ethers,

nitroparaffins or the like; pigments; fillers, dryers; flatting agents; plasticizers; stabilizers; dispersants; surfactants; viscosifiers including polymeric associative thickeners, polysaccharide-based thickeners and so on; suspension agents; flow control agents; defoamers; anti-skinning agents; preservatives; extenders; filming aids; crosslinkers; surface improvers; corrosion inhibitors; and other ingredients useful in latex compositions. In one aspect, latex compositions of the invention having improved rheology and stability are provided by combining the latex polymer and a polysaccharide with water following established procedures. See, e.g., U.S. Patent Nos. 6,372,901; 5,610,225.

In one aspect, in making a pelleted or matrix-comprising composition of the invention comprising an active composition, e.g., an enzyme (e.g., an enzyme of the invention), coated with a latex polymer, e.g., a latex paint, or equivalent, the active composition (e.g., enzyme) is embedded in the body of the pellet (in one aspect, a majority, or all, of the active composition (e.g., enzyme) is embedded in the pellet. Thus, harsh chemicals, e.g., the latex coating, which may be an inactivator of the desired, active ingredient, can be used to coat the surface of the pellet or matrix. The composition of the coating can be broken down by agents such as heat, acid, base, pressure, enzymes, other chemicals and the like, to have a controlled release of the desired enzymatic activity triggered by the exposure to the coating-degrading agent.

In one aspect, an active composition, e.g., an enzyme (e.g., an enzyme of the invention, or another enzyme, e.g., a mannanase), is dispersed in a corn term meal and/or a corn starch matrix (e.g., as a pellet). This mixture (e.g., pellet) disintegrates within ten minutes in room temperature (e.g., about 22°C) water to release all (100%) of the active composition, e.g., releases all of the enzymatic activity. At higher temperatures, the rate of release increases. This is not an acceptable rate of disintegration for many uses.

However, as a delayed release/ controlled release composition of the invention, i.e., when this mixture is coated with a latex polymer, e.g., a latex paint, or equivalent, the disintegration of the mixture (e.g., pellet, matrix) is delayed. The rate and extent of release can be controlled by the thickness of the coating (barrier) applied to the pellet or matrix. For example, a coated particle will release only 30% of the activity after six hours in 22°C water. At 60°C, 50% of the enzyme is released in 90 minutes. At 80°C, 80% of the enzyme is released during one hour.

The invention will be further described with reference to the following examples; however, it is to be understood that the invention is not limited to such examples.

5

EXAMPLES

EXAMPLE 1: Identification and Characterization of Thermostable α -Amylases

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention. This example describes the identification of novel acid amylases of the invention. The screening program was
10 carried out under neutral and low pH conditions. DNA libraries generated from low pH samples were targeted for discovery. This effort afforded the discovery of hundreds of clones having the ability to degrade starch. DNA sequence and bioinformatic analyses classified many of these genes as previously unidentified amylases.

Biochemical studies

15

Biochemical analysis of the amylase genomic clones showed that many had pH optima of less than pH 6. Lysates of these genomic clones were tested for thermal tolerance by incubation at 70°C, 80°C, 90°C or 100°C for 10 minutes and measurement of residual activity at pH 4.5. Those clones retaining >50% activity after heat treatment at 80°C were chosen for further analysis. These clones were incubated at
20 90°C for 10 minutes at pH 6.0 and 4.5 and tested for residual activity at pH 4.5 (Figure 5). A number of clones retained >40% of their activity following this treatment. For comparative purposes, residual activity of an enzyme of the invention (an "evolved" amylase), SEQ ID NO:437 (encoded by SEQ ID NO:436), was equivalent to the best of the second-generation enzymes; the specific activity of SEQ ID NO:437 was greater.

25

Thermal activity of the clones with residual activity after heat treatment at 90°C at pH 4.5 was measured at room temperature, 70°C and 90°C at pH 4.5. Table 1 shows that the hydrolysis rates of SEQ ID NO: 87 (*B. stearothermophilus* amylase) and SEQ ID NO. 113 (*B. licheniformis* amylase) decrease at higher temperatures, whereas the rate for SEQ ID NO:125 continues to increase as the temperature is raised to 70°C and
30 only reduces by around 50% at 90°C.

The exemplary polypeptide having a sequence as set forth in SEQ ID NO:437 (encoded by SEQ ID NO:436) is thermostable, retaining 50% activity after 25 minutes at 100°C in the absence of added calcium, at pH 4.5. This exemplary polypeptide retained 90% activity after 60 minutes at 100°C in the presence of 40 mg/L calcium, pH 4.5. The activity profile of the polypeptide SEQ ID NO:437 is in the range of between about 4.8 and 5.0. Added calcium is not required for activity.

The polypeptide SEQ ID NO:437 can have a light brown to yellow liquid with a specific gravity of 1.1, at pH 10, when formulated with 35% glycerol. Its alpha amylase activity is between about 110 to 115 IAU* / gram (*IAU = INNOVASE™ activity unit). One analytical method used comprised hydrolysis of 4-nitrophenyl-alpha-D-hexa-glucopyranoside (this same method can be used to determine if an enzyme is within the scope of the invention).

Candidate evaluation

Based on residual activity at pH 4.5 after a 90°C heat treatment, specific activity and rate of starch hydrolysis at 90°C when compared with *B. licheniformis* amylase, SEQ ID NO:125 is compared with the enzyme (an "evolved" amylase) of SEQ ID NO:437 in a starch liquefaction assay.

Table 1.	Room temperature	70°C	90°C
SEQ ID NO.:87 ¹	1.25	1.43	0.33
SEQ ID NO.: 113 ²	3.3	1.9	0.39
SEQ ID NO.: 125	1.9	47	19

Table 1 shows rates of dye labeled starch hydrolysis (relative fluorescence units/s) of three genomic clones at pH 4.5 and 3 different temperatures. ¹*B. stearrowthermophilus* amylase, ²*B. licheniformis* amylase.

The following table is a summary of Average Relative Activity (ARA), Thermal Tolerance, Thermal Stability, Specific Activity and Expression (Units / L) for selected exemplary enzymes of the invention (for example, SEQ ID NOS: 125, 126, refers to a polypeptide having a sequence as set forth in SEQ ID NO:126, encoded by SEQ ID NO:125, etc.):

Enzyme	Expression Host	Average Relative Activity (ARA)	pH Optimum	Thermal Tolerance %RA after 5 min** 50, 60, 70, 80, 90°C	Thermal Stability %RA 37, 65, 80°C	Specific Activity (U/mg at pH 5.3, 37°C)	Expression (Units / L)
Benchmark		80					
SEQ ID NOS:			4.0 to 5.5	105, 107, 88, 58, 27	100, 83, 0	82	
125, 126	<i>Pichia</i>	66	4.5 to 6.0	86, 88, 100, 86, 65	100, 347, 553	81	8521
378, 379	<i>Pichia</i>	66	6.0 to 7.0	22, 0, 0, 0, 0		937	183615
416, 417	<i>Pichia</i>	59	4.5 to 5.0	56, 1, 1, 0, 1		39	23256
203, 204	<i>Pichia</i>	61	6.0 to 7.0	18, 2, 3, 2, 3		20	122107
434, 435	<i>Pichia</i>	76	6.0 to 6.5	151, 58, 0, 0, 0		151	17171
420, 421	<i>Pichia</i>	84	5.5 to 7.0	68, 26, 0, 0, 0		75	5005
350, 351	<i>Pichia</i>	59	6.0 to 7.0	6, 0, 0, 0, 0		104	39662
402, 403	<i>Pichia</i>	67	5.5 to 6.0	42, 8, 11, 12, 16		535	75053
336, 337	<i>Pichia</i>	63	4.5 to 5.5	124, 105, 115, 108, 117	100, 0, 0	572	20822
430, 431	<i>Pichia</i>	50	6.0 to 6.5	111, 86, 82, 89, 35		138	6556
127, 128	<i>Pichia</i>	71	5.5 to 6.5	127, 115, 53, 4, 5		17	114999

101, 102	<i>Pichia</i>	63	5.0 to 5.5	124, 164, 145, 120, 144		28	11559
388, 389	<i>Pichia</i>	80	6.0 to 7.0	87, 29, 5, 0, 0		259	163163
539, 540	<i>Pichia</i>	TBD	4.0 to 4.5	102, 100, 31, 12, 3	100, 186, 123	TBD	TBD

A.R.A. is Average Relative Activity. A.R.A. is calculated as the average relative activity of an amylase between pH 4 and pH 7.5.

Approximate units per liter expression is calculated as follows: (total units of amylase present in recovered lyophilized powder) (volume of culture in fermenter)

5 Evaluation of the amylase SEQ ID NO:437

The amylase SEQ ID NO:437 (encoded by SEQ ID NO:436) was evaluated under a variety of conditions. In the following protocols N^o2 yellow dent corn was used as a starch source.

Liquefaction

10 A starch slurry comprising 35% dry solids ("DS") was subjected to primary liquefaction for five minutes under various temperatures in the range of 95°C to 119°C (e.g., at about 110°C), with an enzyme concentration of between 0.2 to 0.8 gram/kilogram (g/kg) starch DS, with added calcium in the range of between zero and 30 parts per million (ppm), at pH 4.0 to pH 5.6. Secondary liquefaction comprised
15 conditions of 120 minutes at 95°C.

Saccharification

Saccharification was initially tested using 35% dry solids ("DS") (starch slurry) and glucoamylase AMG 300L (Novozymes A/S, Denmark) at 0.225 AGU/gram DS (AGU= amyloglucosidase, or glucoamylase, units), pH 4.3, at 60°C for 44 hours.

20 The amylase SEQ ID NO:437 was demonstrated to be useful under the above-described pH conditions, was calcium independent and had a high thermal stability. In one aspect, amylase SEQ ID NO:437, or another amylase of the invention, is used in a dosage range of between 0.5 to 0.7 kg / MT DS starch.

The invention provides methods for making nutritive sweeteners using
25 enzymes of the invention, e.g., processes comprising the above described liquefaction and

saccharification protocols using, e.g., amylase SEQ ID NO:437, or another enzyme of the invention. In one aspect, the dosage range for an enzyme of the invention in these processes is between about 0.5 to 0.7 gram per kg starch DS, a jet temperature (e.g., using a jet cooker) of about 110°C, pH 4.5, no added calcium.

5 Dry Mill Ethanol Production

The invention provides methods for Dry Mill Ethanol Production using enzymes of the invention, e.g., amylase SEQ ID NO:437, or another enzyme of the invention.

In evaluating enzymes of the invention for use in Dry Mill Ethanol
10 Production, particularly, liquefaction of dry mill corn flour, a bench scale reactor was used with corn flour sourced from commercial dry mill. TERMAMYL™ SC (Novozymes A/S, Denmark) amylase was used as a competitive benchmark. Test found optimum conditions to be 85°C, pH 5.7. Five independent variables were studied: temperature (in a range of between 80°C to 100°C), enzyme dose of between 0.2 to 1.0
15 g/kg starch, pH 4.4 to 6.0, calcium in a range between 0 ppm to 200 ppm, and a recycled backset between about 0% to 40%.

At 95°C amylase SEQ ID NO:437 reduces viscosity of dry mill corn flour more rapidly than TERMAMYL™ SC (Novozymes A/S, Denmark) amylase at its optimum conditions, including at 85°C. The rate of viscosity reduction by amylase SEQ
20 ID NO:437 was influenced most by enzyme dose and temperature. The optimal range was found to be in the range of 0.4 to 0.6 g/kg starch, with an optimum temperature at 95°C. The amylase SEQ ID NO:437 was effective at a lower pH and a higher temperature than TERMAMYL™ SC (Novozymes A/S, Denmark) amylase at a pH in the range between pH 4.4 and pH 5.6. Calcium addition had a minimal effect on rate of
25 viscosity reduction at 95°C. The amylase SEQ ID NO:437 was effective in the presence of a 30% recycled backset (e.g., thin stillage, spent wash = recycling byproducts back into liquefaction). Figure 29 shows data summarizing these findings comparing amylase SEQ ID NO:437 with TERMAMYL™ SC (Novozymes A/S, Denmark) amylase in dry mill ethanol processing.

30 In alternative aspects, use of amylase SEQ ID NO:437 in dry mill ethanol processes can provide operational advantages, for example: rapid reduction in viscosity of slurried corn flour, making an increase in dissolved solids and throughput possible

- without additional capital investment; superior thermal stability to best competitor, which eliminates split dosing (amylase SEQ ID NO:437 is a thermostable enzyme and eliminates the need to dose before jet cooking and after), lower viscosities are obtained at higher process temperatures, and provides improved microbial control in slurry tank
- 5 (process is run at higher temperature, so unwanted microbes are killed); lower liquefaction pH, which eliminates need for pH adjustment, decreases scale formation (calcium oxalate precipitate forms on hardware, etc.; if liquefaction done at low pH, there is a higher potential for scale formation) and reduces byproduct formation.

In summary, amylase SEQ ID NO:437 is a thermostable enzyme that can

10 meet key industry needs, for example, under certain conditions, rapidly reduces viscosity of high dry solids corn flour slurry, can be thermostable (optimum temperature 95°C), can be calcium independent, can be active under low pH optimum, and can tolerate up to 30% recycled backset. In one aspect, the recommended dose is in the range of between about 0.4 to 0.6 kg/ MT starch.

15

EXAMPLE 2: Thermostable Amylases Active at Alkaline pH

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, e.g., is a thermostable amylase.

- The initial focus of this example was the evaluation of an existing panel of
- 20 amylases in an commercial automatic dish wash (ADW) formulation. This effort identified two candidates: one with activity at high pH (SEQ ID NO.:115) and another with stability in the ADW formulation (SEQ ID NO.:207). Studies also included the identification of high pH amylases. This effort afforded the discovery of hundreds of clones having the ability to degrade starch. DNA sequence and bioinformatics analyses
- 25 classified many of these genes as previously unidentified amylases. The remaining open reading frames were neopullulanases, amylopullulanases and amylomaltases. Extensive biochemical and applications studies showed that 3 candidates: clone B, SEQ ID NO.:147 and SEQ ID NO.:139) have high specific activity at pH10, but unfortunately lack stability in the ADW formulation. In summary, a panel of novel amylases each having desirable
- 30 phenotypes for the ADW application has been identified.

Biochemical studies

Biochemical analysis of the amylase genomic clones showed that many of them hydrolyzed starch at pH 10 and 50°C. To produce sufficient quantities of enzyme for further biochemical and applications testing, the amylase open reading frames of the 40 most active genomic clones were subcloned into expression vectors. This effort included making 2 constructs for those clones containing a putative signal sequence and establishing the growth and induction conditions for each subclone (plus and minus the amylase signal peptide).

Soluble, active protein was successfully purified to homogeneity from 34 subclones and specific activity (units/mg, where 1 unit = μmol reducing sugars/min) was measured at pH 8 and pH 10 (40°C and 50°C) using 2% starch in buffer. The amylase from *Bacillus licheniformis* (SEQ ID NO.:113) was chosen as the benchmark for these studies. Specific activity was determined by removing samples at various time points during a 30 minute reaction and analyzing for reducing sugars. The initial rate was determined by fitting the progress curves to a linear equation. A comparison of the top candidates is shown in Table 2.

A study to determine the dependence of hydrolysis rate on pH showed that only clone B is an "alkaline amylase" with a pH optimum of approximately 8; all others had pH optima of 7 or less. Nevertheless, it is clear that the panel of hits included several lead amylases with appreciable activity at pH 10 and 50°C.

Table 2. Specific activities (U/mg pure enzyme) of amylases

Enzyme	Specific activity pH 8, 40°C	Specific activity pH 10, 50°C
Clone B	682	20
SEQ ID NO.:139	430	33
SEQ ID NO.:127	250	47
SEQ ID NO.:137	230	3
SEQ ID NO.:113 (<i>B. licheniformis</i>)	228	27
SEQ ID NO.:205	163	4
Remainder	<40	

Stability

Stability in the presence of the ADW formulation was measured for each of the 3 top candidates identified via biochemical analysis. The benchmark for these studies was a commercial enzyme in the formulation matrix. Figure 13 illustrates the

residual activity (measured at pH 8 and 50°C) after a 30 minute incubation at 50°C in the presence of various components of the ADW formulation; pH 8, pH 10.8, ADW solution (with bleach) and ADW solution (without bleach). The measured activity after the incubation is expressed as a percentage of the original activity. The data show that clone
5 B was very sensitive to high temperature, whereas the other amylases were less affected. When the enzymes were incubated at high pH and temperature, the commercial enzyme SEQ ID NO.: 139 became less stable; however, SEQ ID NO.: 127 retained full activity. The apparently anomalous behavior of SEQ ID NO.: 127 after pH 10 incubation vs pH 8 was observed in repeated trials.

10 When amylase activity on dye-labeled starch is measured in the ADW matrix at 50°C, the commercial amylase exhibits roughly 5% of its activity at pH 8. In the same assay, clone B, SEQ ID NO.: 139 and SEQ ID NO.: 127 exhibit <2% of their original activity measured at pH 8.

Wash tests

15 Wash tests using starch coated slides were carried out to gauge the performance of each of the purified enzymes as compared to the commercial amylase. The spaghetti starch coated slides were prepared according to protocol. Two pre-weighed starch coated slides were placed back to back in a 50 mL conical tube and 25 mL of ADW solution, +/- enzyme were added per tube. The tubes were incubated for 20
20 minutes at 50°C with gentle rotation on a vertical carousel. Following the incubation period, the slides were immediately rinsed in water and oven dried overnight. All trials were run in duplicate and the commercial enzyme was run as a positive control. The results (Figure 6) of these experiments are expressed as net % starch removed, e.g. % of starch removed in ADW with enzyme, *minus* the % of starch removed in ADW alone.

25

EXAMPLE 3: Gene Optimization

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, e.g., assessing enzyme performance in the presence of ADW performance.

30 The properties of enzymes may be improved by various evolution strategies, including GeneSiteSaturationMutagenesis (GSSMTM) and GeneReassemblyTM. (Diversa Corporation, San Diego, CA). Such techniques will be applied to the amylase

nucleic acids of the invention in order to generate pools of variants that can be screened for improved performance. In one aspect, parental molecules for evolution include any nucleic acid of the invention, e.g., are one or all of the following: SEQ ID NO.: 113, SEQ ID NO.: 139, SEQ ID NO.: 115 and SEQ ID NO.: 127 (a truncated form of SEQ ID NO.: 127).

A high throughput screen has been developed to assess enzyme performance in the presence of ADW performance. Development of a HTS is of paramount importance in any evolution program The HTS is automated and has showed consistent results for the parental amylases (Figure 7). Parental amylases have measurable activity in the ADW formulation, however highly reduced relative to pH 8 activity.

EXAMPLE 4: Characterization of α -Amylases having Activity at Alkaline pH

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, for example, has alpha-amylase activity at alkaline pH.

Amylases of the invention having activity at alkaline pH were characterized further. Kinetics on 2% starch at pH 8 and 10 (40°C and 50°C) have been performed.

20

Table 4:

<u>Clones, specific activities</u>	<u>pH 8, 40°C</u>	<u>pH 10, 50°C</u>
SEQ ID NO.: 113 (<i>B. licheniformis</i>)	228 units/mg	27 units/mg
Clone B	682 units/mg	31 units/mg
SEQ ID NO.: 139	430 units/mg	33 units/mg
SEQ ID NO.: 127	540 units/mg	50 units/mg
control 0GL5 (<i>E. coli</i>)	1.8 units/mg	0 units/mg

25

1 unit of activity is defined as release of 1 μ mol reducing sugars per minute.

EXAMPLE 5: Amylase Activity Assay: BCA Reducing Ends Assay

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, for example, by a BCA reducing ends

assay. Amylase activity of clones of interest was determined using the following methodology.

1. Prepare 2 substrate solutions, as follows:

- a) 2% soluble starch (potato) pH 8 solution by dissolving 2 gm potato starch in 100 ml 100 mM sodium phosphate pH 8).
- b) 2% soluble starch (potato) pH 10 solution by dissolving 2 gm potato starch in 100 ml 100 mM sodium carbonate.

Heat both solutions in a boiling water bath, while mixing, for 30-40 minutes until starch dissolves.

2. Prepare Solution A from 64 mg/ml sodium carbonate monohydrate, 24 mg/ml sodium bicarbonate and 1.95 mg/ml BCA (4,4'-dicarboxy-2,2'- biquinoline disodium salt (Sigma Chemical cat # D-8284). Added above to dH₂O.

3. Prepare solution B by combining 1.24 mg/ml cupric sulfate pentahydrate and 1.26 mg/ml L-serine. Add mixture to dH₂O.

4. Prepare a working reagent of a 1:1 ration of solutions A and B.

5. Prepare a Maltose standard solution of 10 mM Maltose in dH₂O, where the 10 mM maltose is combined in 2% soluble starch at desired pH to a final concentration of 0, 100, 200, 300, 400, 600 μ M. The standard curve will be generated for each set of time-points. Since the curve is determined by adding 10 μ l of the standards to the working reagent it works out to 0, 1, 2, 3, 4, 6 nmole maltose.

6. Aliquot 1 ml of substrate solution into microcentrifuge tubes, equilibrate to desired temperature (5 min) in heat block or heated water bath. Add 50 μ l of enzyme solution to the inside of the tube lid.

7. While solution is equilibrating mix 5 ml of both solution A & B. Aliquot 100 μ l to 96 well PCR plate. Set plate on ice.

8. After 5 minute temperature equilibration, close lid on tubes, invert and vortex 3 times. Immediately aliquot 10 μ l into plate as t=0 (zero time point). Leave enzyme mixture in heat block and aliquot 10 μ l at each desired time point (e.g. 0, 5, 10, 15, 20, 30 minutes).

9. Ensure that 12 wells are left empty (only working reagent aliquotted) for the addition of 10 μ l of standards, for the standard curve.

10. When all time points are collected and standards are added, cover plate and heated to 80° C for 35 min. Cool plate on ice for 10 min. Add 100 μ l H₂O to

all wells. Mix and aliquot 100 ul into flat bottomed 96-well plate and read absorbance at 560 nm.

11. Zero each sample's time points against its own t=0 (subtract the average t=0 A560 value from other average A560 values). Convert the $A560_{(experimental)}$ to umole (Divide $A560_{(experimental)}$ by the slope of the standard curve (A560/umole). Generate a slope of the time points and the umole (in umole/min), multiply by 100 (as the umole value only accounts for the 10 ul used in the assay, not the amount made in the 1 ml rxn). To get the specific activity divide the slope (in umole/min) by the mg of protein. All points should be done at a minimum in duplicate with three being best. An example standard curve is set forth in Figure 11.

Table 5: Sample data:

		Dilu		(A560exp/std slope)			
Clone	tion	Minutes	A560-1	A560-2	Avg A 560	Zeroed A 560	umole
ENZ	50	0	0.1711	0.1736	0.17235	0	0.0000
		5	0.2104	0.2165	0.21345	0.0411	0.0005
		10	0.2492	0.2481	0.24865	0.0763	0.0009
		15	0.2984	0.2882	0.2933	0.12095	0.0014
		20	0.3355	0.3409	0.3382	0.16585	0.0020
		30	0.3942	0.3805	0.38735	0.215	0.0026
		40	0.4501	0.4412	0.44565	0.2733	0.0033

Activity = 0.008646 umole/min

Divide protein concentration (mg/ml) by any dilution to get mg used in assay.

- 15 Divide the above slope by mg used in assay to get specific activity

Specific Activity = 24.93 umole/min/mg

See for example, Dominic W.S. Wong, Sarah B. Batt, and George H. Robertson (2000) J. Agric. Food Chem. 48:4540-4543; Jeffrey D. Fox and John F. Robyt, (1991) Anal. Biochem. 195, 93-96.

20

EXAMPLE 6: Screening for α -Amylase activity

- The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention. Amylase activity of clones can be assessed by a number of methods known in the art. The following is the general methodology that was used in the present invention. The number of plaques screened, per plate, should be approximately 10,000 pfu's. For each DNA library: at least 50,000
- 25

plaques per isolated library and 200,000 plaques per non-isolated library should be screened depending upon the pfu titer for the λ Zap Express amplified lysate.

Titer determination of Lambda Library

- 1) μ L of Lambda Zap Express amplified library stock added to 600 μ L *E. coli* MRF' cells (OD₆₀₀=1.0). To dilute MRF' stock, 10mM MgSO₄ is used.
- 2) Incubate at 37 °C for 15 minutes.
- 3) Transfer suspension to 5-6mL of NZY top agar at 50 °C and gently mix.
- 4) Immediately pour agar solution onto large (150mm) NZY media plate.
- 5) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.
- 6) Incubate the plate at 39 °C for 8-12 hours.
- 7) Number of plaques is approximated. Phage titer determined to give 10,000 pfu/plate. Dilute an aliquot of Library phage with SM buffer if needed.

Substrate screening

- 1) Lambda Zap Express (50,000 pfu) from amplified library added to 600 μ L of *E. coli* MRF' cells (OD₆₀₀=1.0). For non-environment libraries, prepare 4 tubes (50,000 pfu per tube).
- 2) Incubate at 37 °C for 15 minutes.
- 3) While phage/cell suspensions are incubating, 1.0mL of red starch substrate (1.2% w/v) is added to 6.0mL NZY top agar at 50 °C and mixed thoroughly. Keep solution at 50°C until needed.
- 4) Transfer 1/5 (10,000 pfu) of the cell suspension to substrate/top agar solution and gently mixed.
- 5) Solution is immediately poured onto large (150mm) NZY media plate.
- 6) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.
- 7) Repeat procedures 4-6 4 times for the rest of the cell suspension (1/5 of the suspension each time).
- 8) Incubate plates at 39 °C for 8-12 hours.
- 9) Plate observed for clearing zones (halos) around plaques.
- 10) Plaques with halos are cored out of agar and transferred to a sterile micro tube. A large bore 200 μ L pipette tip works well to remove (core) the agar plug containing the desired plaque.

11) Phages are re-suspended in 500 μ L SM buffer. 20 μ L Chloroform is added to inhibit any further cell growth.

12) Pure phage suspension is incubated at room temperature for 4 hours or overnight before next step.

5 Isolation of pure clones

1) 10 μ L of re-suspended phage suspension is added to 500 μ L of *E. coli* MRF' cells (OD600=1.0).

2) Incubate at 37 °C for 15 minutes.

3) While phage/cell suspension is incubating, 1mL of red starch substrate (1.2% w/v) is
10 added to 6.0mL NZY top agar at 50 °C and mixed thoroughly. Keep solution at 50 °C until needed.

4) Cell suspension is transferred to substrate/top agar solution and gently mixed.

5) Solution is immediately poured onto large (150mm) NZY media plate.

6) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.

15 7) Plate incubated at 39 °C for 8-12 hours.

8) Plate observed for a clearing zone (halo) around a single plaque (pure clone). If a single plaque cannot be isolated, adjust titer and re-plate phage suspension.

9) Single plaque with halo is cored out of agar and transferred to a sterile micro tube. A large bore 200 μ L pipette tip works well to remove (core) the agar plug containing the
20 desired plaque. To amplify the titer, core 5 single active plaques into a micro tube.

10) Phages are re-suspended in 500 μ L SM buffer. 20 μ L Chloroform is added to inhibit any further cell growth.

11) Pure phage suspension is incubated at room temperature for 4 hours or overnight before next step. The pure phage suspension is stored at -80 °C by adding DMSO
25 into the phage suspension (7% v/v).

Excision of pure clone

1) 100 μ L of pure phage suspension is added to 200 μ L *E. coli* MRF' cells (OD600=1.0). To this, 1.0 μ L of EXASSIST helper phage (>1 x 10⁶ pfu/mL; Stratagene) is added. Use 2059 Falcon tube for excision.

30 2) Suspension is incubated at 37 °C for 15 minutes.

3) 3.0 mL of 2 x YT media is added to cell suspension.

4) Incubate at 30 °C for at least 6 hours or overnight while shaking.

- 5) Tube transferred to 70 °C for 20 minutes. The phagemid suspension can be stored at 4 °C for 1 to 2 months.
- 6) 100 µL of phagemid suspension transferred to a micro tube containing 200µL of *E. coli* Exp 505 cells (OD600=1.0).
- 5 7) Suspension incubated at 37 °C for 15 minutes.
- 8) 300µL of SOB is added to the suspension.
- 9) Suspension is incubated at 37 °C for 30 to 45 minutes.
- 10) 100µL of suspension is transferred to a small (90mm) LB media plate containing Kanamycin (LB media with Kanamycin 50µg/mL) for Zap Express DNA libraries or
- 10 Ampicillin (LB media with Kanamycin 100µg/mL) for Zap II DNA libraries.
- 11) The rest of suspension is transferred to another small LB media plate.
- 12) Use sterile glass beads to evenly distribute suspension on the plate.
- 13) Plates are incubated at 30 °C for 12 to 24 hours.
- 14) Plate observed for colonies.
- 15 15) Inoculate single colony into LB liquid media containing suitable antibiotic and incubate at 30 °C for 12 to 24 hours.
- 16) Glycerol stock can be prepared by adding 80% glycerol into liquid culture (15% v/v) and stored at -80 °C.

Activity verification

- 20 1) 50µL of liquid culture is transferred to a micro tube. Add 500µL of 8% pH7 Amylopectin Azure into the same tube. Prepare 2 tubes for each clone.
- 2) Activity is tested at 50 °C for 3 hours and overnight. Use pH 7 buffer as control.
- 3) Cool the test specimen at ice-water bath for 5 minutes.
- 4) Add 750µL of Ethanol and mixed thoroughly.
- 25 5) Centrifuge at 13000 rpm (16000 g's) for 5 minutes.
- 6) Measure OD of the supernatant at 595nm.

RFLP analysis

- 1) 1.0mL of liquid culture is transferred to a sterile micro tube.
- 2) Centrifuge at 13200 rpm (16000 g's) for 1 minute.
- 30 3) Discard the supernatant. Add another 1.0 mL of liquid culture into the same sterile micro tube.
- 4) Centrifuge at 13200 rpm (16000 g's) for 1 minute.

- 5) Discard the supernatant.
- 6) Follow QIAprep spin mini kit protocol for plasmid isolation.
- 7) Check DNA concentration using BioPhotometer.
- 8) Use Sac I and Kpn I for first double digestion. Incubate at 37 °C for 1 hour.
- 5 9) Use Pst I and Xho I for second double digestion. Incubate at 37 °C for 1 hour.
- 10) Add Loading dye into the digested sample.
- 11) Run the digested sample on a 1.0% agarose gel for 1-1.5 hours at 120 volts.
- 12) View gel with gel imager. All clones with a different digest pattern will be sent for sequence analysis.

10 EXAMPLE 7: Assay for Amylases

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention.

Preparation Of Host Cultures

1. Start an overnight culture of XL1-Blue MRF' host cells. Use a single colony from a streak plate to inoculate 10 mL LB supplemented with 20 ug/mL tetracycline.
- 15 Grow overnight culture shaking at 37°C for at least 16 hours.
2. Using aseptic technique, inoculate a fresh 100 mL of LB_{Tet} day culture with XL1-Blue MRF' host from the overnight LB_{Tet} culture.
3. Grow in a 37°C shaker until the OD reaches 0.75 – 1.0.
- 20 4. Pellet host cells at 1000 x g for 10 minutes and gently resuspend in 10 mM MgSO₄ at OD5.
5. Dilute a small amount of host cells to OD1 for use in titering and pintooling.
6. Host preparations can be used for up to 1 week when stored on ice or at 4°C.

-To shorten growth time for the day culture, use ½X the usual Tet concentration in LB (½X = 10 ug/mL), or omit the antibiotic altogether.

-Do not use NZY when selecting with Tetracycline. The high Mg⁺⁺ concentration in NZY medium renders Tet inactive.

Titering Lambda Libraries

7. Place three sterile microfuge tubes in a rack.
- 30 8. Aliquot 995 uL prepared host cells in one tube and 45 uL prepared OD1 host cells into each of the two remaining tubes.

9. Add 5 uL of lambda library to the tube containing 995 uL host cells and mix by vortexing. This results in a dilution factor of 200.
10. Prepare 1/2,000 and 1/20,000 dilutions by consecutively adding 5 uL of previous dilution to the remaining two tubes containing 45 uL prepared host cells. Mix by vortexing after each dilution was made.
11. Allow phage to adsorb to host by incubating at 37°C for 15 minutes.
12. Meanwhile, pipet 100 uL of prepared OD1 host cells to each of three Falcon 2059 tubes.
13. Add 5 uL of each dilution to a separate 2059 tube containing host cells.
14. Plate each by adding 3 mL top agar to each tube and quickly pour over 90 mm NZY plates. Ensure a smooth, even distribution before the top agar hardens.
15. Invert plates and incubate at 37°C overnight.
16. Count plaques and calculate titer of the library stock (in plaque forming units (pfu) per uL).

15 Lambda Microtiter Screening For Amylases

Preparation

1. Prepare a sufficient amount of XLI-Blue MRF' host culture, as described above, for the amount of screening planned. A culture of 100 mL is usually sufficient for screening 2-3 libraries.
- 20 2. Autoclave several bottles compatible with the QFill2 dispenser. These are the wide-mouth Corning bottles, 250 mL containing a sealing ring around the lip.
3. Make sure there are sufficient amounts of plates, top agar, BODIPY starch, red starch solution, etc. available for the screen.
4. Schedule the Day 2 robot run with a representative from Automation.

25 Day 1

1. Label the 1536-well plates (black) with library screen and plate number. Tough-Tags™ tube stickers, cut in half width-wise, are ideal for labeling 1536 well plates.
2. Calculate volumes of library, host cells and NZY medium necessary for the screen. This is easily done with an Excel spreadsheet.
- 30 3. Combine the calculated volumes of lambda library and OD5 host cells in a sterile 250 mL wide-mouth Corning bottle (containing a sealing ring).
4. Allow adsorption to occur at 37°C for 15 minutes.

5. Add the calculated volume of NZY medium and mix well. This is referred to as the cell-phage-medium suspension.
6. Perform a concomitant titer by combining 50 uL of the cell-phage-medium suspension with 250 uL of OD1 host cells in a Falcon 2059 tube, then plating with 9 mL of top agar onto a 150 mm NZY plate. Incubate concomitant titer plate at 37°C overnight.
7. Load the dispenser with the remainder of the suspension and array each labeled 1536-well plate at 4 uL per well. If the dispenser leaves air bubbles in some wells, they can be removed by centrifuging the plates at 200 x g for 1 minute.
8. Add 0.5 uL of positive control phage to well position AD46 of at least two of the assay plates. Use a strong amylase-positive lambda clone for this purpose. The lambda versions of SEQ ID NO.: 113 or SEQ ID NO.: 199 are good choices for positive controls.
9. Incubate assay plates at 37°C overnight in a humidified (≥95%) incubator.

Day 2

1. Count the pfu on the concomitant titer plate and determine the average seed density per well (in pfu per well).
2. Pintool at least 2 plates of each library screen (preferably the 2 containing positive controls) as follows:
 - a) Prepare 2 host lawn plates to act as a surface on which to pintool: combine 250 uL of OD1 host cells with 2 mL 2% red starch and plate with 9 mL top agar onto 150 mm NZY plates. Hold each plate as level as possible as the top agar solidifies in order to produce an even hue of red across the plate.
 - b) Using a twice flame-sterilized 1536 position pintool, replicate at least 2 of the screening plates onto the host lawn plates.
 - c) Place the pintoled recipient plates in a laminar flow hood with the lids off for about 15-30 minutes (to vent off excess moisture).
 - d) Replace the lids and incubate inverted at 37°C overnight.
3. Prepare the 2X BODIPY starch substrate buffer as follows:
 - a) Calculate the total volume of 2X substrate buffer solution needed for all screening plates at 4 uL per well (including any extra deadspace volume

- required by the dispenser) and measure this amount of 100 mM CAPS pH 10.4 into a vessel appropriate for the dispenser used.
- b) Retrieve enough 0.5 mg tubes of BODIPY starch to produce the required volume of 2X substrate buffer [calculated in step a) above] at a final concentration of 20-30 ug/mL.
 - c) Dissolve each 0.5 mg tube in 50 uL DMSO at room temperature, protected from light, with frequent vortexing. This takes more than 15 minutes; some production lots of BODIPY starch dissolve better than others.
 - d) Add 50 uL 100mM CAPS buffer pH 10.4 to each tube and mix by vortexing.
 - e) Pool the contents of all tubes and remove any undissolved aggregates by centrifuging for 1 minute at maximum speed in a microfuge.
 - f) Add the supernatant to the rest of the 100 mM CAPS buffer measured in step a) above.
 - g) Protect the 2X substrate buffer from light by wrapping in foil.
4. Take plates and substrate buffer to the automation room and program the robot with the following parameters:
 - a) dispense 4 uL substrate buffer per well
 - b) 1st read at 1 hour post-substrate, 2nd read at 9 hours, and third read at 17 hours; with 37°C incubation between reads
 - c) excitation filter: 485 nm; emission filter: 535 nm
 - d) set the Spectrafluor gain at 70, or the optimal gain for the batch of 2X substrate buffer prepared.
 - e) ensure that the incubator used will protect assay plates from light.

Day 3

1. Check pintoled plates for clearings in the bacterial lawn at all positions corresponding to wells on the associated assay plate. Also check for clearings in the red starch in any of the pin positions. If plates containing positive controls were used for pintoled, you should be able to see a large clearing zone in the red background. Be wary of contaminants that also form clearing zones in red starch (see comment "Contaminants That Form Clearing Zones in Red Starch" at end of Example 7).
2. Identify putative hits from the data file produced by the robot computer. The KANAL program produced by Engineering simplifies data analysis. As a rule

of thumb, a putative hit is characterized as a well having signal intensity rising at least 1.5 fold over background.

3. For each putative, remove 2 uL from the well and add to a tube containing 500 uL SM buffer and 50 uL CHCl₃. Vortex to mix and store at 4°C. This solution will be referred to hereafter as the 4e-3 stock. The original screening plates should be stored at 4°C, protected from light, at least until breakouts are complete.

This is the recommended method of breaking out putative hits. It is a liquid phase assay that relies on confirmation of activity on BODIPY starch.

- 10 Alternatively, putative hits can be plated directly onto solid phase plates containing red starch such that 2,000-3,000 pfu per hit are examined for clearing zones. However, inability to observe clearing zones on red starch is not necessarily an indication that a putative hit was a false positive. It would then need to be assayed using the format in which it was originally identified (i.e., liquid phase using BODIPY starch as substrate). In addition, very weak positives are more easily identified using the method detailed below.

Day 1

1. In a sterile 50 mL conical tube, combine 0.5 mL OD5 host cells with 45.5 mL NZY. This will be referred to as the host-medium suspension.
2. For each putative hit to be analyzed, aliquot 1 mL of host-medium suspension into each of 3 three sterile microfuge tubes.
3. Set the 12-channel pipetman in multidispense mode with an aliquot size of 20 uL and an aliquot number of 2x. Mount the pipetman with a clean set of sterile tips.
4. Pour about 1 mL of host-medium suspension into a new sterile solution basin and load the multichannel pipetman.
- 25 5. Dispense 20 uL per well into the last row (row P) of a black 384-well plate (12 channels x 2 = 24 wells). This row will be used later for the controls.
6. Expel the remaining liquid in the tips by touching the tips against the surface of the basin and pressing the RESET button on the pipetman. Lay the pipetman down in a way to prevent contamination of the tips. There is no need to change the tips at this point.
- 30 7. Pour the remainder of the fluid in the basin into a waste container (like a beaker) taking care to avoid splash-back contamination.

8. For the first putative to be analyzed, take 111 uL of the $4e-3$ stock (see Day 2 in *Lambda Microtiter Screening for Amylases*) and add it to the first in a set of three tubes containing 1 mL host-medium suspension (step 2). Vortex to mix. This is *Dilution A*.
- 5 9. Take 111 uL of Dilution A and add to the next tube in the set. Vortex to mix. This is *Dilution B*.
- 10 10. Take 111 uL of Dilution B and add to the last tube in the set. Vortex to mix. This is *Dilution C*. You should now have three dilutions of phage, where concentrations of each differ by a factor of 10.
- 10 11. Pour the contents of Dilution C (the most dilute of the 3 samples) into the solution basin and load the multichannel pipetman.
12. Dispense 20 uL per well into the first row of the 384-well plate (12 channels x 2 = 24 wells).
13. Expel the remaining liquid in the tips by touching the tips against the surface of
15 the basin and pressing the RESET button on the pipetman. Lay the pipetman down in a way to prevent contamination of the tips. There is no need to change the tips at this point.
14. Empty the basin as described above.
15. Pour the contents of Dilution B into the same basin and load the multichannel
20 pipetman.
16. Dispense 20 uL per well into the second row of the 384-well plate.
17. Perform steps 13-16 similarly to dispense Dilution A into the third row of the plate.
18. After all three dilutions have been arrayed into the first 3 rows of the plate, discard
25 all tips and the solution basin into the biohazardous waste container.
19. Mount the pipetman with a clean set of sterile tips and open a new sterile solution basin.
20. Repeat steps 8-19 for each remaining putative hit, using remaining rows on the plate up to row O. Five putative hits can be analyzed on one 384-well plate, with
30 the last row (row P) saved for the controls.
21. Add 0.5 uL of each control to a separate well. Use at least 2-3 separate controls, preferably covering a range of activity.
22. Incubate assay plates at 37°C overnight in a humidified ($\geq 95\%$) incubator.

Day 2

1. Pintool all breakout plates onto a host lawn with red starch using the same method described for Day 2 in *Lambda Microtiter Screening for Amylases*, except that a 384 position pintool is used.
- 5 2. Prepare the 2X BODIPY starch substrate buffer as follows:
 - a) Calculate the total volume of 2X substrate buffer solution needed for all breakout plates at 20 uL per well (including any extra deadspace volume required by the dispenser) and measure this amount of 100 mM CAPS pH 10.4 into a vessel appropriate for the dispenser used.
 - 10 b) Retrieve enough 0.5 mg tubes of BODIPY starch to produce the required volume of 2X substrate buffer [calculated in step a) above] at a final concentration of 20-30 ug/mL.
 - c) Dissolve each 0.5 mg tube in 50 uL DMSO at room temperature, protected from light, with frequent vortexing. This takes more than 15 minutes; some production lots of BODIPY starch dissolve better than others.
 - 15 d) Add 50 uL 100mM CAPS buffer pH 10.4 to each tube and mix by vortexing.
 - e) Pool the contents of all tubes and remove any undissolved aggregates by centrifuging for 1 minute at maximum speed in a microfuge.
 - f) Add the supernatant to the rest of the 100 mM CAPS buffer measured in step
 - 20 a) above.
 - g) Protect the 2X substrate buffer from light by wrapping in foil.
3. Dispense 20 uL per well into all breakout plates.
4. Wrap all plates in aluminum foil and incubate at room temperature for 2-6 hours.
5. Read each plate in the Spectrafluor with the following settings:
 - 25 a) fluorescence read (excitation filter: 485 nm; emission filter: 535 nm)
 - b) plate definition: 384 well black
 - c) read from the top
 - d) optimal gain
 - e) number of flashes: 3
- 30 6. On the resulting Excel spreadsheet, chart each putative's 3 rows in a separate graph and check for activity. Ensure that the positives controls produced signals over background.

7. For each putative that appears to have a real signal among the wells, harvest a sample from a positive well as follows:
 - a) Select a positive well from a row representing the highest initial dilution.
 - b) Transfer 2 uL from that well into a tube containing 500 uL SM and 50 uL CHCl₃. This is referred to as the breakout stock.
 - c) Store at 4°C.
 8. Using methods previously described, plate about 10 uL of each breakout stock onto 150 mm NZY plates using red starch. The objective is to obtain several (at least 20) well-separated plaques from which to core isolates.
- Day 3
1. Check pintoled plates for an acceptable incidence of clearings in the bacterial lawn corresponding to wells on the associated assay plate. Also check for clearings in the red starch in the positive controls and in any tested putatives. Be wary of contaminants that also form clearing zones in red starch (see below).
 2. From the solid phase plates containing dilutions of breakout stocks, core several isolated plaques, each into 500 uL SM with 50 uL CHCl₃. This is referred to as the isolate stock.
 3. The isolate stocks can then be individually tested on BODIPY starch using methods described above. This step can be skipped if the plaque that was cored in step 2 produced a clearing zone in the red starch background. The isolate stocks were then be individually tested on BODIPY starch using methods described above. However, this step may be skipped if the plaque that was cored in step 2 produced a clearing zone in the red starch background.

Excisions

- Day 1
1. In a Falcon 2059 tube, mix 200 uL OD1 XL1-Blue MRF' host, 100 uL lambda isolate stock and 1 uL ExAssist phage stock.
 2. Incubate in 37°C shaker for 15 minutes.
 3. Add 3 mL NZY medium.
 4. Incubate in 30°C shaker overnight.
- Day 2
1. Heat to excision tube to 70°C for 20 minutes.

2. Centrifuge 1000 x g for 10 minutes.
3. In a Falcon 2059 tube, combine 50 uL supernatant with 200 uL EXP505 OD1 host.
4. Incubate in 37°C shaker for 15 minutes.
- 5 5. Add 300 uL SOB medium.
6. Incubate in 37C shaker for 30-45 minutes.
7. Plate 50 uL on large LB_{Kan50} plate using sterile glass beads. If the plates are "dry", extra SOB medium can be added to help disburse the cells.
8. Incubate plate at 30°C for at least 24 hours.
- 10 9. Culture an isolate for sequencing and/or RFLP.

Growth at 30°C reduces plasmid copy number and is used to mitigate the apparent toxicity of some amylase clones.

Contaminants That Form Clearing Zones in Red Starch

- When using red starch on solid medium to assay phage for amylase
- 15 activity, it is common to see contaminating colony forming units (cfu) that form clearing zones in the red starch. For pintoed plates, it is important to distinguish amylase-positive phage clones from these contaminants whenever they align with a particular well position. The source of the contaminating microbes is presumably the 2% red starch stock solution, which cannot be sterilized by autoclaving or by filtering after preparation. It is
 - 20 thought that they are opportunistic organisms that survive by metabolizing the red starch. In order to reduce these contaminants, use sterile technique when making 2% red starch solutions and store the stocks either at 4°C or on ice.

EXAMPLE 8: Bioinformatic Analysis

- 25 The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, e.g., by Bioinformatic Analysis.

An initial bioinformatic analysis was made with the known hyper-thermophilic α -amylase sequences. Figure 14a shows an alignment of the sequences some of which have been deposited at the NCBI database. This analysis revealed the

- 30 potential for designing degenerate primers to PCR the entire gene minus its signal sequence (see Figure 14a), yielding potentially novel full-length alpha amylases from a library.

The following libraries were screened by PCR from genomic DNA:

Table 6:

Library #	Name	PCR positive	Subcloned
5	<i>A. lithotropicus</i>	No	
13	<i>Pyrodictium occultum</i>	No	
17	<i>Pyrodictium TAG11</i>	No	Yes
113	<i>Deep sea enrichment</i>	Yes	Yes
170	<i>Deep sea enrichment</i>	Yes	Yes
198	<i>Archaeglobus</i>	No	
206	<i>Acidianus sp</i>	No	
453	<i>Mixed iceland enrich</i>	No	
455	<i>Mixed iceland enrich</i>	Yes	Yes

Figure 14b shows an alignment of the identified sequences and Table 7, illustrated in Figure 18, lists their relative percent identities.

The amino acid identity ranges from about 85-98% identity. Accordingly, these sequences are useful in shuffling of genes as described herein.

Figure 14c shows the nucleic acid alignment of the corresponding polypeptide sequences above. Expression of these amylases in the expression vector pSE420 and the host cell line XL1-Blue showed 1703 and 1706 to have amylase activity.

EXAMPLE 9: Characterization of Library 63 GP-1 alpha amylase pH optimum and specific activity determination

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, e.g., by alpha amylase activity pH optimum and specific activity determination.

In initial experiments, the SEQ ID NO: 81 from *Thermococcus* showed that it was effective in both starch liquefaction for corn wet milling and desizing for textiles. This enzyme has a pH optimum of 4.5 to 5.0. At this lower pH, it is possible to use little or no calcium which lowers overall operating costs and less byproduct formation. In addition, at this low pH, there is decreased chemical usage and ion

exchange load. The industry standard *B. licheniformis* amylase is suboptimal in both thermostability and pH optimum. The 63GP-1 amylase has a higher application specific activity compared to *B. licheniformis* amylase and therefore much less enzyme is required to hydrolyze a ton of starch (as much as 20-fold less enzyme can be used).

5 The pH optimum for the hydrolysis of starch was determined by reacting 50 μ L of the GP-1, 0.35 U/ml, with a 100ml of 1% soluble starch solution (0.0175U/g of starch) for 30 minutes at 95 degrees C. The reducing ends generated in the liquefied starch solution were measured by the neocupronine assay, described herein. The percent hydrolysis of cornstarch was determined by measuring the number of sugar reducing ends
10 produced with the neocupronine assay. Seventy grams of buffer solution (pH4-7) was weighed and 100ppm of calcium was added. Thirty grams of cornstarch was mixed into the buffer solution to form a starch slurry. The enzyme was added and the vessels sealed and incubated at 95 degrees C for 30 minutes with an initial heating rate of six degrees C per minute. A 1 ml sample was extracted from the reaction beakers and analyzed by the
15 neocupronine assay. The optimum for GP-1 was between pH 4.5 and 5, while the commercial *B. licheniformis* amylase performed optimally at about pH 6.0.

EXAMPLE 10: Amylase Ligation Reassembly

The following example describes, inter alia, exemplary methods for
20 determining if a polypeptide is within the scope of the invention, e.g., by the assays described below.

Assay Using RBB-starch

75 μ L of RBB-starch substrate (1% RBB-insoluble corn starch in 50mM NaAc buffer, pH=4.5) was added into each well of a new 96-well plate (V-bottom). Five
25 micro-liters of enzyme lysate was transferred into each well with substrate using Biomek or Zymark. The plates were sealed with aluminum sealing tape and shaken briefly on the shaker. The plates were incubated at 90°C for 30 minutes, followed by cooling at room temperature for about 5 to 10 minutes. One hundred micro-liters of 100% ethanol was added to each well, the plates sealed and shaken briefly on the shaker. The plates were
30 then centrifuged 4000 rpm for 20 minutes using bench-top centrifuge. 100 μ L of the supernatant was transferred into a new 96-well plate (flat bottom) by Biomek and read OD₅₉₅. Controls: SEQ ID NO:81, SEQ ID NO:77, SEQ ID NO:79.

Assay using FITC-starch

Added 50µl of substrate (0.01% FITC-starch in 100mM NaAc buffer, pH=4.5) into each well of a new 384-well plate. Transferred 5µl of enzyme lysate into each well with substrate and incubated the plate at room temperature overnight. The polarization change of the substrate, excitation 485nm, emission 535nm, was read for each well. Controls: SEQ ID NO.: 81, SEQ ID NO.: 77, SEQ ID NO.: 79. Preferably 96 well plates are used for all assays.

Confirmation of new active clones

Each positive clone from screening was grown and induced using a standard protocol. Each clone was examined for growth (i.e., cell density over time), activity at per cell level (RBB-starch assay and liquefaction assay), expression (protein gel) and solubility of protein (by microscope analysis). The confirmed new elevated clones were transferred for fermentation.

15

Example 11: Exemplary protocol for liquefying starch and measuring results

The following example described and exemplary protocol for liquefying starch using selected amylases of the invention.

20 Amylases having a sequence as set forth in SEQ ID NO:10 and SEQ ID NO:4 demonstrated activity on liquefied starch at pH 4.5 or 6.5 using the reaction conditions show below.

Reaction Conditions: 100 mM PO₄ pH 6.5, 1% (w/w) liquefied starch DE 12 at 55°C. Both TLC and HPLC assays were done to verify activity. The data from both 25 assays showed that the clones were active.

pH profiles for the amylases having a sequence as set forth in SEQ ID NO:4 and SEQ ID NO:10 were run using phosphate buffer pHed from 3.0 - 6.5, at 55°C. From the amount of observable hydrolysis, it could be visually said that the clones were more active at certain pH values than at other values at the above indicated reaction 30 conditions:

SEQ ID NO:4 - active from pH 5.0 - 6.5

SEQ ID NO:10 - active from pH 4.5 - 6.5

An exemplary protocol for the saccharification of liquefied starch at pH 6.5:

- Adjust the pH of the liquefied starch to the pH at which the saccharification(s) will be performed. Liquefy starch in 100 mM sodium acetate buffer, pH 4.5 with 100 mM sodium phosphate salts added so that before saccharification, the pH could be adjusted to pH 6.5.
- Weigh 5 gram samples of liquefied starch into tared bottles.
- Use 0.04% (w/w) Optidex L-400 or approximately 400 mL of 1-10 diluted stock Optidex L-400 per 100 grams of liquefied starch.
- Calculate the milligrams of Optidex L-400 contained in the 400 mL of 1-10 diluted stock Optidex L-400. Next, calculate the volume of lysates needed to give the same concentration of enzyme as the Optidex L-400.
- Add enzymes to liquefied starch samples and incubate at desired temperature (50C°). After 18 hours determine DE and prepare a sample for HPLC analysis.

An exemplary DE Determination:

Exemplary Neocuproine Assay:

A 100ml sample was added to 2.0ml of neocuproine solution A (40g/L sodium carbonate, 16g/L glycine, 0.45g/L copper sulfate). To this was added 2.0 ml of neocuproine solution B (1.2g/L neocuproine hydrochloride-Sigma N-1626). The tubes were mixed and heated in a boiling water bath for 12 minutes; cooled, diluted to 10ml volume with DI water and the OD read at 450nm on the spectrophotometer. The glucose equivalent in the sample was extrapolated from the response of a 0.2mg/ml glucose standard run simultaneously.

Exemplary HPLC Analysis:

Saccharification carbohydrate profiles are measured by HPLC (Bio-Rad Aminex HPX-87A column in silver form, 80°C) using refractive index detection. Mobile phase is filtered Millipore water used at a flow rate of 0.7 ml/min. Saccharification samples are diluted 1-10 with acidified DI water (5 drops of 6 M HCl into 200 mL DI water) then filtered through a 0.45 mm syringe filter. Injection volume is 20 uL.

Exemplary TLC:

Reaction products were w/d at hourly timepoints and spotted and dried on a TLC plate. The Plate was then developed in 10:90 water:isopropanol and visualized

with either a vanillin stain or CAM stain and then heated to show reducible sugars. The liquefied starch was partially hydrolyzed to glucose in cases where activity was observed.

EXAMPLE 12: Starch Liquefaction using Amylases of the Invention

5 This example describes an exemplary method of the invention for liquefying starch using amylases of the invention.

Amylase concentrate was prepared from fermentation broths by heat treatment, cell washing, alkaline extraction using microfiltration and ultrafiltration (48% overall yield). The UF concentrate was neutralized with acetic acid and formulated with
10 30% glycerol at pH 4.5. The activity level of the slurry formulation was representative of a commercial product (120U^l/g – 0.5kg/ ton starch).

Standard Amylase Activity Assay

A 1 mL cuvette containing 950 μ L of 50 mM MOPS pH 7.0 containing 5 mM PNP- α -D—hexa-(1 \rightarrow 4)-glucopyranoside was placed in the Peltier temperature
15 controller of the Beckman DU-7400 spectrophotometer preheated to 80°C. The spectrophotometer was blanked at 405nm and 50 μ L of the enzyme solution was added to the cuvette, mixed well and the increase in the OD_{405nm} was monitored over a one-minute interval. The Δ OD_{405nm/min} rate is converted to a standard unit of μ mole/minute from the OD_{405nm} response of 50 μ L of 1 μ mole/mL PNP in 950 mL 50 mM MOPS at pH 7.0 -
20 80°C. One standard Diversa unit of thermostable alpha amylase (DTAA) is equal to the amount of enzyme that will catalyze the release of 1 μ mole/mL/minute of pNP under the defined conditions of the assay.

25 Standard Glucoamylase Activity Assay

A 1 mL cuvette containing 950 μ L of 50 mM MOPS pH 7.0 containing 5 mM pNP- α -D-glucopyranoside was placed in the Peltier temperature controller of the Beckman DU-7400 spectrophotometer preheated to 60°C. The spectrophotometer was blanked at 405nm and 50 μ L of the enzyme solution was added to the cuvette, mixed well
30 and the increase in the OD_{405nm} was monitored over a one-minute interval. The Δ OD_{405nm}/min rate is converted to a standard unit of μ mole/minute from the OD_{405nm}

response of 50 μ L of 1 μ mole/mL pNP in 950 mL 50 mM MOPS at pH 7.0-60°C. One standard Diversa unit of glucoamylase (DGA) is equal to the amount of enzyme that will catalyze the release of 1 μ mole/mL/minute of pNP under the defined conditions of the assay.

5 Dextrose Equivalent Determination

The neocuproine method was used to measure the DE. Selected samples were measured by both the Invention procedure and by a GPC analyst using the GPC Fehlings procedure.

Neocuproine Assay

10 A 100 μ L sample was added to 2.0 ml of neocuproine solution A (40 g/L sodium carbonate, 16g/L glycine, 0.45g/L copper sulfate). To this was added 2.0 ml of neocuproine solution B (1.2 g/L neocuproine hydrochloride-Sigma N-1626). The tubes were mixed and heated in a boiling water bath for 12 minutes; cooled, diluted to 10ml volume with DI water and the OD read at 450 nm on the spectrophotometer. The glucose
15 equivalent in the sample was extrapolated from the response of a 0.2mg/ml glucose standard run simultaneously.

 The starch sample is diluted ~1 to 16 with DI water with the exact dilution recorded. Ten milliliters of the diluted sample was added to 20 mls of DI water. Ten milliliters of Fehlings solution A and B were added to the diluted starch. The sample was
20 boiled for 3 minutes and cooled on ice. Ten milliliters of 30% KI and 10ml of 6N H₂SO₄ was added. The solution was titrated against 0.1N sodium thiosulfate. The titrant volume is recorded and used to calculate the DE.

Residual Starch Determination

 Post-saccharification samples were checked for residual starch using the
25 Staley iodine procedure.

 Twenty grams of sample was weighed into a large weigh dish. 45 μ L of Iodine solution is added to the weigh dish and the starch solution is mixed well. Dark blue indicates the presence of starch, a light blue-green indicates slight starch, light green indicates a trace of starch and yellow-red, absence of starch. Iodine solution is prepared
30 by dissolving 21.25 grams of iodine and 40.0 grams of potassium iodide in one liter of water.

Oligosaccharide Profile

Liquefaction and saccharification carbohydrate profiles were measured by HPLC (Bio-Rad Aminex HPX-87C column in calcium form – 80°C) using refractive index detection.

Gel Permeation Chromatography

The molecular weight distribution was determined by chromatography on a PL Aquagel-OH column with mass detection by refractive index (Waters Model 2410). A Viscotek Model T60 detector was used for continuous viscosity and light scattering measurements.

Capillary Electrophoresis

Beckman Coulter P/ACE MDQ Glycoprotein System – separation of APTS derivatized oligosaccharides on a fused silica capillary - detection by laser-induced fluorescence.

Primary Liquefaction

Line starch directly from the GPC process is pumped into a 60 liter feed tank where pH, DS (dry solids) and calcium level can be adjusted before liquefaction. The amylase is added to the slurry. The 32% DS slurry is pumped at 0.7 liter/minute by a positive displacement pump to the jet - a pressurized mixing chamber where the starch slurry is instantaneously heated to greater than 100C by steam injection. The gelatinized partially liquefied starch is pumped through a network of piping (still under pressure) to give the desired dwell time (5 minutes) at temperature. The pressure is released into a flash tank and samples can be taken. Samples were taken in duplicate.

Secondary Liquefaction

The liquefied starch was collected in one liter glass bottles and held in a water bath at 95C for 90 minutes.

Saccharification

Liquefied starch was cooled to 60C, the pH adjusted to 4.5 and the samples treated with glucoamylase. Saccharification progress was monitored over time by HPLC.

Saccharification

The liquefied syrups produced with each amylase were adjusted to approximately pH 2.5 with 6N HCl immediately after the 90 minute secondary liquefaction to inactivate any residual amylase. The syrups were then adjusted to pH 4.5, placed in a 60°C water bath and saccharified with three levels of glucoamylase. The extent of saccharification was monitored by HPLC at 18-88 hour time points.

The liquefied syrups were saccharified with the standard dosage – 0.04% of a double-strength glucoamylase - and two lower dosages (50% and 25%) to monitor any differences in the saccharification progress.

10 Saccharification Progress - % dextrose development vs time – 0.04% glucoamylase

Amylase	18 hr	24 hr	40 hr	44 hr	88 hr
Commercial	70.2	78.4	86.1	86.7	94.2
SEQ ID NO:437	79	88.6	92.5	92.8	95.3
SEQ ID NO:6	74.1	85.9	91.9	91.6	94.8

Saccharification Progress - % dextrose development vs time – 0.02% glucoamylase

15

Amylase	18 hr	24 hr	40 hr	44 hr	88 hr
<i>B.licheniformis</i> Amylase	54.5	66.7	76.1	77.2	90.9
SEQ ID NO:437	60.1	72	84.8	85.3	93.6
SEQ ID NO:6	57.1	70	84	86.5	92.5

Post-Saccharification sugar profile

In these studies and all previous saccharification studies, the final glucose level achieved after saccharification by amylases of the invention and *B. licheniformis* in liquefied syrups is essentially identical. The DP2 (maltose) level is also similar. These large fragments are poor substrates for glucoamylase and tend to be converted slowly, if at all, into smaller fragments and ultimately, glucose.

	Glucose	DP2	DP3	>DP7
SEQ ID NO:437	95.25	2.39	1.13	0.91
Commercial	94.16	2.10	0.39	2.91
SEQ ID NO:6	94.77	2.27	1.48	0.82

Molecular weight distribution

The molecular weight distribution of syrups liquefied to DE's of 12 and 18 by the exemplary amylases of the invention SEQ ID NO:6 and SEQ ID NO:437, and commercial *Bacillus licheniformis* and commercial *Bacillus stearothermophilus*, were measured by gel permeation chromatography using detection by refractive index, light scattering and viscosity. Both the *B. licheniformis* and *B. stearothermophilus* amylases generate a bimodal distribution – the primary peak centered at 2000, a secondary peak at 32,000 with a shoulder extending past the 160,000 range. The lower molecular weight peak represents approximately 60% of the total mass of the sample. The exemplary amylases of the invention exhibit a single peak at 2000 with very little above 30,000.

HPLC

The DE 12 and 18 syrups produced by the exemplary amylases of the invention SEQ ID NO:6 and SEQ ID NO:437 and commercial *Bacillus licheniformis* and commercial *Bacillus stearothermophilus* amylases were analyzed by HPLC. Both techniques produce fingerprints characteristic of each class of amylase; the oligosaccharide patterns are different for *B. licheniformis* amylase vs *B. stearothermophilus* amylase vs the exemplary amylases of the invention. The liquefied syrups of the invention (e.g., syrups made by methods of the invention and/or made by enzymes of the invention) exhibit evidence of greater branching in the oligosaccharides.

HPLC only resolve the oligosaccharides in the <DP15 range – larger fragments are not visible in these techniques. *Bacillus* amylases are known to liquefy starch in a manner such that the amylopectin fraction is hydrolyzed less extensively than the amylose fraction. These >DP30 amylopectin fragments are contained in the high molecular weight
5 fraction centered at 32,000 and consequently, little evidence of branching is seen in the HPLC analyses of the *Bacillus* liquefied syrups. The <DP15 oligosaccharides from Invention amylases contain fragments from both amylose and amylopectin.

EXAMPLE 13: Starch Liquefaction at acidic conditions using amylases of the invention

10 The invention provides methods for liquefying starch using amylases of the invention, including amylases active under acidic conditions, e.g., between about pH 4.0 and 5.0, e.g., pH 4.5. The conversion of starch to glucose can be catalyzed by the sequence action of two enzymes: alpha-amylases of the invention to liquefy the starch (e.g., the hydrolysis of high molecular weight glucose polymers to oligosaccharides
15 consisting of 2 to 20 glucose units, typically a dextrose equivalent of 10 to 12, by an amylase of the invention), followed by saccharification with a glycoamylase (which can be a glycoamylase of the invention). In one aspect, processing is in a corn wet milling plant producing a starch slurry having a pH or about 4.0 to 4.5. In one aspect, the pH is raised, e.g., to 5.8 to 6.0 before liquefaction to accommodate an alpha amylase with a low
20 pH activity and stability (which can be an alpha amylase of the invention). In one aspect, amylases of the invention can liquefy starch at pH 4.5 to dextrose equivalents ranging from 12 to 18; in one aspect, using alpha amylases of the invention at levels of about 3 to 6 grams per ton of starch. In this aspect, use of alpha amylases of the invention enables starch liquefaction to be conducted at pH 4.5.

25 In one aspect, starch liquefaction is conducted at pH 4.5 for 5 minutes at 105°C to 90 minutes at 95°C using amylases of the invention. The quantity of enzyme was adjusted in order to adjust a target DE of 12 to 15 after liquefaction. In one aspect, the liquefied starch is then saccharified with a glucoamylase, e.g., an *Aspergillus* glucoamylase, for about 48 hours at about pH 4.5 and 60°C. If the saccharified syrup did
30 not contain at least 95% glucose, the target liquefaction DE was raised and the saccharification repeated until the liquefaction eventually did produce a saccharified syrup containing more than 95% glucose. The amylase protein required to produce a suitable liquefied feedstock for saccharification was determined by PAGE.

EXAMPLE 14: Starch Liquefaction using amylases of the Invention

This example describes an exemplary method for liquefying starch using amylases of the invention and characterizes liquefaction oligosaccharide patterns of the
5 exemplary enzymes of the invention SEQ ID NO:6 and SEQ ID NO:437 (encoded by SEQ ID NO:436) vs commercial *Bacillus licheniformis* and *Bacillus stearothermophilus* amylases. These results compare the saccharification progress and final dextrose levels from syrups generated by enzymes of the invention and commercial amylases.

Three commercial enzymes, Genencor Spezyme AA, and two others all
10 required more than double the recommended dosage to achieve the target Dextrose equivalent (DE). Dextrose equivalent (DE) is the industry standard for measuring the concentration of total reducing sugars, calculated as D-glucose on a dry weight basis. Unhydrolyzed granular starch has a DE of virtually zero, whereas the DE of D-glucose is defined as 100.

These results confirm the "double dosage" effect for all *Bacillus* amylases
15 and gives more credence to the proposal that the observed dosage for SEQ ID NO:437 in the trials is also twice the value which would be required under more normal conditions. The projected "normal" dosage, 60-70 Units/kilo starch at pH 4.5 to reach a 19 DE, is consistent with the laboratory liquefaction data.

The oligosaccharide pattern generated by amylases of the invention is
20 different from the *Bacillus* profiles. The molecular weight distribution for the *Bacillus* amylases (gel permeation chromatography with detection by light scattering and viscosity) is bimodal with a substantial fraction at the very high molecular weight range (>300,000) even at an 18DE. The SEQ ID NO:437 at 18DE exhibits a uniform
25 distribution with nothing greater than 20,000. This is consistent with the lower viscosity for syrups of the invention (e.g., syrups made by methods of the invention, or, made using enzymes of the invention). The DP (degrees of polymerization) profiles as measured by HPLC also reflects this difference in action pattern.

In this study, as well as in the previous studies, the final glucose level after
30 saccharification of amylases of the invention liquefied syrups vs the *Bacillus* syrups is the same for both cases. However, saccharification data from, e.g., GPC studies, confirm that the non-dextrose residuals for the amylases of the invention are different from the *Bacillus* amylase syrups. Although the dextrose and maltose levels are essentially the

same for both, the amylases of the invention have a higher DP3 fraction but lower amount of the “highers” vs. the *Bacillus* enzyme. Consistent with the absence of high molecular weight fragments after liquefaction, the post saccharification syrups of the invention have a lower content of the >DP7 fraction.

5

	Glucose	DP2	DP3	>DP7
SEQ ID NO:2	95.25	2.39	1.13	0.91
Commercial	94.16	2.10	0.39	2.91
SEQ ID NO:6	94.77	2.27	1.48	0.82

SEQ ID NO:437 amylase concentrate was prepared from fermentation broths by heat treatment, cell washing, alkaline extraction using microfiltration and ultrafiltration (48% overall yield). The UF concentrate was neutralized with acetic acid and formulated with 30% glycerol at pH 4.5. The activity level of the slurry formulation was representative of a commercial product (120U1/g – 0.5kg/ ton starch).

Example 15: Alkaline Amylases for Laundry and Autodishwash Applications

In one aspect, the invention provides detergents comprising amylases of the invention, including amylases active under alkaline conditions, and methods of making and using them.

Three alkali-stable amylase enzymes of the invention were compared to and outperformed a commercial benchmark enzyme with respect to features important in laundry and automatic dishwashing (ADW) applications:

- Amylase having a sequence as set forth in SEQ ID NO:212 (encoded by SEQ ID NO:211) outperformed the purified commercial benchmark enzyme in the ADW wash test on starch-coated slides and was very resistant to hydrogen peroxide.
- Amylase having a sequence as set forth in SEQ ID NO:210 (encoded by SEQ ID NO:209) and SEQ ID NO:212 (encoded by SEQ ID NO:211) outperformed the purified commercial benchmark enzyme in the presence of a laundry/ADW formulation using a soluble substrate.
- In the presence of chelators, amylase having a sequence as set forth in SEQ ID NO:439 (encoded by SEQ ID NO:438) was very stable and

amylase having a sequence as set forth in SEQ ID NO:441 (encoded by SEQ ID NO:440) was moderately stable.

- Amylase having a sequence as set forth in SEQ ID NO:210 (encoded by SEQ ID NO:209) and amylase having a sequence as set forth in SEQ ID NO:212 (encoded by SEQ ID NO:211) and amylase having a sequence as set forth in SEQ ID NO:441 (encoded by SEQ ID NO:440) have very alkaline pH optima in the range of pH 10 to 11. Amylase having a sequence as set forth in SEQ ID NO:445 (encoded by SEQ ID NO:444) and having a sequence as set forth in SEQ ID NO:439 (encoded by SEQ ID NO:438) have pH optima around 8 while retaining significant activity at pH 10.
- Amylase having a sequence as set forth in SEQ ID NO:441 (encoded by SEQ ID NO:440) and having a sequence as set forth in SEQ ID NO:439 (encoded by SEQ ID NO:438) were thermophilic, performing best at 65° to 70°C.

Biochemical characterization

Five amylases of the invention, three with alkaline pH optima, were characterized for pH optimum and temperature optimum, as described in Table 1. "SEQ ID NOS:209, 210" refers to an amylase having a sequence as set forth in SEQ ID NO:110, encoded by SEQ ID NO:209, etc.

Table 1

Amylase	pH optimum	Temp. optimum (°C)*
SEQ ID NOS:209, 210	11	55
SEQ ID NOS:211, 212	10	50
SEQ ID NOS:440, 441	10	70
SEQ ID NOS:444, 445	8	40
SEQ ID NOS:438, 439	8	65

Temperature optima were determined at pH 10 for the amylase having a sequence as set forth in SEQ ID NO:210, encoded by SEQ ID NO:209 ("SEQ ID NOS:209, 210"); SEQ ID NOS:211, 212; and SEQ ID NOS:440, 441 and at pH 8 for SEQ ID NOS:444, 445 and SEQ ID NOS:438, 439.

The pH profiles for amylases of the invention compared to the

benchmark enzyme currently used in a commercial laundry/ADW product are presented in Figure 1. All of the enzymes of the invention demonstrated optimal activity between pH 8 and 10, whereas the commercial benchmark enzyme was most active at pH below 8 and had only residual activity at pH 10. Figure 19 shows the pH profile of the tested amylases of the invention and the commercial benchmark enzyme. Purified protein was added to buffers of the indicated pH containing soluble substrate and the activity was measured. Initial rates were calculated over 10 min and converted to a percentage of the maximum rate.

The temperature profiles of enzymes of the invention are presented in Figure 20. Three were most active between temperatures 45°C and 55°C, while the amylase having a sequence as set forth in SEQ ID NO:441 (encoded by SEQ ID NO:440) ("SEQ ID NOS:440, 441") and SEQ ID NOS:438, 439 had optimum activity between 60°C and 70°C. Figure 20 shows the temperature activity profiles of the tested amylases of the invention. Activity of purified protein was measured at pH 10 (SEQ ID NOS:209, 210, SEQ ID NOS:211, 212, SEQ ID NOS:440, 441) or pH 8 (SEQ ID NOS:444, 445, SEQ ID NOS:438, 439) at the indicated temperature. Activity was measured either by a reducing sugar assay or by monitoring the fluorescence at 520 nm (485 nm excitation) when BODIPY-starch was used. Initial rates were calculated and converted to a percentage of the maximum rate.

Application testing

Experiments were designed to assess the activity and stability of the tested alkaline amylases of the invention in laundry/ADW formulations and with the components individually. Figures 21, 22 and 23 present the results of experiments using a soluble starch substrate. Figure 24 presents results using a solid substrate - the industry-standard starch-coated slides.

Amylase having a sequence as set forth in SEQ ID NO:439 (encoded by SEQ ID NO:438) ("SEQ ID NOS:438, 439") was very resistant to the chelator EDTA (Figure 21) and SEQ ID NOS:211, 212 displayed significant resistance to hydrogen peroxide (Figure 22). In contrast, the commercial benchmark enzyme was not functional in the presence of either component under the conditions of the experiments. In the presence of the complete laundry/ADW formulation, SEQ ID NOS:209, 210 and SEQ ID

NOS:211, 212 were much more active on soluble substrate than the commercial benchmark enzyme (Figure 23).

Figure 21 shows enzyme activity in the presence of EDTA. Purified proteins were incubated at 50°C in the presence or absence of 5mM EDTA for the indicated time, after which residual amylase activity was measured using soluble substrate. Activity in the presence of EDTA is expressed as the % of activity in the absence of chelator. Figure 22 shows enzyme activity in the presence of peroxide hydroxide. Purified proteins were incubated at 50°C in the presence or absence of 1M H₂O₂ for the indicated time after which amylase activity was measured using soluble starch. Activity in the presence of peroxide hydroxide is presented as the % of activity in the absence of H₂O₂. Figure 23 shows enzyme activity in the ADW solution (distilled water, hardening solution, bleach, chelators, surfactants) with soluble substrate (BODIPY-starch). Purified proteins reacted with the soluble starch at 40°C in the presence of laundry/ADW formulation. Initial rates were calculated over 5 minutes and expressed as fluorescent units (FU)/s per ng of protein.

The lead performers emerging from the tests on soluble substrate were the amylase having a sequence as set forth in SEQ ID NO:210 (encoded by SEQ ID NO:209) ("SEQ ID NOS:209, 210") and SEQ ID NOS:211, 212. These amylases, along with SEQ ID NOS:440, 441, were compared with the commercial benchmark enzyme in the industry-standard wash test on the starch-coated slides. Results of these experiments are presented in Figure 24. The enzyme having a sequence as set forth in SEQ ID NO:212 (encoded by SEQ ID NO:211) consistently outperformed the purified benchmark enzyme in this test although the formulated benchmark enzyme showed better performance. The nature of the benchmark commercial formulation is unknown, but the purified benchmark enzyme displayed two-fold increase in activity in the presence of Bovine Serum Albumin (BSA). Figure 24 shows the results of the wash tests with starch-coated slides. Purified proteins were incubated with slides at 50°C for 30 min in the presence of ADW solution (distilled water, water hardening solution, bleach, chelators, surfactants). Starch removal was measured comparing weight loss after the enzyme treatment to the initial weight of the slide.

Summary of the characterization of exemplary amylases

The gene encoding the amylase having a sequence as set forth in SEQ ID

NO:212 (encoded by SEQ ID NO:211) ("SEQ ID NOS:211, 212") was isolated from an environmental library collected from a biotope with a pH of 11.0 and temp of 41°C. The amylase encoded by this gene belongs to Family I and does not contain any known Starch/Carbohydrate Binding Domains. The protein has been expressed with and without
5 a C-terminal histidine tag, and in non-glycosylating and a glycosylating host. Enzyme expressed in all of these Host/His tag combinations have pH optima around 10 and temperature optima around 50°C (experiments represented by Figures 19 and 20). The enzyme expressed in the glycosylating host with a His tag was used for the experiments represented by Figures 21 through 24. The presence of the His tag does not seem to
10 affect specific activity, however, glycosylation appears to result in a slightly lower specific activity than that without glycosylation.

In summary:

- The best performer in these application assays was the amylase having a sequence as set forth in SEQ ID NO:212 (encoded by SEQ ID NO:211) ("SEQ ID
15 NOS:211, 212").
- pH and temperature optima of SEQ ID NOS:211, 212 meet the requirements for laundry/ADW applications and SEQ ID NOS:211, 212, with proper formulation, should exceed the performance of the commercial benchmark enzyme.

20 Example 16: Identification and characterization of a thermostable glucoamylase

The following example describes the identification and characterization of an exemplary thermostable amylase of the invention having glucoamylase activity.

Nucleic Acid Extraction: The filamentous fungus *Thermomyces lanuginosus* ATCC 200065 was grown in liquid culture in Potato Dextrose Medium
25 (Difco, BD, Franklin Lakes, NJ). Biomass was collected and high molecular weight genomic DNA was isolated using DNEASY™ (DNeasy) Plant Maxi Kit (Qiagen, Valencia, CA) using standard protocols. Total RNA was also isolated using RNEASY™ (RNeasy) Plant Mini Kit (Qiagen) using standard protocols.

Library Construction: *Thermomyces* genomic DNA was partially digested
30 with restriction enzymes and fragments between 1-10 kb were purified for construction of a genome library. The fragments were ligated into the vector Lambda Zap Express™ (Stratagene, San Diego, CA) and packaged into infectable phage as per manufacturer's instructions.

Library Screening: The above lambda library was used to infect XL1 Blue MRF' cells (Stratagene) in top agar. Approximately 50,000 pfu of phage was added to 600 ul of cells OD600=1. The mixture was incubated at 37°C for 15 minutes in a water bath and then added to 6 ml melted 0.7% top agar and plated onto NZY agar plates. The plate was then incubated overnight at 39°C. A nylon circle (F. Hoffmann-La Roche Ltd., Basel Switzerland) was laid on top of the resulting plaque lawn and lifted back up with some of the phage adhering to the nylon. The nylon was submerged in 1.5M NaCl, 0.5M NaOH for 2 minutes, 1.5M NaCl, 0.5M Tris pH 7.6 for 5 minutes and 2X SSC, 0.2M Tris pH7.6 for 30 seconds. The nylon filter was then UV crosslinked in a Stratagene crosslinker.

A 639 bp PCR fragment from the glucoamylase gene of *Aspergillus niger* was generated from *Aspergillus* genomic DNA for use as a probe. The primers (5'-GCGACCTTGGATTCATGGTTGAGCAAC-3' (SEQ ID NO:595) and 5'-CACAATAGAGACGAAGCCATCGGCGAA-3') (SEQ ID NO:596) were used in the PCR reaction that utilized the Expand High Fidelity PCR Kit™ (Roche) using 30 cycles of 95°C for 20 seconds, 55°C for 30 seconds, and 72°C for 1 minute in a thermal cycler. This PCR fragment is composed of exons 1-4 of the *Aspergillus* glucoamylase gene. The isolated PCR fragment was prepared as a radioactive probe using the Prime It Kit™ (Stratagene) following manufacturer's instructions.

The library filter lifts were washed in a prehybridization solution (DIG Easy Hyb™, Roche) for two hours at 42°C in a hybridization oven (Robbins). The probe was added to 15ml fresh DIG Easy Hyb™ and used to replace the prehybridization solution. The filter was washed with probe overnight at 45°C. The probe was then removed and the filter washed once with 2X SSC, 0.1% SDS for 15 minutes, and twice with 0.1X SSC, 0.1% SDS for 15 minutes each. The nylon filter was then exposed to x-ray film overnight at -80C. Following developing, hybridization spots on the x-ray film were used to identify clones from the original plate. An agar plug was taken from the plate where the spots lined up and suspended in SM buffer to release the phage into solution. Several isolated plaques corresponding to *Thermomyces* genomic fragments containing all or part of the glucoamylase gene were thus isolated.

100 ul of isolated phage stock was added to 200ul XL-1 Blue MRF' cells (Stratagene) and 1 ul ExAssist™ helper phage (Stratagene). The mixture was incubated at 37C for 15 minutes, and 3 ml of 2X YT media was added. This was then incubated at

37°C with shaking for 2.5 hours. The mix was then heated for 20 minutes at 70°C and cooled on ice. 100 ul of the mix was removed and added to 200 ul SOLR cells (Stratagene) and incubated at 37°C for 15 minutes. 50 ul was plated on LB kanamycin (50 ug/ml) plates and incubated overnight at 37°C. Resulting colonies contain cloned

5 genomic fragments in the plasmid pBK-CMV.

Sequencing: DNA sequencing on candidate clones were performed with the BigDye Terminator Cycle Sequencing Version 2.0 Kit™ (Applied Biosystems, Foster City, CA) and a 3700 DNA Analyzer™ (Applied Biosystems) using manufacturer's protocols. A genomic clone was identified with a 4.1 kb insert that contained the entire
10 glucoamylase gene and flanking sequence, as set forth in SEQ ID NO:587. Potential introns were identified by comparing this sequence with consensus sequences for introns in *Aspergillus*. The *Thermomyces lanuginosus* nucleotide sequence has an open reading frame encoding a protein of 617 amino acids, interrupted by four introns of 64 bp, 61 bp, 80 bp, and 57 bp respectively.

15 cDNA Synthesis: The primers 5'-

ATGTTATTCCAACCGACTTTGTGCGC-3' (SEQ ID NO:597) and 5'-

TCATCGCCACCAAGAATTCACGGTG-3' (SEQ ID NO:598) were used in a cDNA synthesis reaction using a Thermoscript rtPCR Kit™ (Invitrogen) using manufacturer's protocols. The template for synthesis was total RNA isolated from *Thermomyces*

20 *lanuginosus* cells growing on potato dextrose media (Difco). An 1854 bp fragment from the reaction was isolated, cloned and sequenced, with the nucleic acid sequence set forth in SEQ ID NO:593.

Expression Cloning: Primers were designed for overexpression of *Thermomyces* glucoamylase in the host *Pichia pastoris*. The primers 5'-

25 GTCTCGAGAAAAGAGCAACGGGCTCGCTCGAC-3' (SEQ ID NO:599) and 5'-

GTTCTAGATCATCGCCACCAAGAATTCACGGT-3' (SEQ ID NO:600) were used to generate a PCR fragment using the cDNA clone as a template using 30 cycles of 95°C for 20 seconds, 55°C for 30 seconds, 72°C for 2 minutes, using Expand High Fidelity PCR Kit™ (Roche) and manufacturer's protocols. The PCR fragment was digested with the

30 restriction enzymes Xho I and Xba I and ligated into the corresponding restriction sites of the plasmid pPIC Z alpha (Invitrogen). The construct was transformed into *Pichia pastoris* Strain X-33 (Invitrogen) where the construct integrates stably into the *Pichia* chromosome. Selection was based on resistance to zeocin. This construct was designed

such that the *Pichia* clone can be induced with methanol to secrete the mature *Thermomyces* glucoamylase into the media.

A 1-liter culture of the *Pichia* expression clone was inoculated with an overnight starter culture in BMGY and grown overnight at 30°C in a shake flask. The cells were collected by centrifugation the following day and resuspended in 1 liter of BMMY. The cells were cultured at 30°C in a shake flask for 3 days with methanol added to 0.5% final every 24 hours. The media containing the expressed glucoamylase enzyme was then collected and tested in a glucoamylase activity assay and SDS PAGE electrophoresed using standard protocols to determine the protein size.

Primers were also designed for overexpression of the *Thermomyces* glucoamylase gene in *Escherichia coli*. The primers (SEQ ID NO:601) 5'-GTCCATGGCAACGGGCTCGCTCGAC-3' and (SEQ ID NO:602) 5'-GTTCTAGATCATCGCCACCAAGAATTCACGGT-3' were used to generate a PCR product as before, from the cDNA template. The PCR fragment was digested with the restriction enzymes Nco I and Xba I and ligated into corresponding restriction sites of the plasmid pSE420 (Invitrogen). The construct was transformed into *Escherichia coli* Strain XL-1 Blue MR (Stratagene). Selection for the plasmid was based on ampicillin resistance. The glucoamylase gene is under the control of the lac-z promoter in this plasmid vector and is induced with IPTG (isopropyl-thio-galactopyranoside). The construct was designed such that the mature glucoamylase gene will be expressed within the *Escherichia* cell and will contain an extra methionine residue at the N-terminus.

Standard assay: Enzyme aliquots were added to a solution of 5 mM buffer, 3 mM malto-oligosaccharides (Sigma, M-3639) in a waterbath. 100 ul aliquots removed at time points to 200 ul glucose oxidase reagent (Sigma, GAGO-20) and incubated 37°C, 30 min. The reaction was stopped with addition of 12 N sulfuric acid and the absorbance at 540 nm determined. The full-length version of the enzyme (SEQ ID NO:594) was tested for pH, temperature and substrate utilization. As noted below, data demonstrated that the pH optimum to be around pH 5.5. Data demonstrated that the enzyme (SEQ ID NO:8) is stable at 70°C with a rapid irreversible loss of activity between 70°C and 75°C. Data demonstrated that the enzyme (SEQ ID NO:594) hydrolyses oligosaccharides down to maltose with the rate of hydrolysis being higher for longer saccharides. The rate in cleaving 1,6 linkages is much slower than 1,4 as observed in the substrate panose which has a 1,6 linkage at the non-reducing end. The catalytic domain version appears to be less

thermostable. The enzyme (SEQ ID NO:594) has a good rate of hydrolysis at 50°C but appears to die at 70°C.

Activity Assay: Enzyme (SEQ ID NO:594) activity was measured by the release of free glucose from an oligo-dextrin substrate. The liberated glucose was then
5 oxidized in a coupled reaction resulting in a colored product. An enzyme (SEQ ID NO:594) aliquot added to solution of 5mM buffer, 3mM malto-oligosaccharides (Sigma, M-3639) in a water bath. 100 ul aliquots removed at time points to 200ul glucose oxidase reagent (Sigma, GAGO-20) and incubated 37°C, 30 min. The reaction was stopped with addition of 12 N sulfuric acid and the absorbance at 540 nm determined. Time points
10 were then plotted to determine the relative rate for the reaction.

pH Profile: Acetate buffer (pH 4.0, 4.5, 5.0, and 5.4) as well as phosphate buffer (pH 6.2, 7.0, 8.1) were used in an activity assay to determine the relative rate for the glucoamylase (SEQ ID NO:594) at each pH. The rates were then plotted, as illustrated in Figure 5. The enzyme (SEQ ID NO:594) appears to have maximal activity
15 around pH 5.5.

Temperature Profile: The relative rate of the enzyme (SEQ ID NO:594) at various temperatures (50°C, 60°C, 70°C, 80°C, and 85°C) was determined in acetate buffer pH 5.3. The rates are plotted in Figure 6. The enzyme (SEQ ID NO:594) appears to have maximal activity at 70°C, above which there is a rapid loss of activity.

20 Temperature Stability Data: Enzyme (SEQ ID NO:594) was added to 5 mM acetate buffer at the indicated temperature. Enzyme (SEQ ID NO:594) aliquots were removed to ice at 4 minute intervals. The aliquots were then tested for activity on substrate for 20 minutes at 70°C, and the data is illustrated in Figure 7.

Substrate Utilization: The dextrans maltose (G2), maltotriose (G3), panose
25 (Pan), maltotetraose (G4), and maltoheptaose (G7), were substituted for the malto-oligosaccharides in the activity assay to test for substrate utilization of the glucoamylase (SEQ ID NO:594). Rate of glucose release for various substrates tested in 5 mM acetate buffer, 70°C. Substrates tested: maltose, maltotriose, panose, maltotetraose, and maltoheptaose, were all at 3 mM. The assay was then plotted in Figure 8. Then enzyme
30 (SEQ ID NO:594) was able to hydrolyze straight-chain (1,4 linkages) dextrans down to maltose with a higher rate for the longer dextrans. The enzyme (SEQ ID NO:594) demonstrated low activity on 1,6 linkages as demonstrated by the substrate panose.

EXAMPLE 17: Glucoamylase Activity Assay: BCA Reducing Ends Assay

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention, for example, by a BCA reducing ends assay. Glucoamylase activity can be determined using the following methodology.

1. Prepare 2 substrate solutions, as follows:

- a) 2% soluble starch (potato) pH 8 solution by dissolving 2 gm potato starch in 100 ml 100 mM sodium phosphate pH 8).
- b) 2% soluble starch (potato) pH 10 solution by dissolving 2 gm potato starch in 100 ml 100 mM sodium carbonate.

Heat both solutions in a boiling water bath, while mixing, for 30-40 minutes until starch dissolves.

2. Prepare Solution A from 64 mg/ml sodium carbonate monohydrate, 24 mg/ml sodium bicarbonate and 1.95 mg/ml BCA (4,4'-dicarboxy-2,2'-biquinoline disodium salt (Sigma Chemical cat # D-8284). Added above to dH₂O.

3. Prepare solution B by combining 1.24 mg/ml cupric sulfate pentahydrate and 1.26 mg/ml L-serine. Add mixture to dH₂O.

4. Prepare a working reagent of a 1:1 ration of solutions A and B.

5. Prepare a Maltose standard solution of 10 mM Maltose in dH₂O, where the 10 mM maltose is combined in 2% soluble starch at desired pH to a final concentration of 0, 100, 200, 300, 400, 600 μ M. The standard curve will be generated for each set of time-points. Since the curve is determined by adding 10 μ l of the standards to the working reagent it works out to 0, 1, 2, 3, 4, 6 nmole maltose.

6. Aliquot 1 ml of substrate solution into microcentrifuge tubes, equilibrate to desired temperature (5 min) in heat block or heated water bath. Add 50 μ l of enzyme solution to the inside of the tube lid.

7. While solution is equilibrating mix 5 ml of both solution A & B. Aliquot 100 μ l to 96 well PCR plate. Set plate on ice.

8. After 5 minute temperature equilibration, close lid on tubes, invert and vortex 3 times. Immediately aliquot 10 μ l into plate as t=0 (zero time point). Leave enzyme mixture in heat block and aliquot 10 μ l at each desired time point (e.g. 0, 5, 10, 15, 20, 30 minutes).

9. Ensure that 12 wells are left empty (only working reagent aliquotted) for the addition of 10 ul of standards, for the standard curve.

10. When all time points are collected and standards are added, cover plate and heated to 80° C for 35 min. Cool plate on ice for 10 min. Add 100 ul H₂O to
5 all wells. Mix and aliquot 100 ul into flat bottomed 96-well plate and read absorbance at 560 nm.

11. Zero each sample's time points against its own t=0 (subtract the average t=0 A₅₆₀ value from other average A₅₆₀ values). Convert the A₅₆₀_(experimental) to umole (Divide A₅₆₀_(experimental) by the slope of the standard curve (A₅₆₀/umole).

10 Generate a slope of the time points and the umole (in umole/min), multiply by 100 (as the umole value only accounts for the 10 ul used in the assay, not the amount made in the 1ml rxn). To get the specific activity divide the slope (in umole/min) by the mg of protein. All points should be done at a minimum in duplicate with three being best. Divide
protein concentration (mg/ml) by any dilution to get mg used in assay. Divide the above
15 slope by mg used in assay to get specific activity. See for example, Wong (2000) J. Agric. Food Chem. 48:4540-4543; Fox (1991) Anal. Biochem. 195, 93-96.

EXAMPLE 18: Screening for Glucoamylase activity

The following example describes an exemplary method for determining if a polypeptide is within the scope of the invention. Glucoamylase activity of clones can
20 be assessed by a number of methods known in the art. The following is the general methodology that can be used.

The number of plaques screened, per plate, can be approximately 10,000 pfu's. For each DNA library: about 50,000 plaques per isolated library and 200,000 plaques per non-isolated library can be screened depending upon the pfu titer for the λ
25 Zap Express amplified lysate.

Titer determination of Lambda Library

8) μ L of Lambda Zap Express amplified library stock added to 600 μ L *E. coli* MRF' cells (OD₆₀₀=1.0). To dilute MRF' stock, 10mM MgSO₄ is used.

9) Incubate at 37°C for 15 minutes.

30 10) Transfer suspension to 5-6mL of NZY top agar at 50 °C and gently mix.

11) Immediately pour agar solution onto large (150mm) NZY media plate.

12) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.

13) Incubate the plate at 39 °C for 8-12 hours.

14) Number of plaques is approximated. Phage titer determined to give 10,000 pfu/plate.

Dilute an aliquot of Library phage with SM buffer if needed.

Substrate screening

5 13) Lambda Zap Express (50,000 pfu) from amplified library added to 600µL of *E. coli* MRF' cells (OD600=1.0). For non-environment libraries, prepare 4 tubes (50,000 pfu per tube).

14) Incubate at 37 °C for 15 minutes.

10 15) While phage/cell suspension are incubating, 1.0mL of red starch substrate (1.2% w/v) is added to 6.0mL NZY top agar at 50 °C and mixed thoroughly. Keep solution at 50°C until needed.

16) Transfer 1/5 (10,000 pfu) of the cell suspension to substrate/top agar solution and gently mixed.

17) Solution is immediately poured onto large (150mm) NZY media plate.

15 18) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.

19) Repeat procedures 4-6 four times for the rest of the cell suspension (1/5 of the suspension each time).

20) Incubate plates at 39°C for 8-12 hours.

21) Plate observed for clearing zones (halos) around plaques.

20 22) Plaques with halos are cored out of agar and transferred to a sterile micro tube. A large bore 200µL pipette tip works well to remove (core) the agar plug containing the desired plaque.

23) Phages are re-suspended in 500µL SM buffer. 20µL Chloroform is added to inhibit any further cell growth.

25 24) Pure phage suspension is incubated at room temperature for 4 hours or overnight before next step.

Isolation of pure clones

12) 10µL of re-suspended phage suspension is added to 500µL of *E. coli* MRF' cells (OD600=1.0).

30 13) Incubate at 37°C for 15 minutes.

- 14) While phage/cell suspension is incubating, 1mL of red starch substrate (1.2% w/v) is added to 6.0mL NZY top agar at 50 °C and mixed thoroughly. Keep solution at 50 °C until needed.
- 15) Cell suspension is transferred to substrate/top agar solution and gently mixed.
- 5 16) Solution is immediately poured onto large (150mm) NZY media plate.
- 17) Allow top agar to solidify completely (approximately 30 minutes), then invert plate.
- 18) Plate incubated at 39°C for 8-12 hours.
- 19) Plate observed for a clearing zone (halo) around a single plaque (pure clone). If a single plaque cannot be isolated, adjust titer and re-plate phage suspension.
- 10 20) Single plaque with halo is cored out of agar and transferred to a sterile micro tube. A large bore 200µL pipette tip works well to remove (core) the agar plug containing the desired plaque. To amplify the titer, core 5 single active plaques into a micro tube.
- 21) Phages are re-suspended in 500µL SM buffer. 20µL Chloroform is added to inhibit any further cell growth.
- 15 22) Pure phage suspension is incubated at room temperature for 4 hours or overnight before next step. The pure phage suspension is stored at -80 °C by adding DMSO into the phage suspension (7% v/v).

Excision of pure clone

- 17) 100µL of pure phage suspension is added to 200µL *E. coli* MRF' cells (OD600=1.0).
- 20 To this, 1.0µL of ExAssist helper phage (>1 x 10⁶ pfu/mL; Stratagene) is added. Use 2059 Falcon tube for excision.
- 18) Suspension is incubated at 37°C for 15 minutes.
- 19) 3.0 mL of 2 x YT media is added to cell suspension.
- 20) Incubate at 30 °C for at least 6 hours or overnight while shaking.
- 25 21) Tube transferred to 70°C for 20 minutes. The phagemid suspension can be stored at 4°C for 1 to 2 months.
- 22) 100 µL of phagemid suspension transferred to a micro tube containing 200µL of *E. coli* Exp 505 cells (OD600=1.0).
- 23) Suspension incubated at 37 °C for 15 minutes.
- 30 24) 300µL of SOB is added to the suspension.
- 25) Suspension is incubated at 37°C for 30 to 45 minutes.

- 26) 100 μ L of suspension is transferred to a small (90mm) LB media plate containing Kanamycin (LB media with Kanamycin 50 μ g/mL) for Zap Express DNA libraries or Ampicillin (LB media with Kanamycin 100 μ g/mL) for Zap II DNA libraries.
- 27) The rest of suspension is transferred to another small LB media plate.
- 5 28) Use sterile glass beads to evenly distribute suspension on the plate.
- 29) Plates are incubated at 30°C for 12 to 24 hours.
- 30) Plate observed for colonies.
- 31) Inoculate single colony into LB liquid media containing suitable antibiotic and incubate at 30 °C for 12 to 24 hours.
- 10 32) Glycerol stock can be prepared by adding 80% glycerol into liquid culture (15% v/v) and stored at -80 °C.

Activity verification

- 7) 50 μ L of liquid culture is transferred to a micro tube. Add 500 μ L of 8% pH7 Amylopectin Azure into the same tube. Prepare 2 tubes for each clone.
- 15 8) Activity is tested at 50°C for 3 hours and overnight. Use pH 7 buffer as control.
- 9) Cool the test specimen at ice-water bath for 5 minutes.
- 10) Add 750 μ L of Ethanol and mixed thoroughly.
- 11) Centrifuge at 13000 rpm (16000 g's) for 5 minutes.
- 12) Measure OD of the supernatant at 595nm.

20 RFLP analysis

- 13) 1.0mL of liquid culture is transferred to a sterile micro tube.
- 14) Centrifuge at 13200 rpm (16000 g's) for 1 minute.
- 15) Discard the supernatant. Add another 1.0 mL of liquid culture into the same sterile micro tube.
- 25 16) Centrifuge at 13200 rpm (16000 g's) for 1 minute.
- 17) Discard the supernatant.
- 18) Follow QIAprep spin mini kit protocol for plasmid isolation.
- 19) Check DNA concentration using BioPhotometer.
- 20) Use Sac I and Kpn I for first double digestion. Incubate at 37 °C for 1 hour.
- 30 21) Use Pst I and Xho I for second double digestion. Incubate at 37 °C for 1 hour.
- 22) Add Loading dye into the digested sample.
- 23) Run the digested sample on a 1.0% agarose gel for 1-1.5 hours at 120 volts.

24) View gel with gel imager. All clones with a different digest pattern will be sent for sequence analysis.

EXAMPLE 19: Assay for glucoamylases

The following example describes an exemplary method for determining if
5 a polypeptide is within the scope of the invention.

Preparation Of Host Cultures

5. Start an overnight culture of XL1-Blue MRF' host cells. Use a single colony from a streak plate to inoculate 10 mL LB supplemented with 20 ug/mL tetracycline. Grow overnight culture shaking at 37°C for at least 16 hours.
- 10 6. Using aseptic technique, inoculate a fresh 100 mL of LB_{Tet} day culture with XL1-Blue MRF' host from the overnight LB_{Tet} culture.
7. Grow in a 37°C shaker until the OD reaches 0.75 – 1.0.
8. Pellet host cells at 1000 x g for 10 minutes and gently resuspend in 10 mM MgSO₄ at OD5.
- 15 9. Dilute a small amount of host cells to OD1 for use in titering and pintoooling.
10. Host preparations can be used for up to 1 week when stored on ice or at 4°C.

-To shorten growth time for the day culture, use ½X the usual Tet concentration in LB (½X = 10 ug/mL), or omit the antibiotic altogether.

-Do not use NZY when selecting with Tetracycline. The high Mg⁺⁺
20 concentration in NZY medium renders Tet inactive.

Titering Lambda Libraries

11. Place three sterile microfuge tubes in a rack.
12. Aliquot 995 uL prepared host cells in one tube and 45 uL prepared OD1 host cells into each of the two remaining tubes.
- 25 13. Add 5 uL of lambda library to the tube containing 995 uL host cells and mix by vortexing. This results in a dilution factor of 200.
14. Prepare 1/2,000 and 1/20,000 dilutions by consecutively adding 5 uL of previous dilution to the remaining two tubes containing 45 uL prepared host cells. Mix by vortexing after each dilution was made.
- 30 15. Allow phage to adsorb to host by incubating at 37°C for 15 minutes.

16. Meanwhile, pipet 100 uL of prepared OD1 host cells to each of three Falcon 2059 tubes.

17. Add 5 uL of each dilution to a separate 2059 tube containing host cells.

18. Plate each by adding 3 mL top agar to each tube and quickly pour over 90 mm NZY plates. Ensure a smooth, even distribution before the top agar hardens.

19. Invert plates and incubate at 37°C overnight.

20. Count plaques and calculate titer of the library stock (in plaque forming units (pfu) per uL).

Lambda Microtiter Screening For glucoamylases

Preparation

5. Prepare a sufficient amount of XL1-Blue MRF' host culture, as described above, for the amount of screening planned. A culture of 100 mL is usually sufficient for screening 2-3 libraries.

6. Autoclave several bottles compatible with the QFill2 dispenser. These are the wide-mouth Corning bottles, 250 mL containing a sealing ring around the lip.

7. Make sure there are sufficient amounts of plates, top agar, BODIPY starch, red starch solution, etc. available for the screen.

8. Schedule the Day 2 robot run with a representative from Automation.

Day 1

10. Label the 1536-well plates (black) with library screen and plate number. Tough-Tags™ tube stickers, cut in half width-wise, are ideal for labeling 1536 well plates.

11. Calculate volumes of library, host cells and NZY medium necessary for the screen. This is easily done with an Excel spreadsheet.

12. Combine the calculated volumes of lambda library and OD5 host cells in a sterile 250 mL wide-mouth Corning bottle (containing a sealing ring).

13. Allow adsorption to occur at 37°C for 15 minutes.

14. Add the calculated volume of NZY medium and mix well. This is referred to as the cell-phage-medium suspension.

15. Perform a concomitant titer by combining 50 uL of the cell-phage-medium suspension with 250 uL of OD1 host cells in a Falcon 2059 tube, then plating with 9 mL of top agar onto a 150 mm NZY plate. Incubate concomitant titer plate at 37°C overnight.

16. Load the dispenser with the remainder of the suspension and array each labeled 1536-well plate at 4 uL per well. If the dispenser leaves air bubbles in some wells, they can be removed by centrifuging the plates at 200 x g for 1 minute.

5 17. Add 0.5 uL of positive control phage to well position AD46 of at least two of the assay plates. Use a strong glucoamylase-positive lambda clone for this purpose. The lambda versions of SEQ ID NO.: 113 or SEQ ID NO.: 199 are good choices for positive controls.

18. Incubate assay plates at 37°C overnight in a humidified ($\geq 95\%$) incubator.

10 Day 2

21. Count the pfu on the concomitant titer plate and determine the average seed density per well (in pfu per well).

22. Pintool at least 2 plates of each library screen (preferably the 2 containing positive controls) as follows:

15 a) Prepare 2 host lawn plates to act as a surface on which to pintool: combine 250 uL of OD1 host cells with 2 mL 2% red starch and plate with 9 mL top agar onto 150 mm NZY plates. Hold each plate as level as possible as the top agar solidifies in order to produce an even hue of red across the plate.

20 b) Using a twice flame-sterilized 1536 position pintool, replicate at least 2 of the screening plates onto the host lawn plates.

c) Place the pintoled recipient plates in a laminar flow hood with the lids off for about 15-30 minutes (to vent off excess moisture).

d) Replace the lids and incubate inverted at 37°C overnight.

23. Prepare the 2X BODIPY starch substrate buffer as follows:

25 a) Calculate the total volume of 2X substrate buffer solution needed for all screening plates at 4 uL per well (including any extra deadspace volume required by the dispenser) and measure this amount of 100 mM CAPS pH 10.4 into a vessel appropriate for the dispenser used.

30 b) Retrieve enough 0.5 mg tubes of BODIPY starch to produce the required volume of 2X substrate buffer [calculated in step a) above] at a final concentration of 20-30 ug/mL.

- 5 c) Dissolve each 0.5 mg tube in 50 uL DMSO at room temperature, protected from light, with frequent vortexing. This takes more than 15 minutes; some production lots of BODIPY starch dissolve better than others.
- d) Add 50 uL 100mM CAPS buffer pH 10.4 to each tube and mix by vortexing.
- 5 e) Pool the contents of all tubes and remove any undissolved aggregates by centrifuging for 1 minute at maximum speed in a microfuge.
- f) Add the supernatant to the rest of the 100 mM CAPS buffer measured in step a) above.
- g) Protect the 2X substrate buffer from light by wrapping in foil.
- 10 24. Take plates and substrate buffer to the automation room and program the robot with the following parameters:
- a) dispense 4 uL substrate buffer per well
- b) 1st read at 1 hour post-substrate, 2nd read at 9 hours, and third read at 17 hours; with 37°C incubation between reads
- 15 c) excitation filter: 485 nm; emission filter: 535 nm
- d) set the Spectrafluor gain at 70, or the optimal gain for the batch of 2X substrate buffer prepared.
- e) ensure that the incubator used will protect assay plates from light.

Day 3

- 20 4. Check pintoled plates for clearings in the bacterial lawn at all positions corresponding to wells on the associated assay plate. Also check for clearings in the red starch in any of the pin positions. If plates containing positive controls were used for pintoled, you should be able to see a large clearing zone in the red background. Be wary of contaminants that also form clearing zones in red starch
- 25 (see comment "Contaminants That Form Clearing Zones in Red Starch").
5. Identify putative hits from the data file produced by the robot computer. The KANAL program produced by Engineering simplifies data analysis. As a rule of thumb, a putative hit is characterized as a well having signal intensity rising at least 1.5 fold over background.
- 30 6. For each putative, remove 2 uL from the well and add to a tube containing 500 uL SM buffer and 50 uL CHCl₃. Vortex to mix and store at 4°C. This solution will be referred to hereafter as the 4e-3 stock. The original screening

plates should be stored at 4°C, protected from light, at least until breakouts are complete.

This is the recommended method of breaking out putative hits. It is a liquid phase assay that relies on confirmation of activity on BODIPY starch.

- 5 Alternatively, putative hits can be plated directly onto solid phase plates containing red starch such that 2,000-3,000 pfu per hit are examined for clearing zones. However, inability to observe clearing zones on red starch is not necessarily an indication that a putative hit was a false positive. It would then need to be assayed using the format in which it was originally identified (i.e., liquid phase using BODIPY starch as substrate).
- 10 In addition, very weak positives are more easily identified using the method detailed below.

Day 1

25. In a sterile 50 mL conical tube, combine 0.5 mL OD5 host cells with 45.5 mL NZY. This will be referred to as the host-medium suspension.
- 15 26. For each putative hit to be analyzed, aliquot 1 mL of host-medium suspension into each of 3 three sterile microfuge tubes.
27. Set the 12-channel pipetman in multidispense mode with an aliquot size of 20 uL and an aliquot number of 2x. Mount the pipetman with a clean set of sterile tips.
28. Pour about 1 mL of host-medium suspension into a new sterile solution basin and
20 load the multichannel pipetman.
29. Dispense 20 uL per well into the last row (row P) of a black 384-well plate (12 channels x 2 = 24 wells). This row will be used later for the controls.
30. Expel the remaining liquid in the tips by touching the tips against the surface of the basin and pressing the RESET button on the pipetman. Lay the pipetman down
25 in a way to prevent contamination of the tips. There is no need to change the tips at this point.
31. Pour the remainder of the fluid in the basin into a waste container (like a beaker) taking care to avoid splash-back contamination.
32. For the first putative to be analyzed, take 111 uL of the 4e-3 stock (see Day 2 in
30 *Lambda Microtiter Screening for glucoamylases*) and add it to the first in a set of three tubes containing 1 mL host-medium suspension (step 2). Vortex to mix. This is *Dilution A*.

33. Take 111 uL of Dilution A and add to the next tube in the set. Vortex to mix. This is *Dilution B*.
34. Take 111 uL of Dilution B and add to the last tube in the set. Vortex to mix. This is *Dilution C*. You should now have three dilutions of phage, where concentrations of each differ by a factor of 10.
35. Pour the contents of Dilution C (the most dilute of the 3 samples) into the solution basin and load the multichannel pipetman.
36. Dispense 20 uL per well into the first row of the 384-well plate (12 channels x 2 = 24 wells).
37. Expel the remaining liquid in the tips by touching the tips against the surface of the basin and pressing the RESET button on the pipetman. Lay the pipetman down in a way to prevent contamination of the tips. There is no need to change the tips at this point.
38. Empty the basin as described above.
39. Pour the contents of Dilution B into the same basin and load the multichannel pipetman.
40. Dispense 20 uL per well into the second row of the 384-well plate.
41. Perform steps 13-16 similarly to dispense Dilution A into the third row of the plate.
42. After all three dilutions have been arrayed into the first 3 rows of the plate, discard all tips and the solution basin into the biohazardous waste container.
43. Mount the pipetman with a clean set of sterile tips and open a new sterile solution basin.
44. Repeat steps 8-19 for each remaining putative hit, using remaining rows on the plate up to row O. Five putative hits can be analyzed on one 384-well plate, with the last row (row P) saved for the controls.
45. Add 0.5 uL of each control to a separate well. Use at least 2-3 separate controls, preferably covering a range of activity.
46. Incubate assay plates at 37°C overnight in a humidified ($\geq 95\%$) incubator.
- Day 2
47. Pintool all breakout plates onto a host lawn with red starch using the same method described for Day 2 in *Lambda Microtiter Screening for glucoamylases*, except that a 384 position pintool is used.

48. Prepare the 2X BODIPY starch substrate buffer as follows:
- a) Calculate the total volume of 2X substrate buffer solution needed for all breakout plates at 20 uL per well (including any extra deadspace volume required by the dispenser) and measure this amount of 100 mM CAPS pH 10.4 into a vessel appropriate for the dispenser used.
 - b) Retrieve enough 0.5 mg tubes of BODIPY starch to produce the required volume of 2X substrate buffer [calculated in step a) above] at a final concentration of 20-30 ug/mL.
 - c) Dissolve each 0.5 mg tube in 50 uL DMSO at room temperature, protected from light, with frequent vortexing. This takes more than 15 minutes; some production lots of BODIPY starch dissolve better than others.
 - d) Add 50 uL 100mM CAPS buffer pH 10.4 to each tube and mix by vortexing.
 - e) Pool the contents of all tubes and remove any undissolved aggregates by centrifuging for 1 minute at maximum speed in a microfuge.
 - f) Add the supernatant to the rest of the 100 mM CAPS buffer measured in step a) above.
 - g) Protect the 2X substrate buffer from light by wrapping in foil.
49. Dispense 20 uL per well into all breakout plates.
50. Wrap all plates in aluminum foil and incubate at room temperature for 2-6 hours.
51. Read each plate in the Spectrafluor with the following settings:
- a) fluorescence read (excitation filter: 485 nm; emission filter: 535 nm)
 - b) plate definition: 384 well black
 - c) read from the top
 - d) optimal gain
 - e) number of flashes: 3
52. On the resulting Excel spreadsheet, chart each putative's 3 rows in a separate graph and check for activity. Ensure that the positives controls produced signals over background.
53. For each putative that appears to have a real signal among the wells, harvest a sample from a positive well as follows:
- a) Select a positive well from a row representing the highest initial dilution.
 - b) Transfer 2 uL from that well into a tube containing 500 uL SM and 50 uL CHCl₃. This is referred to as the breakout stock.

c) Store at 4°C.

54. Using methods previously described, plate about 10 uL of each breakout stock onto 150 mm NZY plates using red starch. The objective is to obtain several (at least 20) well-separated plaques from which to core isolates.

5 Day 3

55. Check pintoled plates for an acceptable incidence of clearings in the bacterial lawn corresponding to wells on the associated assay plate. Also check for clearings in the red starch in the positive controls and in any tested putatives. Be wary of contaminants that also form clearing zones in red starch (see below).

10 56. From the solid phase plates containing dilutions of breakout stocks, core several isolated plaques, each into 500 uL SM with 50 uL CHCl₃. This is referred to as the isolate stock.

15 57. The isolate stocks can then be individually tested on BODIPY starch using methods described above. This step can be skipped if the plaque that was cored in step 2 produced a clearing zone in the red starch background. The isolate stocks were then be individually tested on BODIPY starch using methods described above. However, this step may be skipped if the plaque that was cored in step 2 produced a clearing zone in the red starch background.

Excisions

20 Day 1

58. In a Falcon 2059 tube, mix 200 uL OD1 XL1-Blue MRF⁺ host, 100 uL lambda isolate stock and 1 uL ExAssist phage stock.

59. Incubate in 37°C shaker for 15 minutes.

60. Add 3 mL NZY medium.

25 61. Incubate in 30°C shaker overnight.

Day 2

10. Heat to excision tube to 70°C for 20 minutes.

11. Centrifuge 1000 x g for 10 minutes.

12. In a Falcon 2059 tube, combine 50 uL supernatant with 200 uL EXP505 OD1
30 host.

13. Incubate in 37°C shaker for 15 minutes.

14. Add 300 uL SOB medium.

15. Incubate in 37C shaker for 30-45 minutes.

16. Plate 50 uL on large LB_{Kan50} plate using sterile glass beads. If the plates are "dry", extra SOB medium can be added to help disburse the cells.

17. Incubate plate at 30°C for at least 24 hours.

5 18. Culture an isolate for sequencing and/or RFLP.

Growth at 30°C reduces plasmid copy number and is used to mitigate the apparent toxicity of some glucoamylase clones.

Contaminants That Form Clearing Zones in Red Starch

When using red starch on solid medium to assay phage for glucoamylase
10 activity, it is common to see contaminating colony forming units (cfu) that form clearing zones in the red starch. For pintoed plates, it is important to distinguish glucoamylase-positive phage clones from these contaminants whenever they align with a particular well position. The source of the contaminating microbes is presumably the 2% red starch stock solution, which cannot be sterilized by autoclaving or by filtering after preparation. It is
15 thought that they are opportunistic organisms that survive by metabolizing the red starch. In order to reduce these contaminants, use sterile technique when making 2% red starch solutions and store the stocks either at 4°C or on ice.

20 Assay Using RBB-starch

75µl of RBB-starch substrate (1% RBB-insoluble corn starch in 50mM NaAc buffer, pH=4.5) can be added into each well of a new 96-well plate (V-bottom). Five micro-liters of enzyme lysate can be transferred into each well with substrate using Biomek or Zymark. The plates can be sealed with aluminum sealing tape and shaken
25 briefly on the shaker. The plates can be incubated at 90°C for 30 minutes, followed by cooling at room temperature for about 5 to 10 minutes. One hundred micro-liters of 100% ethanol is added to each well, the plates sealed and shaken briefly on the shaker. The plates are then centrifuged 4000rpm for 20 minutes using bench-top centrifuge. 100µl of the supernatant is transferred into a new 96-well plate (flat bottom) by Biomek and read
30 OD₅₉₅.

Assay using FITC-starch

Add 50µl of substrate (0.01% FITC-starch in 100mM NaAc buffer, pH=4.5) into each well of a new 384-well plate. Transfer 5µl of enzyme lysate into each well with substrate and incubate the plate at room temperature overnight. The polarization change of the substrate, excitation 485nm, emission 535nm, is read for each well. 96 well plates can be used for all assays.

Example 20: Exemplary protocol for liquefying starch and measuring results

The following example described and exemplary protocol for liquefying starch. Reaction Conditions: 100 mM PO₄ pH 6.5, 1% (w/w) liquefied starch DE 12 at 55°C. Both TLC and HPLC assays can be done to verify activity.

An exemplary protocol for the saccharification of liquefied starch at pH 6.5:

- Adjust the pH of the liquefied starch to the pH at which the saccharification(s) will be performed. Liquefy starch in 100 mM sodium acetate buffer, pH 4.5 with 100 mM sodium phosphate salts added so that before saccharification, the pH could be adjusted to pH 6.5.
- Weigh 5 gram samples of liquefied starch into tared bottles.
- Use 0.04% (w/w) Optidex L-400 or approximately 400 mL of 1-10 diluted stock Optidex L-400 per 100 grams of liquefied starch.
- Calculate the milligrams of Optidex L-400 contained in the 400 mL of 1-10 diluted stock Optidex L-400. Next, calculate the volume of lysates needed to give the same concentration of enzyme as the Optidex L-400.
- Add enzymes to liquefied starch samples and incubate at desired temperature (50C°). After 18 hours determine DE and prepare a sample for HPLC analysis.

An exemplary DE Determination:Exemplary Neocuproine Assay:

A 100ml sample can be added to 2.0ml of neocuproine solution A (40g/L sodium carbonate, 16g/L glycine, 0.45g/L copper sulfate). To this can be added 2.0 ml of neocuproine solution B (1.2g/L neocuproine hydrochloride-Sigma N-1626). The tubes can be mixed and heated in a boiling water bath for 12 minutes; cooled, diluted to 10ml volume with DI water and the OD read at 450nm on the spectrophotometer. The glucose

equivalent in the sample can be extrapolated from the response of a 0.2mg/ml glucose standard run simultaneously.

Exemplary HPLC Analysis:

- Saccharification carbohydrate profiles are measured by HPLC (Bio-Rad Aminex HPX-87A column in silver form, 80°C) using refractive index detection. Mobile phase is filtered Millipore water used at a flow rate of 0.7 ml/min. Saccharification samples are diluted 1-10 with acidified DI water (5 drops of 6 M HCl into 200 mL DI water) then filtered through a 0.45 mm syringe filter. Injection volume is 20 uL.

Exemplary TLC:

- Reaction products can be w/d at hourly timepoints and spotted and dried on a TLC plate. The plate can be then developed in 10:90 water:isopropanol and visualized with either a vanillin stain or CAM stain and then heated to show reducible sugars. The liquefied starch can be partially hydrolyzed to glucose in cases where activity was observed.

15 EXAMPLE 21: Starch Liquefaction using glucoamylases

- This example describes an exemplary method of the invention for liquefying starch using glucoamylases of the invention. Glucoamylase concentrate can be prepared from fermentation broths by heat treatment, cell washing, alkaline extraction using microfiltration and ultrafiltration (48% overall yield). The UF concentrate can be neutralized with acetic acid and formulated with 30% glycerol at pH 4.5. The activity level of a commercial product can be about 120 U¹/g – 0.5 kg/ ton starch.

Exemplary glucoamylase activity assay

- A 1 mL cuvette containing 950 µL of 50 mM MOPS pH 7.0 containing 5 mM PNP-α- D—hexa-(1→4)-glucopyranoside is placed in the Peltier temperature controller of the Beckman DU-7400 spectrophotometer preheated to 80°C. The spectrophotometer is blanked at 405nm and 50 µL of the enzyme solution is added to the cuvette, mixed well and the increase in the OD_{405nm} is monitored over a one-minute interval. The ΔOD_{405nm/min} rate is converted to a standard unit of µmole/minute from the OD_{405nm} response of 50 µL of 1 µmole/mL PNP in 950 mL 50 mM MOPS at pH 7.0 - 80°C. One standard unit of thermostable alpha glucoamylase (DTAA) is equal to the

amount of enzyme that will catalyze the release of 1 $\mu\text{mole/mL/minute}$ of pNP under the defined conditions of the assay.

Standard Glucoamylase Activity Assay

A 1 mL cuvette containing 950 μL of 50 mM MOPS pH 7.0 containing 5
 5 mM pNP- α -D-glucopyranoside is placed in the Peltier temperature controller of the Beckman DU-7400 spectrophotometer preheated to 60°C. The spectrophotometer is blanked at 405nm and 50 μL of the enzyme solution is added to the cuvette, mixed well and the increase in the $\text{OD}_{405\text{nm}}$ is monitored over a one-minute interval. The $\Delta\text{OD}_{405\text{nm}}/\text{min}$ rate is converted to a standard unit of $\mu\text{mole/minute}$ from the $\text{OD}_{405\text{nm}}$
 10 response of 50 μL of 1 $\mu\text{mole/mL}$ pNP in 950 mL 50 mM MOPS at pH 7.0-60°C. One standard Diversa unit of glucoamylase (DGA) is equal to the amount of enzyme that will catalyze the release of 1 $\mu\text{mole/mL/minute}$ of pNP under the defined conditions of the assay.

Dextrose Equivalent Determination

15 The neocuproine method is used to measure the DE. Selected samples were measured by both the Invention procedure and by a GPC analyst using the GPC Fehlings procedure.

Neocuproine Assay

A 100 μL sample is added to 2.0 ml of neocuproine solution A (40 g/L
 20 sodium carbonate, 16g/L glycine, 0.45g/L copper sulfate). To this is added 2.0 ml of neocuproine solution B (1.2 g/L neocuproine hydrochloride-Sigma N-1626). The tubes were mixed and heated in a boiling water bath for 12 minutes; cooled, diluted to 10ml volume with DI water and the OD read at 450 nm on the spectrophotometer. The glucose equivalent in the sample is extrapolated from the response of a 0.2mg/ml glucose standard
 25 run simultaneously.

The starch sample is diluted ~1 to 16 with DI water with the exact dilution recorded. Ten milliliters of the diluted sample is added to 20 mls of DI water. Ten milliliters of Fehlings solution A and B were added to the diluted starch. The sample is boiled for 3 minutes and cooled on ice. Ten milliliters of 30% KI and 10ml of 6N H_2SO_4
 30 is added. The solution is titrated against 0.1N sodium thiosulfate. The titrant volume is recorded and used to calculate the DE.

Residual Starch Determination

Post-saccharification samples were checked for residual starch using the Staley iodine procedure.

Twenty grams of sample is weighed into a large weigh dish. 45 μ L of
5 Iodine solution is added to the weigh dish and the starch solution is mixed well. Dark blue indicates the presence of starch, a light blue-green indicates slight starch, light green indicates a trace of starch and yellow-red, absence of starch. Iodine solution is prepared by dissolving 21.25 grams of iodine and 40.0 grams of potassium iodide in one liter of water.

10 Oligosaccharide Profile

Liquefaction and saccharification carbohydrate profiles were measured by HPLC (Bio-Rad Aminex HPX-87C column in calcium form – 80°C) using refractive index detection.

Gel Permeation Chromatography

15 The molecular weight distribution is determined by chromatography on a PL Aquagel-OH column with mass detection by refractive index (Waters Model 2410). A Viscotek Model T60 detector is used for continuous viscosity and light scattering measurements.

20 Capillary Electrophoresis

Beckman Coulter P/ACE MDQ Glycoprotein System – separation of APTS derivatized oligosaccharides on a fused silica capillary - detection by laser-induced fluorescence.

Primary Liquefaction

25 Line starch directly from the GPC process is pumped into a 60 liter feed tank where pH, DS (dry solids) and calcium level can be adjusted before liquefaction. The glucoamylase is added to the slurry. The 32% DS slurry is pumped at 0.7 liter/minute by a positive displacement pump to the jet - a pressurized mixing chamber where the starch slurry is instantaneously heated to greater than 100°C by steam injection.
30 The gelatinized partially liquefied starch is pumped through a network of piping (still under pressure) to give the desired dwell time (5 minutes) at temperature. The pressure is released into a flash tank and samples can be taken. Samples were taken in duplicate.

Secondary Liquefaction

The liquefied starch is collected in one liter glass bottles and held in a water bath at 95°C for 90 minutes.

Saccharification

5 Liquefied starch is cooled to 60°C, the pH adjusted to 4.5 and the samples treated with glucoamylase. Saccharification progress is monitored over time by HPLC.

Saccharification

The liquefied syrups produced with each glucoamylase were adjusted to approximately pH 2.5 with 6N HCl immediately after the 90 minute secondary
10 liquefaction to inactivate any residual glucoamylase. The syrups were then adjusted to pH 4.5, placed in a 60°C water bath and saccharified with three levels of glucoamylase. The extent of saccharification is monitored by HPLC at 18 to 88 hour time points.

The liquefied syrups were saccharified with the standard dosage – 0.04% of a double-strength glucoamylase - and two lower dosages (50% and 25%) to monitor
15 any differences in the saccharification progress.

Saccharification Progress - % dextrose development vs time – 0.04% glucoamylase.

20 EXAMPLE 22: Starch Liquefaction at pH 4.5 using glucoamylases

The conversion of starch to glucose can be catalyzed by the sequence action of two enzymes: amylases (e.g., alpha-amylases), including enzymes of the invention, to liquefy the starch (e.g., the hydrolysis of high molecular weight glucose polymers to oligosaccharides consisting of 2 to 20 glucose units, typically a dextrose
25 equivalent of 10 to 12, by a glucoamylase of the invention), followed by saccharification with a glucoamylase (which can be a glucoamylase of the invention, e.g., SEQ ID NO:594). In one aspect, processing is in a corn wet milling plant producing a starch slurry having a pH or about 4.0 to 4.5. In one aspect, the pH is raised, e.g., to 5.8 to 6.0 before liquefaction to accommodate a glucoamylase with a low pH activity and stability.
30 In one aspect, glucoamylases of the invention can liquefy starch at pH 4.5 to dextrose equivalents ranging from 12 to 18; in one aspect, using glucoamylases of the invention at levels of about 3 to 6 grams per ton of starch. In this aspect, use of glucoamylases of the

invention enables starch liquefaction to be conducted at pH 4.5.

In one aspect, starch liquefaction is conducted at pH 4.5 for 5 minutes at 105°C to 90 minutes at 95°C using glucoamylases of the invention. The quantity of enzyme is adjusted in order to adjust a target DE of 12 to 15 after liquefaction. In one
5 aspect, the liquefied starch is then saccharified with a glucoamylase, e.g., an *Aspergillus* glucoamylase, for about 48 hours at about pH 4.5 and 60°C. If the saccharified syrup did not contain at least 95% glucose, the target liquefaction DE is raised and the saccharification repeated until the liquefaction eventually did produce a saccharified
10 syrup containing more than 95% glucose. The glucoamylase protein required to produce a suitable liquefied feedstock for saccharification is determined by PAGE.

A number of embodiments of the invention have been described. Nevertheless, it will be understood that various modifications may be made without departing from the spirit and scope of the invention. Accordingly, other embodiments are
15 within the scope of the following claims.

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THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. An isolated, synthetic, or recombinant nucleic acid comprising:
 - (a) a nucleic acid sequence having at least 85% sequence identity to the sequence SEQ ID NO:605, wherein the nucleic acid encodes at least one polypeptide having an alpha-amylase activity; or
 - (b) a nucleic acid sequence encoding a polypeptide having at least 85% sequence identity to the sequence SEQ ID NO:606, or an enzymatically active fragment thereof having alpha-amylase activity; or
 - (c) a nucleic acid sequence encoding a polypeptide having at least 85% sequence identity to the sequence SEQ ID NO: 606, that hybridizes under stringent conditions to a nucleic acid comprising SEQ ID NO:605, wherein the nucleic acid encodes a polypeptide having an alpha amylase activity, and the stringent conditions comprise a wash step comprising a wash in 0.2X SSC at a temperature of about 65°C for about 15 minutes; or
 - (d) the nucleic acid sequence SEQ ID NO:605, wherein the nucleic acid encodes at least one polypeptide having an alpha-amylase activity; or
 - (e) a nucleic acid sequence of any one of (a) to (d) encoding an alpha-amylase lacking a signal sequence; or
 - (f) a nucleic acid sequence of any one of (a) to (e) encoding an alpha-amylase having a heterologous sequence or signal sequence; or
 - (g) a fully complementary nucleic acid sequence to any one of (a) to (f).
2. An expression cassette, a vector, or a cloning vehicle comprising a nucleic acid:
 - (a) comprising a sequence according to claim 1; or
 - (b) comprising a sequence according to claim 1, wherein the cloning vehicle comprises a viral vector, a plasmid, a phage, a phagemid, a cosmid, a fosmid, a bacteriophage or an artificial chromosome; or
 - (c) comprising a sequence according to claim 1, wherein the viral vector comprises an adenovirus vector, a retroviral vector or an adeno-associated viral vector; or

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(d) comprising a sequence according to claim 1, comprising a bacterial artificial chromosome (BAC), a plasmid, a bacteriophage P1-derived vector (PAC), a yeast artificial chromosome (YAC), or a mammalian artificial chromosome (MAC).

3. A transformed cell:

(a) comprising a heterologous nucleic acid that comprises a sequence according to claim 1, or an expression cassette, vector, or cloning vehicle according to claim 2; or

(b) comprising a heterologous nucleic acid that comprises a sequence according to claim 1, or an expression cassette, vector, or cloning vehicle according to claim 2, wherein the cell is a bacterial cell, a mammalian cell, a fungal cell, a yeast cell, an insect cell or a plant cell.

4. An isolated, synthetic, or recombinant polypeptide having an alpha-amylase activity comprising:

(a) an amino acid sequence having at least 85% sequence identity to the sequence SEQ ID NO:606, or an enzymatically active fragment thereof having an alpha-amylase activity, wherein the fragment is 40 or more amino acid residues in length; or

(b) an amino acid sequence encoded by a nucleic acid according to claim 1; or

(c) an amino acid sequence according to (a) or (b) lacking a signal sequence; or

(d) an amino acid sequence according to (a), (b) or (c) further comprising a heterologous sequence or heterologous signal sequence; or

(e) an amino acid sequence according to any one of (a) to (d) further comprising a polysaccharide, or wherein the polypeptide comprises at least one glycosylation site.

5. A protein preparation comprising a polypeptide of claim 4, wherein the protein preparation comprises a liquid, a solid, or a gel.

6. A method for hydrolyzing a starch, or removing, or liquefying the starch, comprising the following steps:

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- 5 (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;
- (b) providing a composition comprising a starch; and
- (c) contacting the polypeptide of step (a) with the composition of step (b) under conditions wherein the polypeptide hydrolyzes the starch, or removes or liquefies the starch.
7. A detergent composition comprising a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1.
- 10 8. A food supplement, feed supplement, feed or a food comprising a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1.
9. A composition comprising a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1.
10. A method for textile processing or desizing comprising the following steps:
- 15 (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;
- (b) providing a textile; and
- (c) contacting the polypeptide of step (a) and the textile of step (b) under conditions
20 wherein the alpha-amylase can process or desize the textile.
11. A method for treating a paper, a paper product, a paper pulp or a fiber comprising the following steps:
- (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a
25 nucleic acid of claim 1;
- (b) providing a composition comprising a paper, a paper product, a paper pulp or a fiber; and

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- (c) contacting the polypeptide of step (a) and the composition of step (b) under conditions wherein the polypeptide can treat the paper, paper product, paper pulp or fiber.

5 12. A method for producing a high-maltose or a high-glucose syrup comprising the following steps:

- (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;
- (b) providing a composition comprising a starch; and
- 10 (c) contacting the polypeptide of step (a) and the fabric of step (b) under conditions wherein the polypeptide of step (a) can hydrolyze the composition of step (b), thereby producing a high-maltose or a high-glucose syrup.

13. A method for improving the flow of the starch-containing production fluids, comprising the following steps:

- 15 (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;
- (b) providing a production fluid comprising a polysaccharide; and
- (c) contacting the polypeptide of step (a) and the production fluid of step (b) under
20 conditions wherein the alpha-amylase can hydrolyze the polysaccharide in the production fluid, thereby improving its flow by decreasing its density.

14. A method for using amylase in brewing or alcohol production comprising the following steps:

- 25 (a) providing a polypeptide comprising a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;
- (b) providing a composition used for brewing or in alcohol production comprising a starch; and

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(c) combining the polypeptide of step (a) with the composition of the step (b) under conditions wherein the polypeptide can hydrolyze the starch in the composition used for brewing or alcohol production.

5 15. An oil well drilling fluid comprising a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1.

16. A method for changing the viscosity of a composition comprising treating the composition with a polypeptide of claim 4.

10 17. A method for aiding in the carrying away of drilling mud comprising treating the drilling mud with a composition comprising a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1.

18. A method for bio-bleaching a composition comprising treating the composition with a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1.

19. A method for making an ethanol-based fuel comprising the following steps:

(a) providing an alpha-amylase enzyme of claim 4;

15 (b) providing a composition comprising a starch; and

(c) contacting the amylase of (a) with the composition of (b) under conditions wherein the amylase hydrolyzes the starch.

20. A method for producing a food or feed comprising a recombinant alpha-amylase, the method comprising the steps of:

20 (a) providing a polypeptide having an alpha-amylase activity, wherein the polypeptide comprises a polypeptide of claim 4, or a polypeptide encoded by a nucleic acid of claim 1;

(b) providing a composition comprising a food or feed;

(c) expressing the nucleic acid to produce a recombinant alpha-amylase; and

25 (d) mixing the recombinant alpha-amylase and the feed-comprising composition, thereby producing a food or feed comprising a recombinant amylase.

21. A baking process:

(a) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1; or

(b) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1, wherein the baking process further comprises use of a second polypeptide having an amylase activity, an alpha amylase activity, or a beta amylase, or another enzyme.

22. A corn wet milling process:

(a) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1; or

(b) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1, wherein the corn wet milling process further comprises use of a second polypeptide having amylase activity, an alpha amylase activity, or a beta amylase, or another enzyme.

23. A dry milling process:

(a) comprising use of a polypeptide of claim 4 or a polypeptide encoded by the nucleic acid of claim 1; or

(b) comprising use of a polypeptide of claim 4 or a polypeptide encoded by the nucleic acid of claim 1, wherein the process further comprises use of a second polypeptide having amylase activity, an alpha amylase activity, or a beta amylase, or another enzyme.

24. A drilling process:

(a) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1; or

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(b) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1, wherein the process further comprises use of a second polypeptide having amylase activity, an alpha amylase activity, or a beta amylase, or another enzyme.

5 25. A method for making an alcohol:

(a) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1; or

10 (b) comprising use of a polypeptide of claim 4 or a polypeptide encoded by a nucleic acid of claim 1, wherein the process further comprises use of a second polypeptide having amylase activity, an alpha amylase activity, or a beta amylase, or another enzyme.

26. An isolated, synthetic, or recombinant nucleic acid according to claim 1, substantially as hereinbefore described.

15 27. An isolated, synthetic, or recombinant polypeptide having alpha-amylase activity according to claim 4, substantially as hereinbefore described.

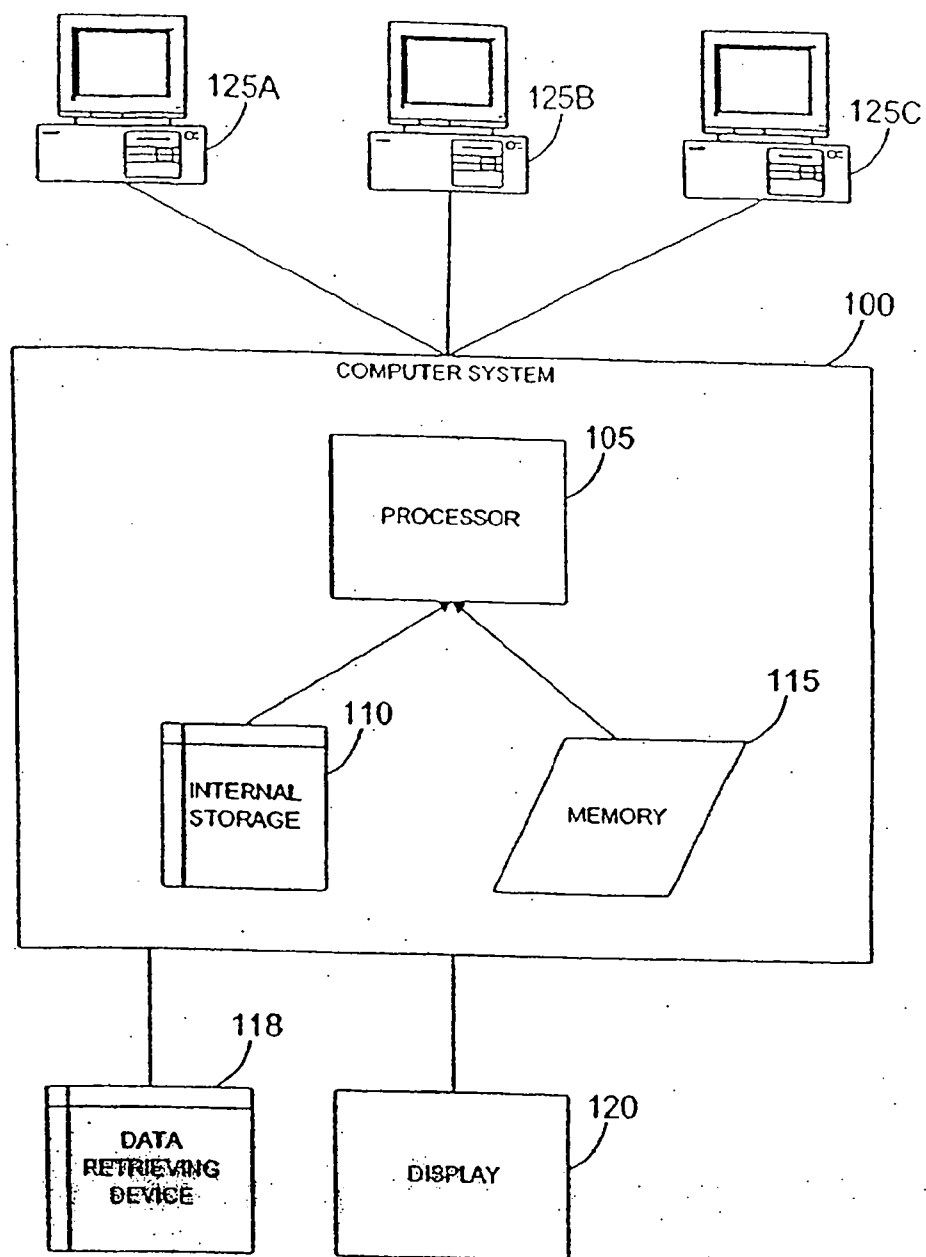


FIGURE 1

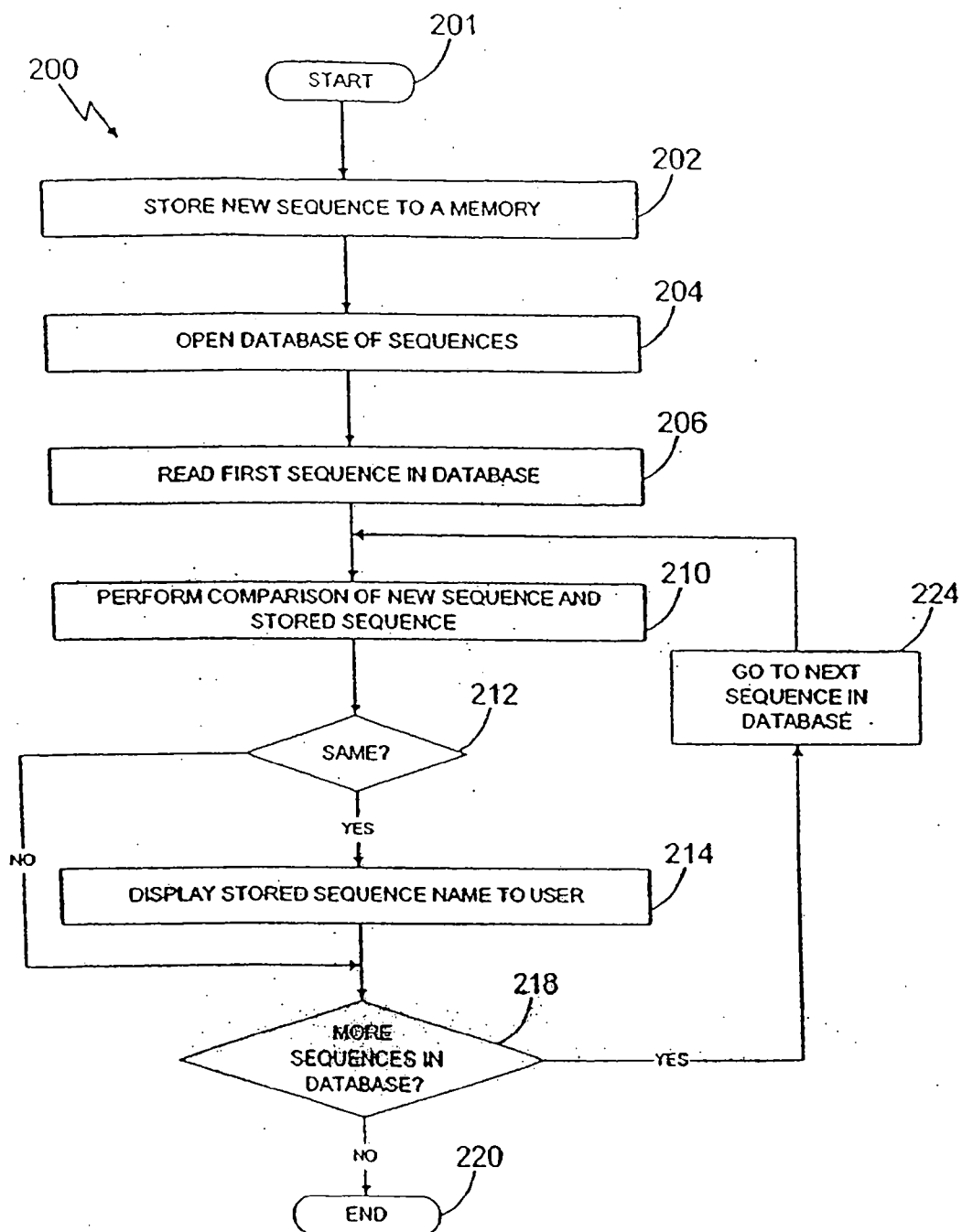


FIGURE 2

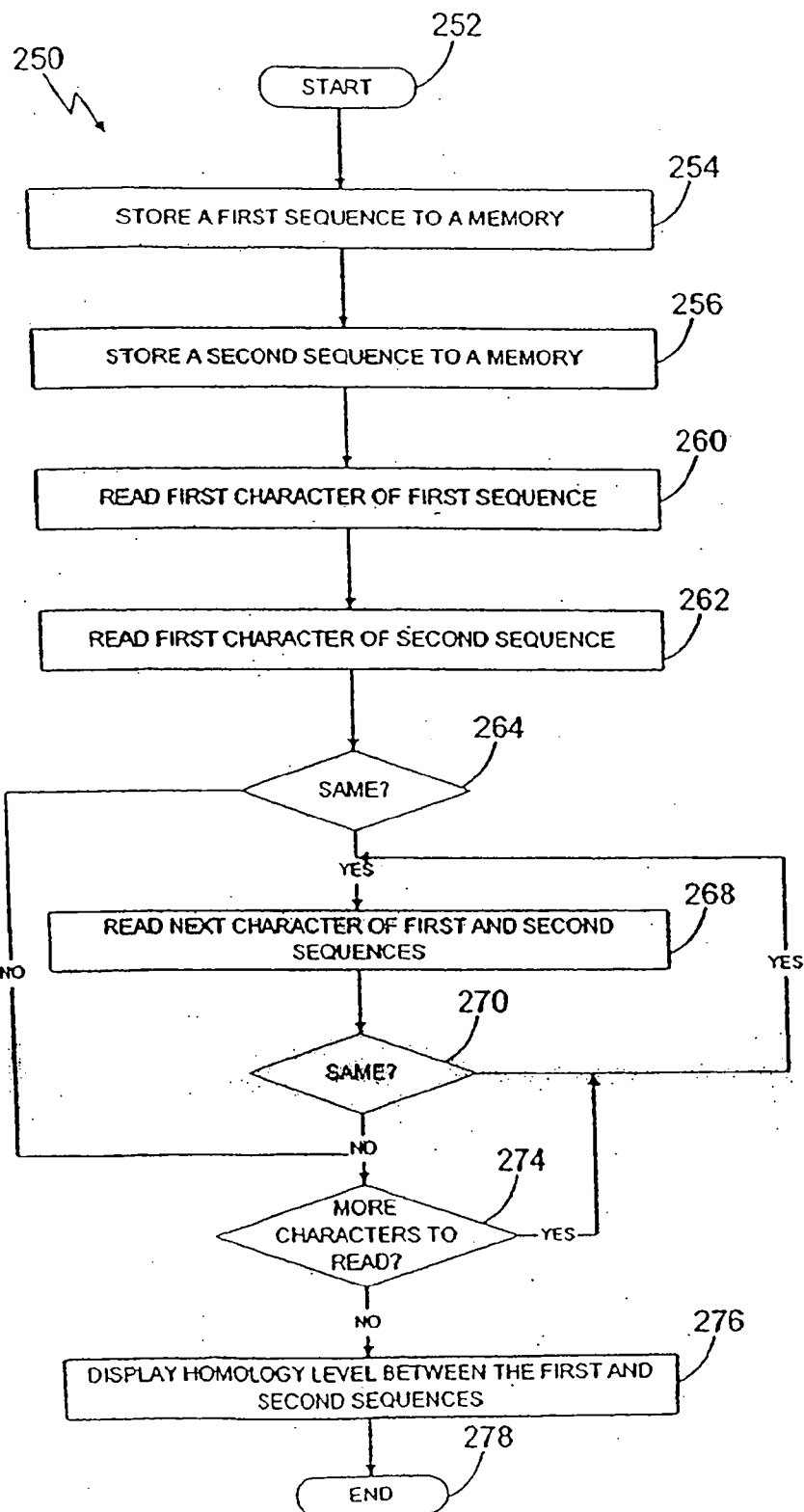


FIGURE 3

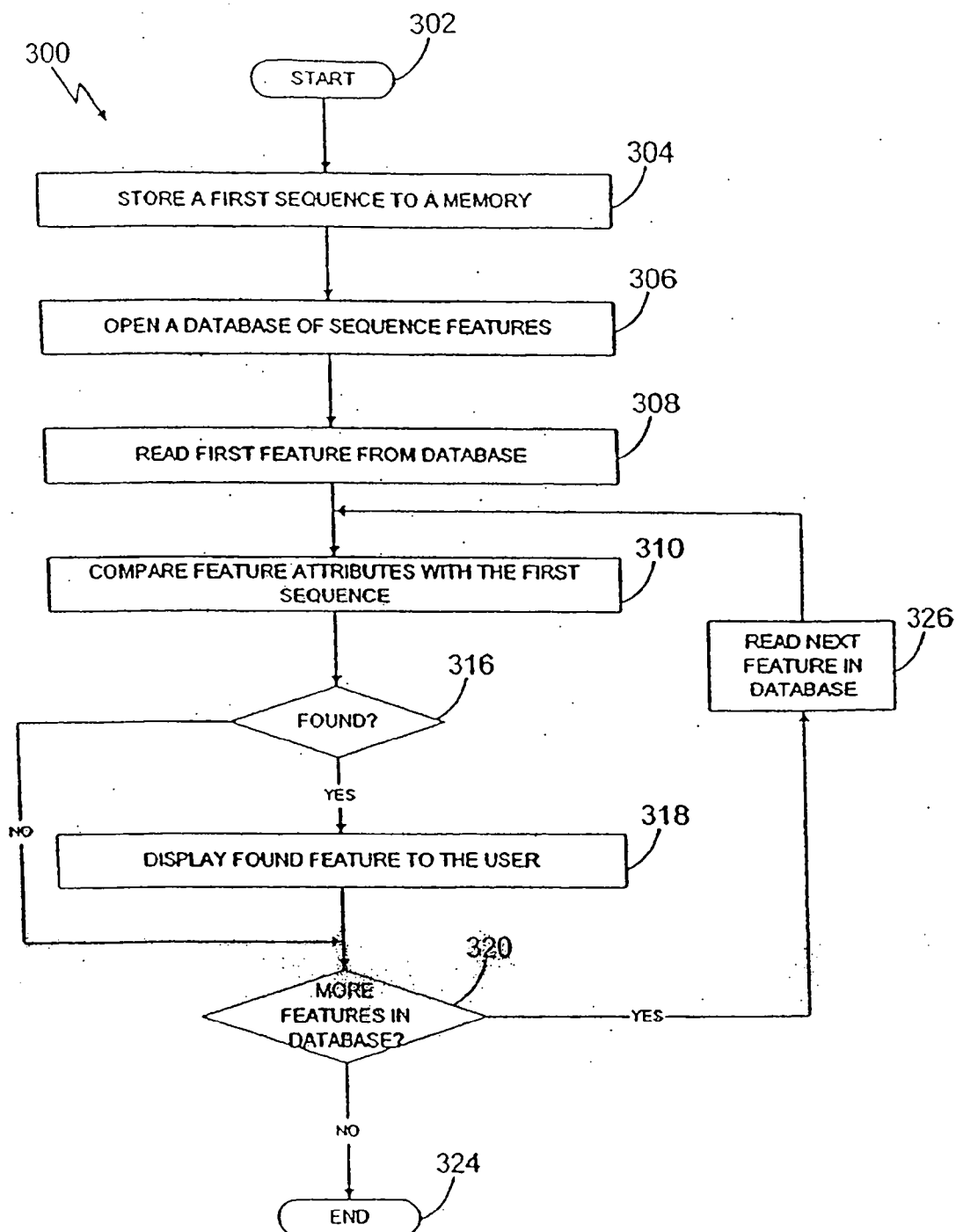


FIGURE 4

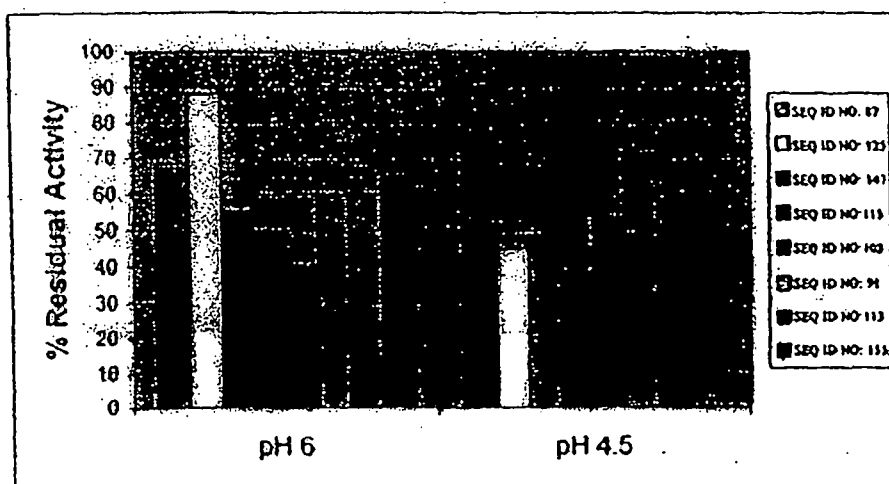


FIGURE 5

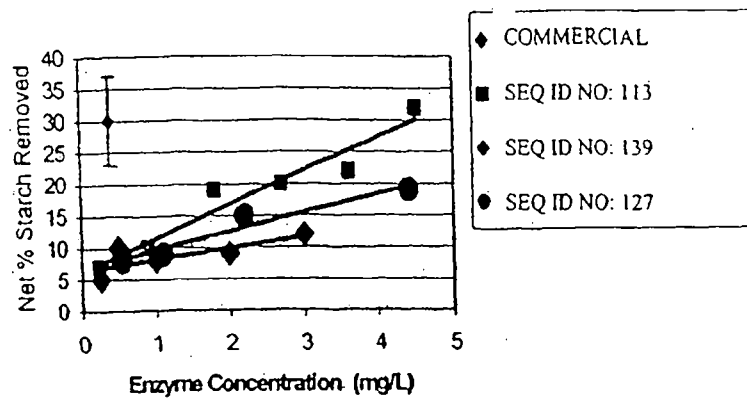


FIGURE 6

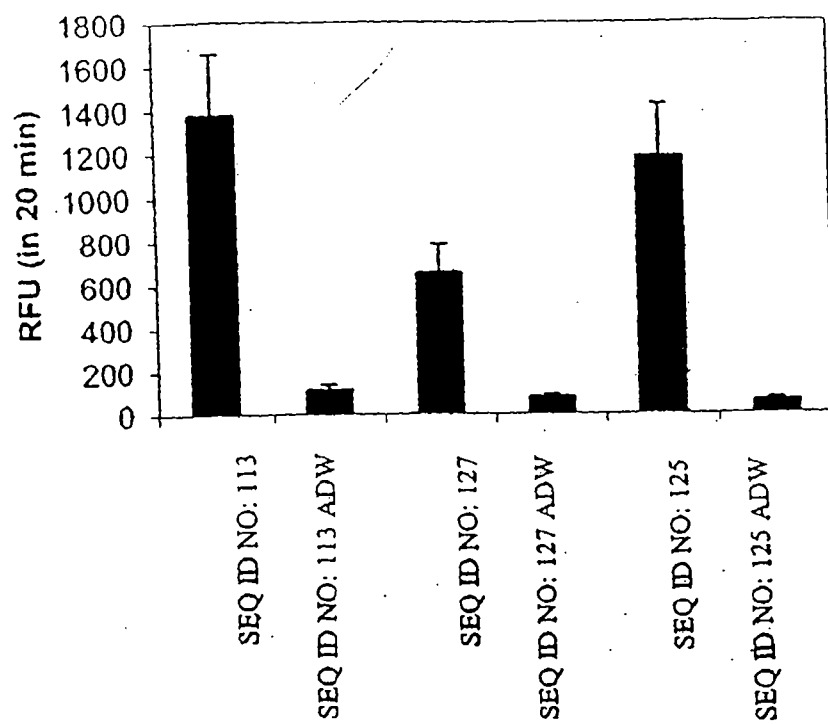


FIGURE 7

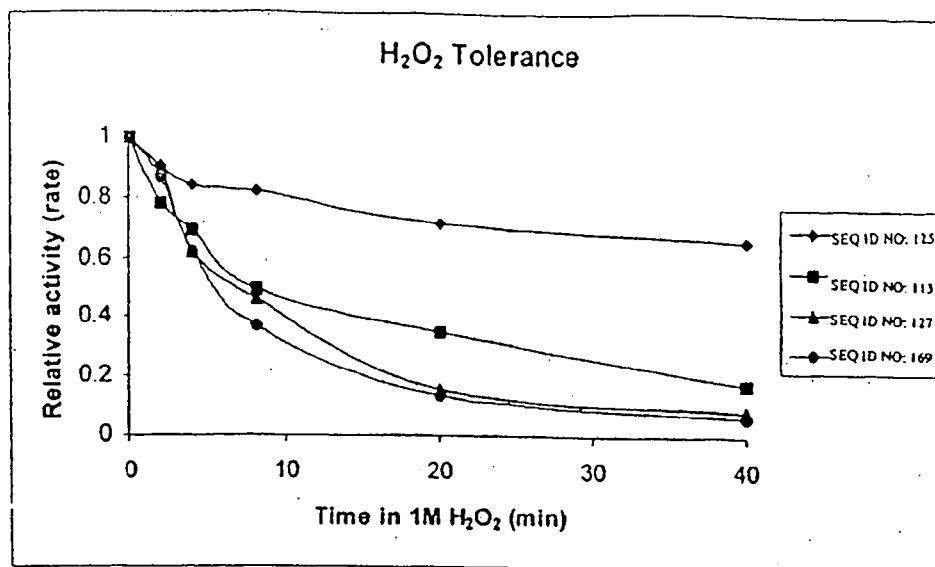


FIGURE 8

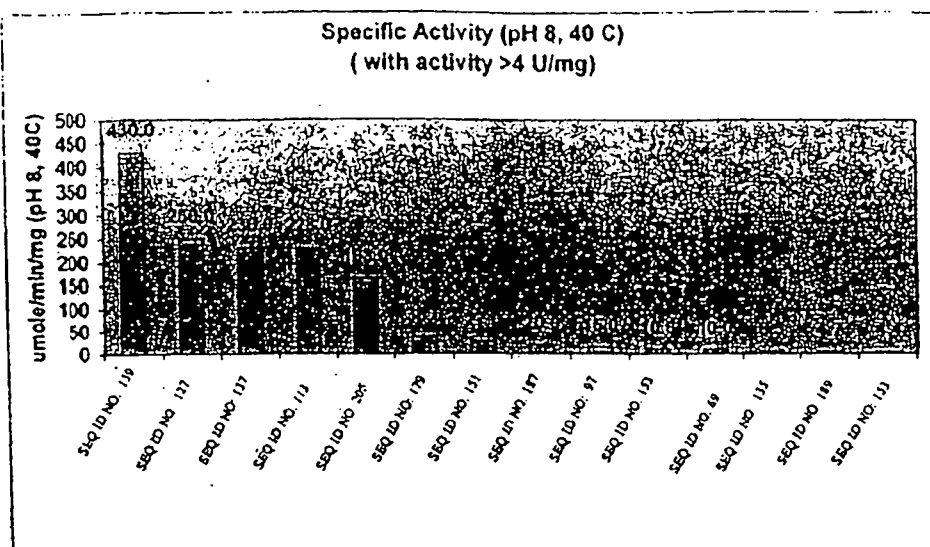


FIGURE 9A

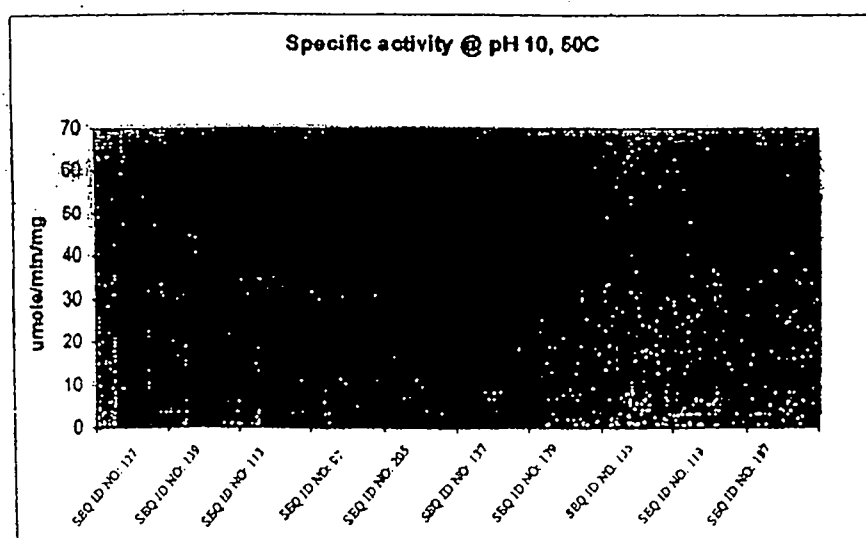


FIGURE 9B

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      1
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SEQ ID NO.:127 (1) -QANTAPVNGTMMQYFEWDLNDGTLWTHVKNEASSLSLQITALWIPPHYKST$-QADVGYGVYLYDAGEINOKSTIR
SEQ ID NO.:115 (1) AKYSELEQGGVINDQARYWDPREGQIWNQDTEKIKIPWYDAQISQWEDPSKCMGGAYSMGSDPYDYFDLGEFYKGTVE

      81
SEQ ID NO.:113 (76) TKYGTGSELOSAINBLHSRQIMVYGEVVENHKGADPTEDVTAVEVDPADRNEVISGDIKIKANTHFHFPGRGSTYSDFK
SEQ ID NO.:127 (79) TKYGTGTQYQAIQAASAGHQVYVAVVFNKQABSTEWDAVEVNPSHRNQSTSTYQIQANTKDFPGRGNTYSSEK
SEQ ID NO.:115 (81) TRESABEELNMISTAHQYGIKVISQDQENQVSSLEWNPYVGVYTRQDPSHVASQYKAHYMDHFN-----

      161
SEQ ID NO.:113 (156) WHWYHFDGTWDESEKLNRYKFGG--KASDDEVSNENGNYOYLMYADSDYDHPVAACIKRWGTWANELOEGFHLA
SEQ ID NO.:127 (159) WRWYHFDGTWDESEKLNRYKFGGTGKADDEVDAENGHYOYLMBADLCMDHPEVVTELKNWGTWYNTNVDGFLA
SEQ ID NO.:115 (150) -----N$TSEGTFGGFPDDELVPPNQYLLASHES-----YAAVRSIGTEAHFY

      241
SEQ ID NO.:113 (234) WHIKYSPLEMDQNHVREKTKGEFTHRYNDGAGESTYKTNPNHSESTVSEHYDFHASTOGGGYDMKLLNG--
SEQ ID NO.:127 (239) WHIKYSFFPDNLTHVRSQTRKGLFAAGQISYDVNKLHIYITKSGMSHMAADHNNFYTSKSSGYDMRYLLN--
SEQ ID NO.:115 (200) WGGYGAWVVDQLQMGG-----HAGGTDNVDAILLAYSIG--AKVSEFHYKMOEAPDNKNIPAHVYAYGCE

      321
SEQ ID NO.:113 (312) WKKKHPRKRIFFDMLHQPQGSLESTVQTFFPLKRWATRESYHOVEAGOMYGTGDSQ--REIPAKKHIEPIL
SEQ ID NO.:127 (317) WKKDQSLKRIFFDMLHQPQGSLSQSWVIFFFPLKRWATROEYHCVNCGHYGIPKYN-----IPKSKRIPLL
SEQ ID NO.:115 (272) WYVRDFFRAKFFIADH-----IINNYPTKRYTYZ-GOHVHNRAYEWLKND-----KLNNA--

      401
SEQ ID NO.:113 (390) KARKQYVSAQHDYFDHRIHGWTHSDSVANSGLAAITDGPGGAKRMVYQONAGBTNDITNRS--EPVVINS
SEQ ID NO.:127 (392) IARBYAYGQDYIDHQIIGWTHSDSPNSGLAAITDGPGGKWMYVKKHAGKVQDITNRS--ETVTINAD
SEQ ID NO.:115 (331) WIHEHLGCHTILYDDDEIEMKZYGDSPGL-IYENLGSDWEEHWNVSKPAGYTIHETYNLGGWVDRYQYD

      481
SEQ ID NO.:113 (468) GGEFHVN-----GGEESIIVOR-----
SEQ ID NO.:127 (470) GGEFKVN-----GGEESIIVKTSQVFTVNNATTISGQNVYVVGNIPELGNWNTANAIKMTPSSYPTWKATIALP
SEQ ID NO.:115 (410) VVKLTAPPHPDANGYYVYVWSAGVG-----

      561
SEQ ID NO.:113 (485) -----
SEQ ID NO.:127 (541) QGKAIEPKFIKQDQSGNVVWESI PNRTYTVPLSTGSYTASWNP
SEQ ID NO.:115 (437) -----

```

FIGURE 10

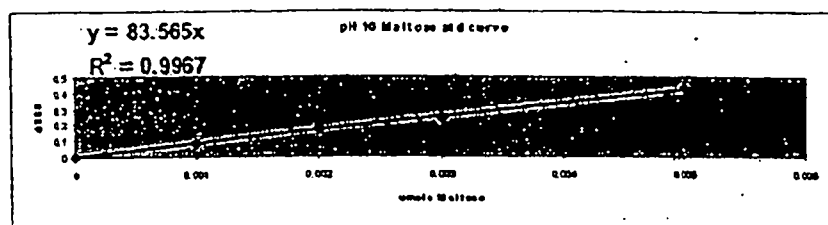


FIGURE 11

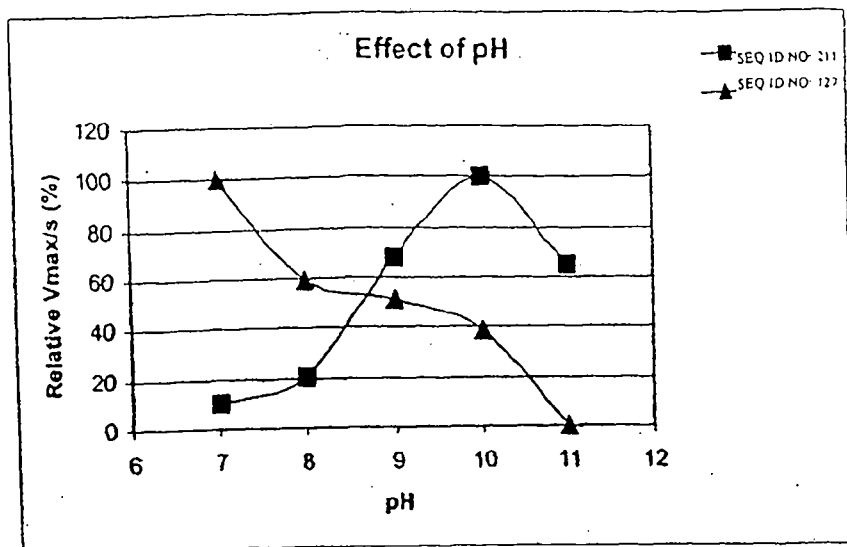


FIGURE 12

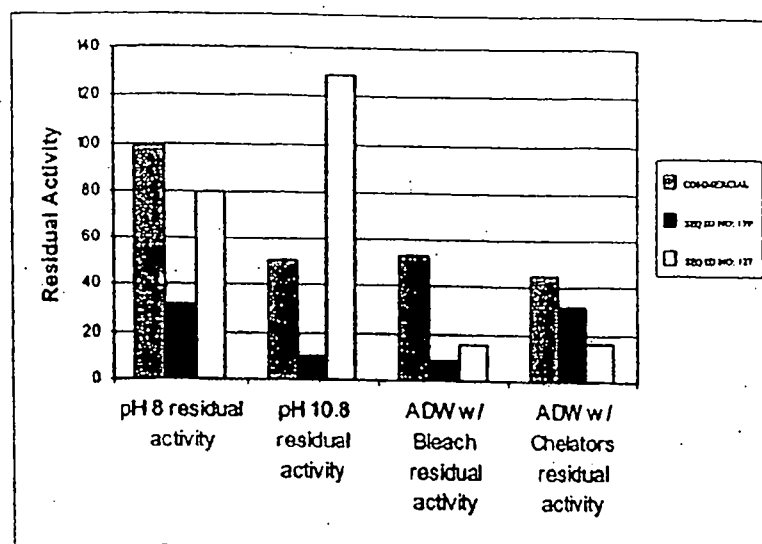


FIGURE 13

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1                               50
SEQ ID NO: 81 -----MKK FVALFITMFF VVSMVV...AQPASAA K
pyro -----MKK FVALLITMFF VVSMVAV...AQPASAA K
pyro2 -----VNIKK LTPLLTLLLF FI...VL...ASPVSAA K
thermo SESQC TATCT WRVVYMSAKK LLALLFV LAV LVGVAVIP AR VGIAPVSAG A
thermo2 -----MA RKVLVALL VF LVVLSVSAV P
Consensus -----SA--

51                               100
SEQ ID NO: 81 YS..ELEEGG VIMQAFYWDV PGGGIWWD TI RSKIP EWYEA GISAIWIPPA
pyro YS..ELEEGG VIMQAFYWDV PGGGIWWD TI RSKIP EWYEA GISAIWIPPA
pyro2 YL..ELEEGG VIMQAFYWDV PGGGIWWD HI RSKIP EWYEA GISAIWLPP P
thermo TSRPSLEEGG VIMQAFYWDV PGGGIWWD TI RSKIPDWA SA GISAIWIPPA
thermo2 AKAETLEGG VIMQAFYWDV PGGGIWWD TI AQKIPDWA SA GISAIWIPPA
Consensus -----LE-GG VIMQAFYWDV P-GGIWWD-I --KIP-W-A GISAIW-PP-
Sense primer

101                               150
SEQ ID NO: 81 SKGMSGGYSM GYDPYDFFDL GEYNQKGTIE TRFGSKQELI NMINTAHAYG
pyro SKGMSGAYS M GYDPYDFFDL GEYNQKGTVE TRFGSKQSLI NMINTAHAYG
pyro2 SKGMSGGYSM GYDPYDFFDL GEYYQKGTVE TRFGSKKEE LV RLIQTAHAYG
thermo SKGMSGAYS M GYDPYDFFDL GEYYQKGTVE TRFGSKQELI NMINTAHAYG
thermo2 SKGMSGGYSM GYDPYDFFDL GEYYQKGSVE TRFGSKKEE LV NMINTAHAYN
Consensus SKGM-G-YSM GYDPYD-FDL GEY-QKG--E TRFGSK-EL- --I-TAH--

151                               200
SEQ ID NO: 81 IKVIADIVIN HRAGGDLEWN PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
pyro IKVIADIVIN HRAGGDLEWN PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
pyro2 IKVIADIVIN HRAGGDLEWN PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
thermo IKVIADIVIN HRAGGDLEWN PFTNSYTWTD FSKVASGKYT ANYLDFHPNE
thermo2 MKVIADIVIN HRAGGDLEWN PFTNSYTWTD FSKVASGKYT ANYLDFHPNE
Consensus -KVIAD-VIN HRAGGDLEWN PF---YT WTD FSKVASGKYT ANYLDFHPNE

201                               250
SEQ ID NO: 81 VKCCDEGTFG GFDPIAHEKS WDQHWLW ASD ESYAAYLR SI GVDWRFDYV
pyro VKCCDEGTFG GFDPIAHEKE WDQHWLW ASD ESYAAYLR SI GVDWRFDYV
pyro2 LHCCDEGTFG GFDPICHKE WDQYWLW KSN ESYAAYLR SI GFDGWRFDYV
thermo VKCCDEGTFG GFDPIAHEKS WDQYWLW ASQ KSYAAYLR SI GVDWRFDYV
thermo2 LHAGD SGTFG GYPDICHDKS WDQHWLW ASN ESYAAYLR SI GIDWRFDYV
Consensus -----D-GTFG G-PDI-H-K- WDQ-WLW-S- -SYAAYLR SI G-D-WRFDYV

251                               300
SEQ ID NO: 81 KGYGAWVVKD WLNWNGWAV GEYWDTNVDA LLNWAYSSGA KVFDFPLYK
pyro KGYGAWVVKD WLNWNGWAV GEYWDTNVDA LLNWAYSSGA KVFDFPLYK
pyro2 KGYGAWVVKD WLNWNGWAV GEYWDTNVDA LLSWAYESGA KVFDFPLYK
thermo KGYGAWVVKD WLNWNGWAV GEYWDTNVDA LLNWAYSSGA KVFDFPLYK
thermo2 KGYAPWVVKD WLNWNGWAV GEYWDTNVDA LLSWAYDSGA KVFDFPLYK
Consensus KGY--WV-- WL--W--AV GEYWDTNVDA LL-WAY-SGA KVFDFPLYK

301                               350
SEQ ID NO: 81 MDEAFDNKNI PALVSA LQNG QTVVSRD PFK AVTFVANHDT DIIWNKYLAY
pyro MDEAFDNTNI PALVDA LQNG GTVVSRD PFK AVTFVANHDT DIIWNKYPAY
pyro2 MDEAFDNKNI PALVYA LQNG QTVVSRD PFK AVTFVANHDT DIIWNKYPAY
thermo MDEAFDNKNI PALVSA LQNG QTVVSRD PFK AVTFVANHDT DIIWNKYPAY
thermo2 MDEAFDNKNI PALVDA LQNG GTVVSRD PFK AVTFVANHDT DIIWNKYPAY
Consensus MDEAFDN-NI PALV-AL-NG -TVVSRD PFK AVTFVANHDT -IIWNKY-A Y

```

FIGURE 14A

```

351                                     400
SEQ ID NO: 81 AFILTYEGQP VIFYRDYEEW LNKDRLNNLI WIHDHLAGGS TSIVYYDSDE
      pyro AFILTYEGQP VIFYRDYEEW LNKDKLNNLI WIHDHLAGGS TSIVYYDSDE
      pyro2 AFILTYEGQP VIFYRDFEEW LNKDKLINLI WIHDHLAGGS TTIVYYDNDS
      thermo AFILTYEGQP VIFYRDYEEW LNKDRLKNLI WIHNNLAGGS TSIVYYDNDS
      thermo2 AFILTYEGQP AIFYRDYEEW LNKDRLRNLI WIHDHLAGGS TDIIYYDSDE
Consensus AFILTYEGQP -IFYRD-EEW LNKD-L-NLI WIH--LAGGS T-I-YYD-DE

401                                     450
SEQ ID NO: 81 MIFVRNGYGS KPGLITYINL GSSKVGRWVY VPKFAGACIH EYTGNLGGWV
      pyro LIFVRNGDSK RPGLITYINL GSSKVGRWVY VPKFAGACIH EYTGNLGGWV
      pyro2 LIFVRNGDSR RPGLITYINL SPNWWGRWVY VPKFAGACIH EYTGNLGGWV
      thermo LIFVRNGYGN KPGLITYINL GSSKVGRWVY VPKFAGSCIH EYTGNLGGWV
      thermo2 LIFVRNGYGD KPGLITYINL GSSKAGR WVY VPKFAGSCIH EYTGNLGGW I
Consensus -IFVRNG--- -PGLITYINL -----GRWVY VPKFAG-CIH EYTGNLGGW-

451                                     486
SEQ ID NO: 81 DKYVYSSGWV YFEAPAYDPA NGQYGYSVWS YCGVG*
      pyro DKYVSSSGWV YLEAPAYDPA SGQYGYTVWS YCGVG*
      pyro2 DKRVDSGWV YLEAPPDPA NGYGYYSVWS YCGVG*
      thermo DKYVGSNGWV YLEAPAHDP A KGQYGYSVWS YCGVG*
      thermo2 DKWVDSGRV YLEAPAHDP A NGQYGYSVWS YCGVG*
Consensus DK-V-S-G-V Y-EAP--DPA -G-YGY-VWS YCGVG*
Antisense primer

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FIGURE 14A
(cont.)

	1		50
SEQ ID NO: 81	-----	-----MKK FVA	LFITMFFVVS MAVVAQPASA
pyro	-----	-----MKK FVA	LLITMFFVVS MAAVAQPASA
SEQ ID NO: 73	-----	-----	-----
thermo2	-----	-----MA	RKVLVALLVF LVLVSVAVP
SEQ ID NO: 75	-----	-----	-----
SEQ ID NO: 77	-----	-----	-----
SEQ ID NO: 83	-----	-----	-----
SEQ ID NO: 85	-----	-----	-----
SEQ ID NO: 79	-----	-----MKP AKL	LVFVLVVS IL AGLYAQPAGA
thermo	SESQCTATCT	WRVVYMSAKK	LLALLFVLAV LVGVAVIPAR VGIAPVSAGA
pyro2	-----	-----VNIKK	LTPLLTLLLF FIVLASPVSA
CLONE A	-----	-----MRRSARV	LVLIIAFFLL AGIYYPSTSA
Consensus	-----	-----	-----
	51		100
SEQ ID NO: 81	AKYSELEEGG	VIMQAFYWDV	PGGGIWWDTI RSKIPWYEA GISAIWIPPA
pyro	AKYSELEEGG	VIMQAFYWDV	PAGGIWWDTI RSKIPWYEA GISAIWIPPA
SEQ ID NO: 73	---MALEEGG	LIMQAFYWDV	PGGGIWWDTI AQKIPDASA GISAIWIPPA
thermo2	AKAETLEGG	VIMQAFYWDV	PGGGIWWDTI AQKIPDASA GISAIWIPPA
SEQ ID NO: 75	---MALEEGG	LIMQAFYWDV	PMGGIWWDTI AQKIPDASA GISAIWIPPA
SEQ ID NO: 77	---MALEEGG	LIMQAFYWDV	PMGGIWWDTI AQKIPDASA GISAIWIPPA
SEQ ID NO: 83	---MALEEGG	LIMQAFYWDV	PGGGIWWDTI AQKIPWASA GISAIWIPPA
SEQ ID NO: 85	---MALEEGG	LIMQAFYWDV	PGGGIWWDTI AQKIPWASA GISAIWIPPA
SEQ ID NO: 79	AKYLELEEGG	VIMQAFYWDV	PSGGIWWDTI RQKIPWYDA GISAIWIPPA
thermo	TSRPSLEEGG	VIMQAFYWDV	PAGGIWWDTI RSKIPDASA GISAIWIPPA
pyro2	AKYLELEEGG	VIMQAFYWDV	PGGGIWWDHI RSKIPWYEA GISAIWLPPP
CLONE A	AKYSELEGG	VIMQAFYWDV	PEGGIWWDTI RQKIPWYDA GISAIWIPPA
Consensus	-----GG	-IMQAFYWDV	P-GGIWWD-I --KIP-W--A GISAIW-PP-
	101		150
SEQ ID NO: 81	SKGMSGGYSM	GYDPYDFFDL	GEYNQKGTIE TRFGSKQELI NMINTAHAYG
pyro	SKGMSGAYSM	GYDPYDFFDL	GEYNQKGTVE TRFGSKQELI NMINTAHAYG
SEQ ID NO: 73	SKGMSGGYSM	GYDPYDFFDL	GEYYQKGSVE TRFGSKEELV NMINTAHAYN
thermo2	SKGMSGGYSM	GYDPYDFFDL	GEYYQKGSVE TRFGSKEELV NMINTAHAYN
SEQ ID NO: 75	SKGMSGGYSM	GYDPYDYFDL	GEYYQKGTVE TRFGSKQELI NMINTAHAYG
SEQ ID NO: 77	SKGMSGGYSM	GYDPYDYFDL	GEYYQKGTVE TRFGSKQELI NMINTAHAYG
SEQ ID NO: 83	SKGMSGGYSM	GYDPYDFFDL	GEYYQKGTVE TRFGSKEELV NMINTAHAYG
SEQ ID NO: 85	SKGMSGGYSM	GYDPYDFFDL	GEYYQKGTVE TRFGSKEELV NMINTAHAYG
SEQ ID NO: 79	SKGMSGAYSM	GYDPYDFFDL	GEYDQKGTVE TRFGSKQELI NMINTAHAYG
thermo	SKGMSGGYSM	GYDPYDYFDL	GEYYQKGTVE TRFGSKEELV NMINTAHAYG
pyro2	SKGMSGAYSM	GYDPYDYFDL	GEYYQKGTVE TRFGSKEELV RLIQTAHAYG
CLONE A	SKGMSGAYSM	GYDPYDYFDL	GEFYQKGTVE TRFGSKEELV NMINTAHAYG
Consensus	SKGM-G-YSM	GYDPYD-FDL	GE--QKG--E TRFGSK-EL- --I-TAH---
	151		200
SEQ ID NO: 81	IKVIADIVIN	HRAGGDLEWN	PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
pyro	IKVIADIVIN	HRAGGDLEWN	PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 73	MKVIADIVIN	HRAGGDLEWN	PFTNSYTWTD FSKVASGKYT ANYLDFHPNE
thermo2	MKVIADIVIN	HRAGGDLEWN	PFTNSYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 75	MKVIADIVIN	HRAGGDLEWN	PFVNDYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 77	MKVIADIVIN	HRAGGDLEWN	PFVNDYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 83	IKVIADIVIN	HRAGGDLEWN	PFVNDYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 85	IKVIADIVIN	HRAGGDLEWN	PFVNDYTWTD FSKVASGKYT ANYLDFHPNE
SEQ ID NO: 79	IKVIADIVIN	HRAGGDLEWN	PFVNDYTWTD FSKVASGKYT ANYLDFHPNE
thermo	IKVIADIVIN	HRAGGDLEWN	PFTNSYTWTD FSKVASGKYT ANYLDFHPNE
pyro2	IKVIADIVIN	HRAGGDLEWN	PFVGDYTWTD FSKVASGKYT ANYLDFHPNE
CLONE A	IKVIADIVIN	HRAGGDLEWN	PYVGDYTWTD FSKVASGKYK AHYMDHPNN
Consensus	-KVIAD-VIN	HRAGG-LEWN	P----YTWTD FSKVASGKY- A-Y-DFHPN-

FIGURE 14B

	201	250
SEQ ID NO: 81	VKCCDEGTFG GFDIAHEKS WDQHWLWASD ESYAAYLR SI GVDARFDY V	
pyro	VKCCDEGTFG GFDIAHEKE WDQHWLWASD ESYAAYLR SI GVDARFDY V	
SEQ ID NO: 73	LHAGDSGTFG GYPDICHDKS WDQHWLWASN ESYAAYLR SI GIDARFDY V	
thermo2	LHAGDSGTFG GYPDICHDKS WDQHWLWASN ESYAAYLR SI GIDARFDY V	
SEQ ID NO: 75	LHAGDSGTFG GYPDICHDKS WDQYWLWASQ ESYAAYLR SI GIDARFDY V	
SEQ ID NO: 77	LHAGDSGTFG GYPDICHDKS WDQYWLWASQ ESYAAYLR SI GIDARFDY V	
SEQ ID NO: 83	LHCCDEGTFG GYPDICHDKS WDQYWLWASS ESYAAYLR SI GVDARFDY V	
SEQ ID NO: 85	LHCCDEGTFG GYPDICHDKS WDQYWLWASS ESYAAYLR SI GVDARCFDY V	
SEQ ID NO: 79	VKCCDEGTFG GFDIAHEKS WDQYWLWASN ESYAAYLR SI GVDARFDY V	
thermo	VKCCDEGTFG GFDIAHEKS WDQYWLWASQ KSYAAYLR SI GIDARFDY V	
pyro2	LHCCDEGTFG GFPDICHHEKE WDQYWLWASN ESYAAYLR SI GFDGWRFDY V	
CLONE A	YSTSDEGTFG GFPDIDHLPV FNQYWLWASN ESYAAYLR SI GIDARFDY V	
Consensus	----D-GTFG G-PDI-H--- --Q-WLW-S- -SYAAYLR SI G-D-W-FDY V	
	251	300
SEQ ID NO: 81	KGYGAWVVVD WLNWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
pyro	KGYGAWVVVD WLNWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
SEQ ID NO: 73	KGYAPWVVKN WLNWNGGWAV GEYWDTNVDA LLSWAYDSGA KVFDFFPLY K	
thermo2	KGYAPWVVKN WLNWNGGWAV GEYWDTNVDA LLSWAYDSGA KVFDFFPLY K	
SEQ ID NO: 75	KGYAPWVVVD WLNWNGGWAV GEYWDTNVDA VLNWAYSSGA KVFDFFPLY K	
SEQ ID NO: 77	KGYAPWVVVD WLNWNGGWAV GEYWDTNVDA VLNWAYSSGA KVFDFFPLY K	
SEQ ID NO: 83	KGYGAWVVVD WLSWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
SEQ ID NO: 85	KGYGAWVVVD WLSWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
SEQ ID NO: 79	KGYGAWVVVD WLDWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
thermo	KGYGAWVVVD WLNWNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
pyro2	KGYGAWVVVD WLNWNGGWAV GEYWDTNVDA LLSWAYESGA KVFDFFPLY K	
CLONE A	KGYGAWVVVD WLSQNGGWAV GEYWDTNVDA LLNWAYSSGA KVFDFFPLY K	
Consensus	KGY--WVV-- WL--W---AV GEYWDTNVDA -L-WAY-S-A KVFDFF-LYY K	
	301	350
SEQ ID NO: 81	MDEAFDNKNI PALVSA LQNG QTVVSRD PFK AVTFVANHDT DIIWNKYLA Y	
pyro	MDEAFDNTNI PALVDA LQNG GTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
SEQ ID NO: 73	MDEAFDNNNI PALVDA LQNG GTVVS RD PFK AVTFVANHDT NIIWNKYPA Y	
thermo2	MDEAFDNNNI PALVDA LQNG GTVVS RD PFK AVTFVANHDT NIIWNKYPA Y	
SEQ ID NO: 75	MDEAFDNNNI PALVDA LRYG QTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
SEQ ID NO: 77	MDEAFDNNNI PALVDA LRYG QTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
SEQ ID NO: 83	MDEAFDNTNI PALVDA LRYG QTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
SEQ ID NO: 85	MDEAFDNTNI PALVYA LQNG GTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
SEQ ID NO: 79	MDEAFDNKNI PALVEA LQNG GTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
thermo	MDEAFDNKNI PALVSA LQNG QTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
pyro2	MDEAFDNNNI PALVYA LQNG QTVVS RD PFK AVTFVANHDT DIIWNKYPA Y	
CLONE A	MDEAFDNKNI PALVYA LQNG GTVVS RD PFK AVTFVANHDT NIIWNKYPA Y	
Consensus	MD-AF DN-NI PALV-A---G -TVVSRD PFK AVTFVANHDT -IIWNKY-A Y	
	351	400
SEQ ID NO: 81	AFILTYEGQP VIFYRDYEEW LNKDRLNLI WINDHLAGGS TSIVYYDSDE	
pyro	AFILTYEGQP VIFYRDYEEW LNKDKLNLI WINDHLAGGS TSIVYYDSDE	
SEQ ID NO: 73	AFILTYEGQP AIFYRDYEEW LNKDRLNLI WINDHLAGGS TDIIYYDSDE	
thermo2	AFILTYEGQP AIFYRDYEEW LNKDRLNLI WINDHLAGGS TDIIYYDSDE	
SEQ ID NO: 75	AFILTYEGQP TIFYRDYEEW LNKDKLNLI WINDHLAGGS TDIVYYDSDE	
SEQ ID NO: 77	AFILTYEGQP TIFYRDYEEW LNKDKLNLI WINDHLAGGS TDIVYYDSDE	
SEQ ID NO: 83	AFILTYEGQP VIFYRDYEEW LNKDKLNLI WINDHLAGGS TDIVYYDSDE	
SEQ ID NO: 85	AFILTYEGQP VIFYRDYEEW LNKDKLNLI WINDHLAGGS TDIVYYDSDE	
SEQ ID NO: 79	AFILTYEGQP TIFYRDYEEW LNKDRLNLI WINDHLAGGS TDIVYYDSDE	
thermo	AFILTYEGQP VIFYRDYEEW LNKDRLNLI WINDHLAGGS TSIVYYDSDE	
pyro2	AFILTYEGQP VIFYRDYEEW LNKDKLNLI WINDHLAGGS TTIVYYDSDE	
CLONE A	AFILTYEGQP VIFYRDYEEW LNKDKLNLI WINDHLAGGS TKILYYDDDE	
Consensus	AFILTYEGQP -IFYRD-EEW LNKD-L-NLI WIH--LAGGS T-I-YD-DE	

FIGURE 14B

(cont.)

	401		450
SEQ ID NO: 81	MIFVRNGYGS	KPGLITYINL	GSSKVGRWVY V.PKFAGACI HEYTGNLGGW
pyro	LIFVRNGDSK	RPGLITYINL	GSSKVGRWVY V.PKFAGACI HEYTGNLGGW
SEQ ID NO: 73	LIFVRNGYGD	KPGLITYINL	GSSKAGR WVY V.PKFAGSCI HEYTGNLGGW
thermo2	LIFVRNGYGD	KPGLITYINL	GSSKAGR WVY V.PKFAGSCI HEYTGNLGGW
SEQ ID NO: 75	LIFVRNGYGS	KPGLITYINL	GSSKAGR WVY V.PKFAGSCI HEYTGNLGGW
SEQ ID NO: 77	LIFVRNGYGS	KPGLITYINL	ASSKAGR WVY V.PKFAGSCI HEYTGNLGGW
SEQ ID NO: 83	LIFVRNGYGT	KPGLITYINL	GSSKVGR WVY V.PKFAGSCI HEYTGNLGGW
SEQ ID NO: 85	LIFVRNGYGT	KPGLITYINL	GSSKAGR WVY V.PKFAGSCI HEYTGSLGGW
SEQ ID NO: 79	LIFVRNGYGD	KPGLITYINL	GSSKAGR WVY V.PKFAGACI HEYTGNLGGW
thermo	LIFVRNGYGN	KPGLITYINL	GSSKVGR WVY V.PKFAGSCI HEYTGNLGGW
pyro2	LIFVRNGDSR	RPGLITYINL	SPNWGR WVY V.PKFAGACI HEYTGNLGGW
CLONE A	LIFMR EGYGD	RPGLITYINL	GSDWAER WVN VGSKPAGYTI HEYTGNLGGW
Consensus	-IF-R-G---	-PGLITYINL	-----RWV- V--KFAG--I HEYTG-LGGW

	451		487
SEQ ID NO: 81	VDKYV YSSGW	VYLEAP AYDP	ANGQYGY SVW SYCGVG*
pyro	VDKYV ESSGW	VYLEAP AYDP	ASGQYGY TVW SYCGVG*
SEQ ID NO: 73	IDKWV DSSGR	VYLEAP AHDP	ANGQYGY SVW SYCGVG*
thermo2	IDKWV DSSGR	VYLEAP AHDP	ANGQYGY SVW SYCGVG*
SEQ ID NO: 75	VDKWV DSSGW	VYLEAP AHDP	ANGQYGY SVW SYCGVG*
SEQ ID NO: 77	VDKWV DSSGW	VYLEAP AHDP	ANGQYGY SVW SYCGVG*
SEQ ID NO: 83	IDKYV SSSGW	VYLEAP AHDP	ANGYYGY SVW SYCGVG*
SEQ ID NO: 85	IDKYV SSSGW	VYLEAP AHDP	ANGQYGY SVW SYCGVG*
SEQ ID NO: 79	VDKWV DSSGW	VYLEAP AHDP	ANGYYGY SVW SYCGVG*
thermo	VDKYV GSNGW	VYLEAP AHDP	AKGQYGY SVW SYCGVG*
pyro2	VDKRV DSSGW	VYLEAP PHDP	ANGYYGY SVW SYCGVG*
CLONE A	VDRYV QYDGW	VKLTAP PHDP	ANGYYGY SVW SYAGVG*
Consensus	-D--V---G-	V---AP--DP	A-G-YGY -VW SY-GVG*

FIGURE 14B
(cont.)

	1					50
SEQ ID NO: 83	-----	-----	-----	-----	-----	-----
SEQ ID NO: 85	-----	-----	-----	-----	-----	-----
SEQ ID NO: 75	-----	-----	-----	-----	-----	-----
SEQ ID NO: 77	-----	-----	-----	-----	-----	-----
SEQ ID NO: 73	-----	-----	-----	-----	-----	-----
SEQ ID NO: 79	---ATGA AGC	CTGCGAAA CT	CCTCGTCCTT T	GTGCTCGTAG	TCTCTATCCT	
SEQ ID NO: 81	---ATGA AGA	AGTTTGTC GC	CCTGTTCAT A	ACCATGTTTT	TCGTAGTGAG	
CLONE A	ATGAGGA GAT	CCGCAAGG GT	TTTGTTCT G	ATTATAGCGT	TTTCCTCTCT	
Consensus	-----	-----	-----	-----	-----	-----
	51					100
SEQ ID NO: 83	-----	-----	-----	-----	ATGGCTCTGG	
SEQ ID NO: 85	-----	-----	-----	-----	ATGGCTCTGG	
SEQ ID NO: 75	-----	-----	-----	-----	ATGGCTCTGG	
SEQ ID NO: 77	-----	-----	-----	-----	ATGGCTCTGG	
SEQ ID NO: 73	-----	-----	-----	-----	ATGGCTCTGG	
SEQ ID NO: 79	CGCGGGG CTC	TACGCCCA GC	CCGCGGGG C	GGCCAAGTAC	CTGGAGCTCG	
SEQ ID NO: 81	CATGGCA GTC	GTTGCACA GC	CAGCTAGCG C	CGCAAAGTAT	TCCGAGCTCG	
CLONE A	GGCGGGG ATT	TACTACCC CT	CCACGAGTG C	CGCAAGTAC	TCCGAGCTGG	
Consensus	-----	-----	-----	-----	-----	-----
	101					150
SEQ ID NO: 83	AAGAGGG CGG	GCTCATAA TG	CAGGCCTTC T	ACTGGGATGT	TCCTGGAGGA	
SEQ ID NO: 85	AAGAGGG CGG	GCTTATAA TG	CAGGCATTCT	ATTGGGACGT	CCCAGGTGGA	
SEQ ID NO: 75	AAGAGGG CGG	GCTTATAA TG	CAGGCATTCT	ACTGGGACGT	CCCCATGGGA	
SEQ ID NO: 77	AAGAGGG CGG	GCTCATAA TG	CAGGCCTTC T	ACTGGGACGT	CCCCATGGGA	
SEQ ID NO: 73	TAGAGGG CGG	GCTTATAA TG	CAGGCCTTC T	ACTGGGACGT	CCCAGGTGGA	
SEQ ID NO: 79	AAGAGGG CGG	CGTCATAA TG	CAGGCGTTC T	ACTGGGACGT	GCCTTCAGGA	
SEQ ID NO: 81	AAGAAGG CGG	CGTTATAA TG	CAGGCCTTC T	ACTGGGACGT	CCCAGGTGGA	
CLONE A	AGCAGGG CGG	AGTCATAA TG	CAGGCCTTC T	ACTGGGACGT	TCCGGAGGGA	
Consensus	-----GG CGG	--T-ATAA TG	CAGGC-TTC T	A-TGGGA-GT	-CC----GGA	
	151					200
SEQ ID NO: 83	GGAATCT GGT	GGGACACA AT	AGCTCAAAA G	ATACCCGAAT	GGGCAAGTGC	
SEQ ID NO: 85	GGAATCT GGT	GGGACACC AT	AGCCCAGAA G	ATACCCGAAT	GGGCAAGTGC	
SEQ ID NO: 75	GGAATCT GGT	GGGACACG AT	AGCCCAGAA G	ATACCCGACT	GGGCAAGCGC	
SEQ ID NO: 77	GGAATCT GGT	GGGACACG AT	AGCCCAGAA G	ATACCCGACT	GGGCAAGCGC	
SEQ ID NO: 73	GGAATCT GGT	GGGACACC AT	AGCCCAGAA G	ATACCCGACT	GGGCGAGCGC	
SEQ ID NO: 79	GGAATAT GGT	GGGACACA AT	ACGGCAGAA G	ATACCCGAGT	GGTACGATGC	
SEQ ID NO: 81	GGAATCT GGT	GGGACACC AT	CAGGAGCAA G	ATACCCGAGT	GGTACGAGGC	
CLONE A	GGAATCT GGT	GGGACACA AT	ACGGCAGAA G	ATCCCTGAAT	GGTACGATGC	
Consensus	GGAAT-T GGT	GGGACAC- AT	-----AAG	AT-CC-GA-T	GG-----GC	
	201					250
SEQ ID NO: 83	AGGAATC TCA	GCGATATGGA	TTCCACCAG C	GAGTAAGGGC	ATGAGCGGTG	
SEQ ID NO: 85	AGGAATC TCA	GCGATATGGA	TTCCACCAG C	GAGTAAGGGA	ATGAGCGGTG	
SEQ ID NO: 75	CGGGATT TCG	GCGATATGGA	TTCCCCCG C	GAGCAAGGGT	ATGAGCGGCG	
SEQ ID NO: 77	CGGGATT TCG	GCGATATGGA	TCCCTCCCG C	GAGCAAGGGT	ATGAGCGGCG	
SEQ ID NO: 73	CGGGATT TCG	GCAATATGGA	TTCTTCCCG C	GAGTAAGGGC	ATGAGCGGCG	
SEQ ID NO: 79	CGGAATC TCC	GCAATATGGA	TTCCCCCG C	GAGCAAGGGC	ATGGGCGGCG	
SEQ ID NO: 81	GGGAATA TCC	GCCATTTGGA	TTCCGCCAG C	CAGCAAGGGG	ATGAGCGGCG	
CLONE A	AGGCATA TCC	GCCATCTGGA	TACCCCGG C	GAGCAAGGGC	ATGGGCGGGG	
Consensus	-GG-AT- TC-	GC-AT-TGGA	T-CC-CC-GC	-AG-AAGGG-	ATG-GCGG-G	

FIGURE 14C

	251		300
SEQ ID NO: 83	GTTATTC CAT GGGCTACG AT CCCTACGAT T TCTTTGACCT	CGGCGAGTAC	
SEQ ID NO: 85	GTTATTC CAT GGGCTACG AT CCCTACGAT T TCTTTGACCT	CGGCGAGTAC	
SEQ ID NO: 75	GCTATTC GAT GGGCTACG AC CCCTACGAT T ATTTTGACCT	CGGTGAGTAC	
SEQ ID NO: 77	GCTATTC GAT GGGCTACG AC CCCTACGAT T ATTTTGACCT	CGGTGAGTAC	
SEQ ID NO: 73	GCTATTC GAT GGGCTACG AC CCCTACGAT T TCTTCGACCT	CGGTGAGTAC	
SEQ ID NO: 79	CCTATTC GAT GGGCTACG AC CCCTACGAT T TCTTTGACCT	CGGTGAGTAC	
SEQ ID NO: 81	GTTACTC GAT GGGCTACG AT CCCTACGAT T TCTTTGACCT	CGGCGAGTAC	
CLONE A	CCTACTC GAT GGGCTACG AC CCCTACGAT T ACTTCGATCT	GGGCGAGTTT	
Consensus	--TA-TC-AT GGGCTACGA- CCCTACGA-T --TT-GA-CT	-GG-GAGT--	
	301		350
SEQ ID NO: 83	TATCAGA AGG GGACAGTT GA GACCGGCTT C GGCTCAAAGG	AAGAACTGGT	
SEQ ID NO: 85	TATCAGA AGG GGACAGTT GA GACCGGCTT C GGCTCAAAGG	AAGAACTGGT	
SEQ ID NO: 75	TACCAGA AGG GAACGGTG GA AACAAGATT C GGCTCAAAGC	AGGAGCTCAT	
SEQ ID NO: 77	TACCAGA AGG GAACGGTG GA AACAAGATT C GGCTCAAAGC	AGGAGCTCAT	
SEQ ID NO: 73	TACCAGA AGG GAACGGTT GA GACCCGCTT C GGATCAAAG	AGGAGCTTGT	
SEQ ID NO: 79	GACCAGA AGG GAACGGTGA GA GACCGGCTT T GGCTCAAAGC	AGGAGCTCGT	
SEQ ID NO: 81	AACCAGA AGG GAACCATC GA AACCGGCTT T GGCTCTAAAC	AGGAGCTCAT	
CLONE A	TACCAGA AGG GAACCGTT GA GACCCGCTT C GGCTCAAAGG	AAGAGCTCGT	
Consensus	-A-CAGA AGG G-A---T-GA -AC--G-TT -GG-TC-AA--	A-GA-CT--T	
	351		400
SEQ ID NO: 83	GAACATG ATA AACACCGC AC ACTCCTACG G CATAAAGGTG	ATAGCAGACA	
SEQ ID NO: 85	GAACATG ATA AACACCGC AC ACTCCTACG G CATAAAGGTG	ATAGCGGACA	
SEQ ID NO: 75	AAACATG ATA AACACCGC CC ACGCCTATG G CATGAAGGTA	ATAGCCGATA	
SEQ ID NO: 77	AAACATG ATA AACACCGC CC ACGCCTATG G CATGAAGGTA	ATAGCCGATA	
SEQ ID NO: 73	GAACATG ATA AACACCGC CC ATGCTCACA A CATGAAGGTC	ATAGCGGACA	
SEQ ID NO: 79	GAACATG ATA AACACCGC CC ACGCCTACG G CATCAAGGTC	ATCGCAGACA	
SEQ ID NO: 81	CAATATG ATA AACACGGC CC ATGCCTACG G CATAAAGGTC	ATAGCGGACA	
CLONE A	CAACATG ATC TCCACGGC CC ACCAGTACG G CATCAAGGTT	ATAGCGGACA	
Consensus	-AA-ATGAT- --CAC-GC-C A-----A---CAT-AAGGT-	AT-GC-GA-A	
	401		450
SEQ ID NO: 83	TAGTCAT AAA CCACCGCG CC GGTGGAGAC C TTGAGTGGAA	CCCCTTCGTG	
SEQ ID NO: 85	TAGTCAT AAA CCACCGCG CC GGTGGAGGC C TCGAGTGGAA	CCCCTTCGTG	
SEQ ID NO: 75	TAGTCAT CAA CCACCGCG CC GGCGGCGAT C TGGAGTGGAA	CCCCTTCGTG	
SEQ ID NO: 77	TAGTCAT CAA CCACCGCG CC GGCGGTGAC C TGGAGTGGAA	CCCCTTCGTG	
SEQ ID NO: 73	TAGTCAT CAA CCACCGCG CC GGCGGCGAC C TGGAGTGGAA	TCCTTTCACC	
SEQ ID NO: 79	TAGTAAT CAA CCACCGCG CC GGAGGAGAC C TTGAGTGGAA	CCCCTTCGTC	
SEQ ID NO: 81	TCGTAT AAA CCACCGCG CA GGCGGAGAC C TCGAGTGGAA	CCCCTTCGTT	
CLONE A	TAGTGAT AAA CCACCGCG CA GGTGGAGAC C TCGAATGGAA	CCCATACGTC	
Consensus	T-GT-AT-AA CCACCGCG C- GG-GG-G--C T-GA-TGGAA	-CC-T-C---	
	451		500
SEQ ID NO: 83	AACGACT ATA CCTGGACA GA CTTCTCAAA A GTCGCCTCGG	GTAAATATAC	
SEQ ID NO: 85	AACGACT ATA CCTGGACA GA CTTCTCAAA A GTCGCCTCGG	GTAAATATAC	
SEQ ID NO: 75	AACGACT ATA CCTGGACC GA CTTCTCGAA G GTCGCTCGG	GTAAATACAC	
SEQ ID NO: 77	AACGACT ATA CCTGGACC GA CTTCTCAAA G GTCGCTCGG	GTAAATACAC	
SEQ ID NO: 73	AACAGCT ACA CCTGGACC GA TTTCTCGAA G GTCGCTCGG	GCAAGTACAC	
SEQ ID NO: 79	AATGACT ACA CCTGGACG GA CTTCTCGAA G GTCGCTTCGG	GCAAGTACAC	
SEQ ID NO: 81	GGGGACT ACA CCTGGACG GA CTTCTCAAA G GTGGCCTCGG	GCAAATATAC	
CLONE A	GGCGACT ATA CCTGGACG GA CTTTTCTAA G GTCGCCTCGG	GGAAATACAA	
Consensus	-----CTA-A CCTGGAC- GA -TT-TC-AA -GT-GC-TC-G	G-AA-TA-A-	

FIGURE 14C
(cont.)

	501		550
SEQ ID NO: 83	GGCCAAC TAC	CTTGACTT CC	ACCCAAACG A GCTTCACTGT TGTGATGAAG
SEQ ID NO: 85	AGCCAAC TAC	CTTGACTT CC	ACCCAAACG A GCTTCACTGT TGTGATGAAG
SEQ ID NO: 75	GGCCAAC TAC	CTCGACTT CC	ACCCGAACG A GCTCCACGCG GGCGATTCCG
SEQ ID NO: 77	GGCCAAC TAC	CTCGACTT CC	ACCCGAACG A GCTCCATGCG GGCGATTCCG
SEQ ID NO: 73	GGCCAAC TAC	CTCGACTT CC	ACCCGAACG A GCTTCACGCG GGCGATTCCG
SEQ ID NO: 79	GGCCAAC TAC	CTCGACTT CC	ACCCCAACG A GGTCAAGTGC TGCGACGAGG
SEQ ID NO: 81	TGCCAAC TAC	CTCGACTT CC	ACCCCAACG A GGTCAAGTGC TGTGACGAGG
CLONE A	GGCCCAAC TAC	ATGGACTT CC	ATCCAAACA A CTACAGCACC TCAGACGAGG
Consensus	-GCC-AC TAC	-T-GACTT CC	A-CC-AAC- A ----- -GA----G
	551		600
SEQ ID NO: 83	GTACCTT TGG	AGGATACC CT	GATATATGT C ACGACAAAAG CTGGGACCAG
SEQ ID NO: 85	GTACCTT TGG	AGGATACC CT	GATATATGT C ACGACAAAAG CTGGGACCAG
SEQ ID NO: 75	GAACATT TGG	AGGCTATC CC	GACATATGC C ACGACAAGAG CTGGGACCAG
SEQ ID NO: 77	GAACATT TGG	AGGCTATC CC	GACATATGC C ACGACAAGAG CTGGGACCAG
SEQ ID NO: 73	GAACATT TGG	AGGCTATC CC	GACATATGC C ACGACAAGAG CTGGGACCAG
SEQ ID NO: 79	GCACCTT TGG	AGGGTTCC CG	GACATAGCC C ACGAGAAGAG CTGGGACCAG
SEQ ID NO: 81	GCACATT TGG	AGGCTTCC CA	GACATAGCC C ACGAGAAGAG CTGGGACCAG
CLONE A	GAACCTT CGG	TGGCTTCC CA	GACATTGAT C ACCTCGTGCC CTTCAACCAG
Consensus	G-AC-TT-GG	-GG-T--CC-	GA-AT----C AC----- CT---ACCAG
	601		650
SEQ ID NO: 83	TACTGGC TCT	GGGCGAGC AG	CGAAAGCTA C GCTGCCTACC TCAGGAGCAT
SEQ ID NO: 85	TACTGGC TCT	GGGCGAGC AG	CGAAAGCTA C GCTGCCTACC TCAGGAGCAT
SEQ ID NO: 75	TACTGGC TCT	GGGCCAGC CA	GGAGAGCTA C GCGGCCTATC TCAGGAGCAT
SEQ ID NO: 77	TACTGGC TCT	GGGCCAGC CA	GGAGAGCTA C GCGGCATATC TCAGGAGCAT
SEQ ID NO: 73	CACTGGC TCT	GGGCCAGC AA	CGAAAGCTA C GCCGCCTACC TCCGAGCAT
SEQ ID NO: 79	TACTGGC TCT	GGGCGAGC AA	CGAGAGCTA C GCCGCCTACC TCAGGAGCAT
SEQ ID NO: 81	CACTGGC TCT	GGGCGAGC GA	TGAGAGCTA C GCCGCCTACC TAAGGAGCAT
CLONE A	TACTGGC TGT	GGGCGAGC AA	CGAGAGCTA C GCCGCCTACC TCAGGAGCAT
Consensus	-ACTGGC T-T	GGGC-AGC --	-GA-AGCTA C GC-GC-TA-C T--GGAGCAT
	651		700
SEQ ID NO: 83	AGGGGTT GAC	GCCTGGCG TT	TCGACTACG T CAAGGGCTAC GGAGCATGGG
SEQ ID NO: 85	AGGGGTT GAC	GCCTGGTG TT	TCGACTACG T CAAGGGCTAC GGAGCCTGGG
SEQ ID NO: 75	CGGCATC GAC	GCCTGGCG CT	TCGACTACG T CAAGGGCTAT GCTCCCTGGG
SEQ ID NO: 77	CGGCATC GAT	GCCTGGCG CT	TCGACTACG T CAAGGGCTAT GCTCCCTGGG
SEQ ID NO: 73	CGGCATC GAC	GCCTGGCG CT	TCGACTACG T CAAGGGCTAC GCTCCCTGGG
SEQ ID NO: 79	CGGCGTT GAC	GCATGGCG CT	TCGACTACG T CAAGGGCTAC GGAGCGTGGG
SEQ ID NO: 81	CGGCGTT GAT	GCCTGGCG CT	TTGACTACG T GAAGGGCTAC GGAGCGTGGG
CLONE A	AGGGATC GAT	GCGTGGCG CT	TTGACTACG T TAAGGGCTAC GGCGCGTGGG
Consensus	-GG--T-GA-	GC-TGG-G-T	T-GACTACG T -AAGGGCTA- G---C-TGGG
	701		750
SEQ ID NO: 83	TTGTTAA CGA	CTGGCTCA GC	TGGTGGGGAG GCTGGGCCGT TGGAGAGTAC
SEQ ID NO: 85	TTGTTAA CGA	CTGGCTCA GC	TGGTGGGGAG GCTGGGCCGT TGGAGAGTAC
SEQ ID NO: 75	TCGTCAA GGA	CTGGCTGA AC	TGGTGGGGAG GCTGGGCAGT TGGAGAGTAC
SEQ ID NO: 77	TCGTCAA GGA	CTGGCTGA AC	TGGTGGGGAG GCTGGGCCGT TGGAGAGTAC
SEQ ID NO: 73	TCGTCAA GAA	CTGGCTGA AC	CGGTGGGGCG GCTGGGCCGT TGGAGAGTAC
SEQ ID NO: 79	TCGTCAA GGA	CTGGCTGG AC	TGGTGGGGAG GCTGGGCCGT CGGGAGTAC
SEQ ID NO: 81	TCGTCAA GGA	CTGGCTCA AC	TGGTGGGGCG GCTGGGCCGT TGGCAGTAC
CLONE A	TCGTCAA GGA	CTGGCTGA GT	CAGTGGGGCG GCTGGGCCGT CGGCGAGTAC
Consensus	T-GT-A- --A	CTGGCT-- --	--GTGGG- G GCTGGGC-GT -GG-GAGTAC

FIGURE 14C

(cont.)

	751		800
SEQ ID NO: 83	TGGGACA CGA	ACGTTGAT GC	ACTCCTCAA C
SEQ ID NO: 85	TGGGACA CTA	ACGTTGAT GC	ACTCCTCAA C
SEQ ID NO: 75	TGGGACA CCA	ACGTCGAC GC	TGTTCTCAA C
SEQ ID NO: 77	TGGGACA CCA	ACGTCGAC GC	TGTTCTCAA C
SEQ ID NO: 73	TGGGACA CCA	ACGTCGAT GC	ACTCCTGAG C
SEQ ID NO: 79	TGGGACA CAA	ACGTTGAT GC	ACTGCTCAA C
SEQ ID NO: 81	TGGGACA CCA	ACGTTGAT GC	ACTCCTCAA C
CLONE A	TGGGACA CCA	ACGTCGAT GC	GCTCCTCAA C
Consensus	TGGGACA C-A	ACGT-GA-GC	--T-CT-A-C
	801		850
SEQ ID NO: 83	CAAGGTC TTT	GACTTCCC GC	TCTACTACA A
SEQ ID NO: 85	CAAGGTC TTT	GACTTCCC GC	TCTACTACA A
SEQ ID NO: 75	CAAGGTC TTT	GACTTCGC CC	TCTACTACA A
SEQ ID NO: 77	CAAGGTC TTT	GACTTCGC CC	TCTACTACA A
SEQ ID NO: 73	TAAAGTC TTC	GACTTCCC GC	TCTACTACA A
SEQ ID NO: 79	AAAAGTC TTC	GACTTCCC GC	TCTACTACA A
SEQ ID NO: 81	CAAGGTC TTC	GACTTCCC GC	TCTACTACA A
CLONE A	CAAGGTC TTC	GACTTCCC GC	TCTACTACA A
Consensus	-AA-GTCTT-	GACTTC-C-C	TCTACTACA A
	851		900
SEQ ID NO: 83	ACACCAA CAT	CCCGGCAT TA	GTGGATGCA C
SEQ ID NO: 85	ATACCAA CAT	CCCGGCTT TG	GTTTACGCC C
SEQ ID NO: 75	ACAACAA CAT	TCCCGCCC TG	GTGGACGCC C
SEQ ID NO: 77	ACAACAA CAT	TCCCGCCC TG	GTGGACGCC C
SEQ ID NO: 73	ACAACAA CAT	CCCGGCCCTC	GTGGACGCC C
SEQ ID NO: 79	ACAAGAA CAT	TCCCGCAC TC	GTCGAGGCC C
SEQ ID NO: 81	ACAAAAA CAT	TCCAGCGC TC	GTCTCTGCC C
CLONE A	ACAAGAA CAT	TCCCGCCC TC	GTTTACGCC A
Consensus	A-A--AA CAT	-CC-GC--T-	GT----GC--
	901		950
SEQ ID NO: 83	GTCAGCC GCG	ATCCCTTCAA	GGCGGTAAC T
SEQ ID NO: 85	GTCAGCC GCG	ACCCATTCAA	GGCGGTAAC T
SEQ ID NO: 75	GTCAGCC GCG	ACCGGTTCAA	GGCTGTGAC G
SEQ ID NO: 77	GTCAGCC GCG	ACCGGTTCAA	GGCTGTGAC G
SEQ ID NO: 73	GTCAGCC GCG	ACCGGTTCAA	AGCCGTGAC C
SEQ ID NO: 79	GTCAGCC GCG	ACCGGTTTAA	GGCCGTAAC C
SEQ ID NO: 81	GTCTCCC GCG	ACCGGTTCAA	GGCCGTAAC C
CLONE A	GTCAGCA GGG	ATCCCTTCAA	GGCGGTTAC C
Consensus	GTC--C-G-G	A-CC-TT-AA	-GE-CT-AC-
	951		1000
SEQ ID NO: 83	AGATATA ATC	TGGAACAA GT	ATCCGGCTT A
SEQ ID NO: 85	AGATATA ATC	TGGAACAA GT	ATCCGGCTT A
SEQ ID NO: 75	CGACATA ATC	TGGAACAA GT	ATCCAGCCT A
SEQ ID NO: 77	CGACATA ATC	TGGAACAA GT	ATCCAGCCT A
SEQ ID NO: 73	CAACATA ATC	TGGAACAA GT	ATCCGGCCT A
SEQ ID NO: 79	GGACATA ATT	TGGAACAA GT	ACCCGGCCT A
SEQ ID NO: 81	CGATATA ATC	TGGAACAA GT	ACCTTGCTT A
CLONE A	GAACATA ATC	TGGAACAA GT	ACCTTGCTT A
Consensus	--A-ATAAT-	TGGAACAA GT	A-C--GC-TA

FIGURE 14C
(cont.)

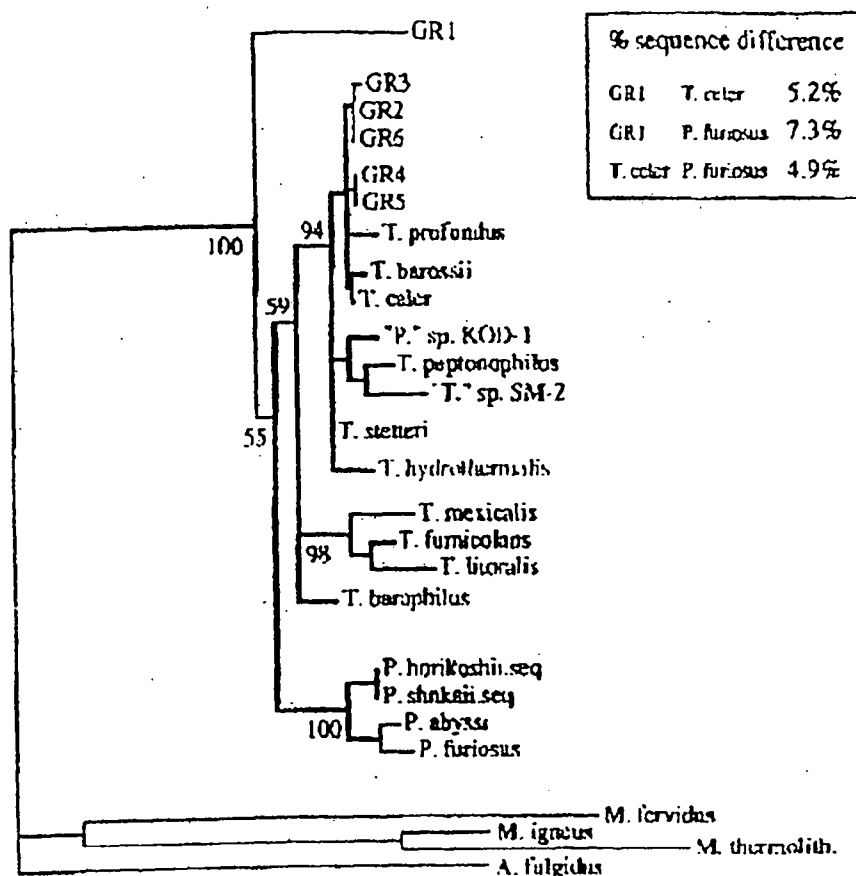
	1001		1050
SEQ ID NO: 83	AGGGACAGCC	TGTTATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 85	AGGGACAGCC	TGTTATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 75	AGGGCCAGCC	GACAATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 77	AGGGCCAGCC	GACAATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 73	AGGGACAGCC	GGCAATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 79	AGGGCCAGCC	GACGATATTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
SEQ ID NO: 81	AAGGCCAGCC	CGTCATATTT	TACCGCGACT ACGAGGAGTG GCTCAACAAG
CLONE A	AAGGTCA GCC	CGTCATCTTC	TACCGCGACT ACGAGGAGTG GCTCAACAAG
Consensus	A-GG-CA GCC	----AT-TT-	TACCGCGACT ACGAGGAGTG GCTCAACAAG
	1051		1100
SEQ ID NO: 83	GATAAGCTTA	ACAACCTCAT	CTGGATACAC GATCACCTTG CTGGAGGGAG
SEQ ID NO: 85	GATAAGCTTA	ACAACCTCAT	CTGGATACAC GATCACCTTG CTGGAGGGAG
SEQ ID NO: 75	GACAAGCTCA	AGAACCTCAT	CTGGATACAT GACAACCTCG CCGGAGGGAG
SEQ ID NO: 77	GATAAGCTCA	AGAACCTCAT	CTGGATACAT GACAACCTCG CCGGAGGGAG
SEQ ID NO: 73	GACAGGCTCA	GGAACTCTAT	CTGGATACAC GACCACCTCG CCGGAGGAAG
SEQ ID NO: 79	GACAGGCTCA	AGAACCTCAT	CTGGATACAC GACCACCTCG CCGGTGGAAG
SEQ ID NO: 81	GACAGGTGTA	ACAACCTCAT	ATGGATACAC GACCACCTCG CAGGTGGAAG
CLONE A	GACAAACTCA	ACAACCTCAT	ATGGATTCA C GAGCACCTGG CAGGGGGAAG
Consensus	GA-A---T-A	--AACCTCAT	-TGGAT-CA - GA--ACCT-G C-GG-GG-AG
	1101		1150
SEQ ID NO: 83	TACTGACATT	GTTTACTACG	ACAGCGACGA GCTTATCTTT GTGAGAAACG
SEQ ID NO: 85	TACTGACATT	GTTTACTACG	ACAGCGACGA GCTTATCTTT GTGAGAAACG
SEQ ID NO: 75	CAC TGACATC	GTTTACTACG	ACAACGACGA GCTGATATTC GTGAGAAACG
SEQ ID NO: 77	CAC TGACATC	GTTTACTACG	ACAACGACGA GCTGATATTC GTGAGAAACG
SEQ ID NO: 73	CACAGACATC	ATCTACTACG	ACAGCGACGA GCTTATCTTC GTGAGAAACG
SEQ ID NO: 79	CACCGACATA	GTCTACTACG	ATAACGATGA ACTCATCTTC GTGAGAAACG
SEQ ID NO: 81	CACGAGCATA	GTTTACTACG	ACAGCGACGA GATGATTTTC GTGAGGAACG
CLONE A	CACCAAGATC	CTCTACTACG	ACGACGATGA GCTCATCTTC ATGAGGGAAG
Consensus	-AC----AT-	-T-TACTACG	A---CGA-GA --T-AT-TT- -T-AG--A-G
	1151		1200
SEQ ID NO: 83	GCTATGGCAC	CAAACCAGGA	CTGATAACCT ATATCAACCT CGGCTCAAGC
SEQ ID NO: 85	GCTATGGCAC	CAAACCAGGA	CTGATAACCT ATATCAACCT CGGCTCAAGC
SEQ ID NO: 75	GCTACGGAAG	CAAGCCGGGA	CTGATAACAT ACATCAACCT CGGCTCAAGC
SEQ ID NO: 77	GCTACGGAAG	CAAGCCGGGA	CTGATAACAT ACATCAACCT CGGCTCAAGC
SEQ ID NO: 73	GCTACGGGA	CAAGCCGGGG	CTTATAACCT ACATCAACCT AGGCTCGAGC
SEQ ID NO: 79	GCTACGGGA	CAAGCCGGGG	CTTATAACTT ACATCAACCT CGGCTCGAGC
SEQ ID NO: 81	GCTATGGGA	CAAGCCGGGC	CTTATAACCT ACATCAACCT CGGTAGCGAC
CLONE A	GCTACGGCGA	CAGGCCGGGG	CTTATAACCT ACATCAACCT CGGTAGCGAC
Consensus	GCTA-GG---	CA--CC-GG-	CT-ATAAC-T A-ATCAACCT -G-----C
	1201		1250
SEQ ID NO: 83	AAAGTTGGAA	GGTGGGTC TA	CGTT...CCA AAGTTCCGG GTTCATGCAT
SEQ ID NO: 85	AAAGCTGGAA	GGTGGGTC TA	CGTT...CCA AAGTTCCGG GTTCATGCAT
SEQ ID NO: 75	AAAGCCGGAA	GGTGGGTT TA	CGTT...CCG AAGTTCCGAG GCTCGTGCAT
SEQ ID NO: 77	AAAGCCGGAA	GGTGGGTT TA	CGTT...CCG AAGTTCCGAG GCTCGTGCAT
SEQ ID NO: 73	AAGGCCGGAA	GGTGGGTC TA	CGTT...CCG AAGTTCCGAG GCTCGTGCAT
SEQ ID NO: 79	AAGGCCGGAA	GGTGGGTC TA	CGTT...CCG AAGTTCCGAG GCTCGTGCAT
SEQ ID NO: 81	AAGGTTGGAA	GGTGGGTT TA	TGTG...CCG AAGTTCCGGG GCGCGTGCAT
CLONE A	TGGGCCGAGA	GATGGGTGAA	CGTTGGCTCA AAGTTCCGGG GCTATACAAT
Consensus	---G--G--A	G-TGGGT--A	-GT-----C- AAGTTCCG-G G-----AT

FIGURE 14C
(cont.)

	1251		1300
SEQ ID NO: 83	CCACGAG TAC ACCGGCAA CC TCGGCGGTT G GATAGACAAG	TACGTCTCCT	
SEQ ID NO: 85	CCACGAG TAC ACCGGCAG CC TCGGCGGTT G GATAGACAAG	TACGTCTCCT	
SEQ ID NO: 75	ACACGAG TAC ACCGGCAA CC TCGGCGGCT G GGTGGACAAG	TGGGTGGACT	
SEQ ID NO: 77	ACACGAG TAC ACCGGCAA TC TCGGCGGCT G GGTGGACAAG	TGGGTGGACT	
SEQ ID NO: 73	ACACGAG TAC ACCGGCAA CC TCGGCGGCT G GATTGACAAG	TGGGTGGACT	
SEQ ID NO: 79	CCACGAG TAC ACCGGCAA CC TCGGCGGCT G GGTGGACAAG	TGGGTGGACT	
SEQ ID NO: 81	CCACGAG TAT ACTGGTAA CC TCGGAGGCT G GGTAGACAAG	TACGTCTACT	
CLONE A	CCACGAA TAC ACCGGAAA CC TCGGCGGCT G GGTGACAGG	TACGTCCAGT	
Consensus	-CACGA- TA- AC-GG-A- -C TCGG-GG-TG G-T-GACA-G	T--GT----T	
	1301		1350
SEQ ID NO: 83	CCAGCGG CTG GGTCTATC TT GAGGCCCCAG CCCACGACCC	GGCGAACGGC	
SEQ ID NO: 85	CCAGCGG CTG GGTCTACC TT GAGGCCCCGG CCCACGACCC	GGCCAATGGC	
SEQ ID NO: 75	CAAGCGG CTG GGTTTACC TC GAGGCTCCT G CCCACGACCC	GGCCAACGGC	
SEQ ID NO: 77	CAAGCGG CTG GGTCTACC TC GAGGCTCCT G CCCACGACCC	GGCCAACGGC	
SEQ ID NO: 73	CAAGCGG TCG GGTCTACC TT GAGGCCCCCG CCCACGACCC	GGCCAACGGC	
SEQ ID NO: 79	CAAGCGG GTG GGTGTACC TC GAGGCCCTG CCCACGACCC	GGCCAACGGC	
SEQ ID NO: 81	CAAGCGG CTG GGTCTATT TC GAAGCTCCAG CTTACGACCC	TGCCAACGGG	
CLONE A	ACGACGG CTG GGTCAAGC TT ACCGCTCCG C CACAGATCC	GGCAAACGGC	
Consensus	----CGG--G GGT--A--T- ---GC-CC-- C--ACGA-CC	-GC-AA-GG-	
	1351		1393
SEQ ID NO: 83	TACTACG GCT ACTCCGTA TG GAGCTACTG C GGGGTGGGT	GA-	
SEQ ID NO: 85	CAGTATG GCT ACTCCGTC TG GAGCTATTG C GGGGTGGGT	GA-	
SEQ ID NO: 75	CAGTACG GCT ACTCCGTT TG GAGCTATTG C GGTGTTGGGT	GA-	
SEQ ID NO: 77	CAGTACG GCT ACTCCGTC TG GAGCTACTG C GGTGTTGGGT	GA-	
SEQ ID NO: 73	CAGTACG GCT ACTCCGTA TG GAGCTACTG C GGTGTTGGGT	GA-	
SEQ ID NO: 79	TATTACG GCT ACTCCGTC TG GAGCTACTG C GGGGTGGGCT	GA-	
SEQ ID NO: 81	CAGTATG GCT ACTCCGTG TG GAGCTATTG C GGTGTTGGGT	GA-	
CLONE A	TATTACG GCT ACTCCGTC TG GAGCTACGC C GGAGTTGGAT	GA-	
Consensus	-A-TA-GGCT ACTC-GT- TG GAGCTA--- C GG-GT-GG-T	GA-	

FIGURE 14C
(cont.)

Neighbor-joining tree for Thermococcales



0.01

bootstrap values for 100 replicates

Summit & Barnet, Deep-Sea Research Pt. II, in press

FIGURE 15

FIGURE 16A

SEQ ID NO.: 1

atggcaaaagtattccgagctcgaagagggcgggctcataatgcaggccttctactgggacgtcccatgggaggaatctgggtggacacgat
 agcccagaagatacccgactgggcaagcgccgggatttcggcgataggattccccggcgagcaaggcatgggcgccgctattcgtatg
 ggctacgaccctacgacttctttgacctgggtgagtagaccagaagggaacggtagagacgcgtttggctccaagcaggagctcgtgaa
 catgataaacaccgccacgcctatggcatgaaggtaatagccgatatagtcataaccacgcgcggcggtgacctggagtggaaacctt
 cgtgaacgactatactggaccgacttctcaaggtcgcgtcgggtaatacacggccaactacctgacttccacccgaacgagctccatgc
 gggcgattccggaacatttggaggctatcccacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctac
 gcggcatatctcaggagcatcggcatcgatgcctggcgcttcgactacgtcaagggtacggagcgtgggtcgtcaaggactggctggactg
 gtggggaggctgggctggcggtgggagtagtgggacacaaacgttgatgcactgctcaactgggctactcagcgtatgaaaagtcttcgactt
 ccgcctactacaagatggagcggcctttgacaacaagaacattcccgactcgtcaggccctcaagaacgggggcacagctcgtcagcc
 gcgaccgtttaaggccgtaaccttcgttgaaccacgacaccgataatactggaacaagtatccagcctacgcgttcacctcacctacgag
 ggccagccgacaatatttaccgcgactacgaggagtggctcaacaaggataagctcaagaacctcatctggatacatgacaacctcggcg
 aggaagcactgacatggttactacgacaacgacgagctgatattcgtgaaacggctacggaagcaagccgggactgataacatacatcaa
 cctcgcctcaagcaagccggaaggtgggtttacgttcgaagttcgcaggctcgtgcatacacgagtaaccggcaatcgcggcggtgggt
 ggacaagtggtggactcaagcggtgggtctacctcaggctcctgccacgaccggccaacggccagtaggctactcgtctggagc
 tactcggtgttggtga

SEQ ID NO.: 2

Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 3

atggccaagtacctggagctcgaagagggcgggctcataatgcaggccttctactgggacgtcccatgggaggaatctgggtggacacgat
 agcccagaagatacccgactgggcaagcgccgggatttcggcgataggattccccggcgagcaaggcatgggcgccgctattcgtatg
 ggctacgaccctacgacttctttgacctgggtgagtagaccagaagggaacggtagagacgcgtttggctccaagcaggagctcgtgaa
 catgataaacaccgccacgcctacggcatcaaggtcgcgacacatagtaatcaaccaccgcgcggaggagaccttgagtggaaacctt
 tcgtcaatgactacacctggacggacttctgaaggctcctccggcaagtagcacggccaattacctgacttccaccgaacgagctccatgc
 gggcgattccggaacatttggaggctatcccacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctac
 gcggcatatctcaggagcatcggcatcgatgcctggcgcttcgactacgtcaagggtatgctcctgggtcgtcaaggactggctgaactggt
 ggggaggctggcggttgagagtagtgggacaccaacgtcgcgctgttctcaactgggcatactcagcgggtgccaaggtctttgacttgc
 cctctactacaagatggatgaggcctttgacaacaaaacattccagcgtcgtctcgtcccttcagaacggccagactgttctccccgcgac
 ccgttcaaggccgtaacctttgtagcaaacacgacaccgataatactggaacaagtagtcagcctacgcgttcactcctacctcaggggcc

FIGURE 16B

agccgacaatatctaccgcgactacgaggagtggtcacaagaaggataagctcaagaacctcatctggatacatgacaacctcgccggagga
 agcactgacatcggttactacgacaacgacgagctgatattcgtgagaaacggctacggaagcaagccgggactgataacatacatcaacctc
 gcctcaagcgaagccggaaggtgggtctacgttccgaagttcgcgggagcgtgcatccacgagtacaccggcaacctcgccggctgggtgg
 acaagtgggtggactcaagcgggtgggtgtacctcagggccctgcccacgaccggccaacggctattacggctactcctgtctggagctatt
 gcgggtgtgggtga

SEQ ID NO.: 4

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Ala Ser Ser Glu Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 5

atggccaagtactccgagctggaagagggcgcggtataatgcaggcccttctactgggacgtccaggtggaggaatctggtgggacaccat
 caggagcaagataccggagtggtacgagggcggaataatccgccatttggattccccggcaagcaagggcatggcgccctattcgtatg
 ggctacgacccccacgactctttgacctcgggtgagtagcaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
 catgataaacaccgcccaacgcctatggcatgaaggtaataagccgatatagtcatcaaacaccgcgcggcggtgacctggagtggaacccctt
 cgtgaacgactatactggaccgactctcaaaaggtcgcgtcggttaatacacggccaactacctgacttccacccgaacgagctccatgc
 gggcgattccggaacatttggaggctatcccgacatatgccacgacaagagctgggaccagtactggctctggggccagccaggagagctac
 gcggcatatctcaggagcatcgccatcgatgcctggcgcttcgactacgcaagggtatgctccctgggtcgtcaaggactggctgaactggt
 ggggaggctggcggttggaggtactgggaacccaagctgacgctgttctcaactgggcatactcgagcggttgcgaaggcttgaacttcg
 cctctactacaagatggatgggctttgacaacaaaasattcagagctcgtctctgcccctcagaacggcgaactgttctccgggae
 ccgttcaaggccgttaactttagcaaacacgacaccgataatactggaacaagtacttgcctatgcttctacacacgaaggccag
 cccgtcatattctaccgcgaccacgaggagtggtcaacaaggacaggttgaacaacctcatatggatacacgaccacctcgcaggtggaag
 caccgacatagtctactacgataacgatgaactcatcttcgtcaggaaaggctacggggacaagccggggcttataacctacatcaacctagggc
 togagcaaggccggaagggtggttatgtgccgaagttcggcgcggtgcatccacgagtatactggaacctcggaggctgggtagacaa
 gtacgtctactcaagcggtgggtctatctgaagctcagcttacgacctgccaacgggcagtagtggtactcctgtggagctactgcggg
 gtgggctga

SEQ ID NO.: 6

Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly

FIGURE 16C

Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp His Glu Glu Trp Leu Asn
 Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 9

atggccaagtactccgagctggaagagggcgggctcataatgcaggccttctactgggacgtcccatgggaggaatctggtggacacgat
 agcccagaagatacccgactgggcaagcgccgggatttcggcgatattgattccccggcgagcaagggcatggcgggcctattcgatg
 ggctacgacccctacgacttcttgacctcggtgagtacgaccagaagggaaacggtagagacgcgtttggctccaagcaggagctcgtgaa
 catgataaacacggcccatgcctacggcataaaggctcatagcggacatcgtcataaaccacgcgcaggcggagacctcgagtggaaacccg
 ttctgtgggactacacctggacggacttctcaaaagggtggcctcgggcaaatatactgccaactacctcgacttccaccgaacgagctccatg
 cggcgatccggaacattggaggctatcccgacatagccacgacaagagctgggaccagtactggctctgggccagccaggagagctac
 gggcgatactcaggagcatcggcatcgtgctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactggt
 ggggaggtcggcggttgagagtactgggacaccaacgtcgacgtgttctcaactgggcatactcgagcgggtgccaaggcttctgacttcg
 ccttactacaagatggacgaggccttcgataacaacaacattcccgccctgggtggacgcccicagatacggtcagacagtgtggtcagccgcg
 acccgttcaaggctgtgacgtttgtagccaaccacgataccgataatctggaacaagtatccagcctacgcgttcacctcaccctacgagggc
 cagccgacaatatctaccgcgactacgaggagtggctcaacaaggataagctcaagaacctcatctggatacatgacaacctcgcggagg
 aagcactgacatcgtttactacgacaacgacgagctgatattcgagaaacggctacggaagcaagccgggactgataacatacatcaacct
 cgctcaagcaaaagccggaaggtgggtttacgttcggaagtcgcaggctcgtgcatacacgagtacaccggcaatctcggcggttgggtgg
 acaagtggtgggactcaagcggtggtctacctcgaggtccttgcacacgacccggccaacggccagtcaggctactccgtctggagctac
 tgcggtgttgggtga

SEQ ID NO.: 10

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 Val Lys Asp Trp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Ala Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu

FIGURE 16D

Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 11

atggccaagtacctggagctcgaggaggcgggctcataatgcaggccttctactgggacgtccccatgggaggaatctggtgggacacgat
agcccagaagatacccgactgggcaagcggcggttgcgatatggattccccggcgagcaaggcgatggcgggcgctattcgtatg
ggctacgacctacgacttcttgacctcggtagtagcaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
catgataaacaccgccacgcctatggcatgaaggtaatagccgatatagtcataaccaccgcggcggtgacctggagtggaacccctt
cgtgaacgactatacctggaccgacttctcaaaaggctcgcgtcgggtaataacacggccaactacctgacttccaccgaacgagctccatgc
ggcggtattccggaacatttggaggctatcccgacatagccacgacaagagctgggaccagtactggctctggccagccaggagagctac
gcggcctatctcaggagcatcggcatcgtatgcttggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactggt
ggggaggctgggctgggtggagagtagtgggacaccaacgtcgcagctgttctcaactgggcatactcgagcgggtgccaaggcttctgacttcg
cccttactacaagatggacgaggccttcgataacaacaactcccgccctggtggacgccctcagatacggtcagacagtggtcagccgcg
accgttcaaggctgtgactgttagccaaccacgataccgataatctggaacaagtatccagcctacgcttcacctacacctacgagggc
cagccgacaatatctaccgcgactacgaggagtggtcaacaaggatagctcaagaacctcatctggatacatgacaacctcggcgagg
aagcagcagcatalgttactacgacagcagcagatgatctcgtgagggaacggctatggaagcaagcctggcctataactacatcaacctc
ggctcagcaaggttgaagggtgggtctacgttccgaagctcggggagcgtgcatccacgagtacaccggcaacctcggcggtggtggtg
acaagtggtggtgactcaagcgggtgggtgtacctcaggccctgccacgacccggccaacggctattacggctactccgtctggagctac
tgcggtgttgctga

SEQ ID NO.: 12

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
Ala Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg
Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
Lys Asp Thr Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr
Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 13

atggccaagtacctggagctcgaggaggcgggctcataatgcaggccttctactgggacgtgccttcaggaggaatattggtgggacacaat
acggcagaagataccggagtggtagtgatccggaatctccgcaatatggattccccggcgagcaaggcgatggcgggcgctattcgtatg
ggctacgacctacgacttcttgacctcggtagtagtagcaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
atgataaacacggcacatgctcagcgcataaaggctatagcggacatcgtcataaccaccgcgcaggcggagacctcagtggaacccgtt
cgttgggactacacctggacggacttctcaaaaggctggcctcgggcaataactgccaactacctgacttccaccccaacgagggtcaagt
ctgtgacgagggcacatttggaggcttccagacatagccacgagaagagctgggaccagcactggctctggcgagcgtatgagagctac
ggcgcttacctaaaggagcatcggcggttggatgcttggcgttcgactacgtcaagggtcagcggcgtgggtcgtcaaggactggctggactg
gtggggaggctgggctcggggagtagtgggacacaaacgtttagtgcactgtctcaazctgggcttactcggcgtatgcaaaagtctcga

FIGURE 16E

cccgtctactacaagatggatgaggcctttgacaacaaaacattccagcgctcgtctctgcccttcagaacggccagactgttgtctcccgcg
 acccggtcaaggccgtaacctttgtgcaaacacgacaccgatataatctggaacaagtatccagcctacgcgttcacctcacctacgaggg
 ccagccgacaatatctaccgcgactacgaggagtggtcaacaaggataagctcaagaacctcatctggatacatgacaacctcgccggag
 gaagcactgacatagctactacgataacgatgaactcatcttcgtcaggaacggctacggggacaagccggggcctataacctacatcaacct
 aggcctcgagcaaggccggaagggtgggttatgtgccgaagttcgccggcgcggtgcattccacgagtatactgtaacctcgagggtggtag
 acaagctactctactcaagcggtgggtctatctcgaagctccagcttacgacctgccaacgggcagtatggctactccgtgtggagctactg
 cgggtgtggctga

SEQ ID NO.: 14

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
 Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
 Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
 Val Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
 Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
 Glu Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
 Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
 Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val
 Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His
 Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu
 Ala Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 15

atggccaagtactccgagctggaagagggcgggctcataatgcaggccttctactgggacgtccccatgggaggaatctggtgggacacgat
 agcccagaagatacccgaactgggcaagcggcggttctggcgatattggattccccggcgagcaagggeatggcgggcgctattcgatg
 ggctacgacccctacgacttctttgacctcggtagtacgaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
 catgataaacacggcccatgcctacggcataaaggctatagcggacatcgtcataaaccaccgcgcaggcgaggacctcgaagtgaacccg
 ttcgttggggactacacctggacggacttctcaaaagggtggcctcgggcaaatatactgccaactacctgacttcaccccaagcagctccatg
 cgggcgattccggaacatttgaaggctatccgacatatgcacgacaagagctgggaccagtaetggctotgggccaagcaggagagctac
 ggggcatalctcaggagcattgggtcgaatgctggcgattggactacgttcagggtcaggagcgtgggttggcgaagcactgggtggactg
 gtggggaggctggggtcggggagttactgggacacaaacgttgatgcactgctcaactgggctactcgaagcgtgcaaaagtcttcgactt
 cccgtctactacaagatggatgaggcctttgacaacaaaacattccagcgctcgtctctgcccttcagaacggccagactgttgtctcccgcg
 acccggtcaaggccgtaacctttgtgcaaacacgacaccgatataatttgaacaagtacccggcctacgccttcacctcacctacgaggg
 ccagccgacgatattctaccgcgactacgaggagtggtcaacaaggacaggctcaagaacctcatctggatacacgaccaccttggcggtg
 gaagcactgacatcgtttactacgacaacgacgagctgatattcgtgagaaacggctacggaagcaagccgggactgataacatacatcaacc
 tcgcctcaagcaaaagcgggaagggtgggttatgtgccgaagttcgccggcgcggtgcattccacgagtatactgtaacctcgagggtggtag
 acaagctactctactcaagcggtgggtctatctcgaagctccagcttacgacctgccaacgggcagtatggctactccgtgtggagctattgc
 ggtgttgggtga

SEQ ID NO.: 16

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp

FIGURE 16F

Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 17

atggccaagtactccgagctggaagggggcgggcataatgcaggccttctactgggacgtcccatgggaggaatctgggtgggacacgat.
 agcccagaagatacccgactgggcaagcgcgggattcggcgatatggattccccggcgagcaagggcatgggcggcgctattcgatg
 ggctacgacctacgacttcttgacctgggtgagtacgaccaggagggaacggtagagacgcgcttggctccaagcaggagctcgtgaa
 catgataaacacggccatgctacggcataaaggatagcggacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccg
 ttctgtggggactacacctggacggacttctcaaaagggtgctcgggcaataactgccaactacctgacttccaccccaacgaggtcaagt
 gctgtgacgagggcacattggaggcttcccagacatagccacgagaagagctgggaccagcactggctctgggcgagcgatgagagcta
 cggcgctacctaaggagcatcgcggtgatgctggcgcttcgactacgtcaaggctacggagcgtgggtcgtcaaggactggctggact
 ggtggggaggctgggcccgggagtgactgggacacaaacgttgatgactgctcaactgggctactcagcgatgcaaaagtcttcgac
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 cgcgacccgttaaggccgtaaccttcgttcaaaaccacgacaccgatataatctggaacaagtatccagcctacgcgttcacctcaccacga
 ggccagccgacaatattctacgcgactacgaggagtggtctcaacaaggataagctcaagaacctcatctggatcatgacaacctcgccg
 gaggagacgagcatagttactacgacagcgacgagatgatcttcgtgaggaaaggctatggaagcaagcctggcctataacttacaacaa
 cctcggctcagcaaggttgaagggtgggttacgttcgaagttcgcaggctcgtgcatacacgagtacaccggcaatctcggcggtgggt
 ggacaagtgggtgactcaagcggtgggttacctcagggtcctgcccacgacccggccaacggccagtagcggctactcctgctggagc
 tactgcggtgttggtga

SEQ ID NO.: 18

Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Glu Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
 Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
 Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
 Val Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
 Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
 Ala Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
 Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
 Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Ser Ile Val

FIGURE 16G

Tyr Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His
 Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu
 Ala Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 19

atggccaagtacctggagctcgaagagggcgggctcataatgcaggccttctactgggacgtcccatgggaggaatctgggtgggacacgat
 agcccaagaatacccgactgggcaagcggcggtatcggcgatatggattcctcccgagcaagggtatgagcggcggtattcggatgg
 gctacgacccctacgattatgttgacctgggtgagtactaccagaagggaacgggtggaacgaggttcggctcaaagcaggagctcataaacat
 gataaacacggcccatgcctacggcataaaggctacagcgacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccgttc
 gttggggactacacctggacggacttctcaaagggtggcctcgggcaataatactgccaactacctgacttccaccgaacgagctccatgcg
 ggcgattccggaacattggaggctatcccgacatatgccagacaagagctgggaccagtactggctctgggccagccaggagagctacgc
 ggcatatctcaggagcatcggcatcgtatcctggcgcttcgactactgcaagggtatgctccctgggtcgtcaaggactggctgaactgggtg
 gggggctggggcggttgagagtactgggacaccaacgtcgactgttctcaactgggcatactcgagcgggtgccaaggctttgacttgcgc
 ctctactacaagatggatgagggcctttgacaacaaaaacattccagcgctcgtctgctccctcagaacggccagactgtgtctccgcgaccc
 gtcaaggccgtaacctttgtagcaaacacgacaccgatataattggaacaagtaaccggcctacgcttcatctcactacgagggccag
 ccgacgatattctaccgcgactacgaggagtggctcaacaaggacaggctcaagaacctcatctggatacacgaccacctcgccggtggaag
 cactgacatcgtttactacgacaacgacgagctgatattcgtgagaacggctacggaagcaagccgggactgataacatacatcaacctgc
 ctcaagcaaaagccggaagggtgggttatgtgccgaagttcggggcgcggtgcacccagcagcacttggttaacctcgagggtggttagaca
 agtacgtctactcaagcggctgggtctatctcgaagctccagcttacgacctgccaacgggcagtatggctactccgtgtggagctactgcgg
 tgttggtga

SEQ ID NO.: 20

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Glu Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr
 Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu His
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 21

atggccaagtactccgagctggaagagggcggcggtataatgcaggccttctactgggacgtcccagggtggaggaatctgggtgggacacat
 caggagcaagataccggagtggtacgaggcgggaataccgccatttgattcctcccgaggcaagggtatgagcggcggtattcggatgg
 gctacgacccctacgatattggacctgggtgagtactaccagaagggaacgggtggaacgaggttcggctcaaagcaggagctcataaac
 atgataaacacggcccatgcctacggcataaaggctacagcgacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccgtt
 cgttggggactacacctggacggacttctcaaagggtggcctcgggcaaatatactgccazctacctgacttccaccgaacgagctccatgc
 ggcgatccggaacattggaggctatcccgacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctac

FIGURE 16H

gcgggtatatctcaggagcatcgccatcgatgcctggcgcttcgactacgtcaagggctacggagcgtgggtcgtcaaggactggctggactg
gtggggagggtcgccgctcggggagtgactgggacacaaacgttgatgcactgtcaactgggcctactcgagcgtgcaaaagtcttcgactt
cccgtctactacaagatggatgaggccttgacaacaaaaacattccagcgtcgtctctgcccttcagaacggccagactgtgtctcccgcg
acccgtcaaggccgtaacctttgtagcaaacacgacaccgatataatttgaacaagtacccggcctacgccttcacctcacctacgaggg
ccagccgacgatattctaccgactacgaggagtggtcaacaaggacaggctcaagaacctcatctggatacacgactacctcgccggtg
gaagcactgacatcggttactacgacaacgacgagctgattctgtgaaacggctacggagcaagccgggactgataacatacatcaacc
tcgctcaagcaagccggaggtggtttatgtccgaagttcggggcgctgcatccacgagtatactgtaacctcggaggctgggtg
acaagtacgtctactcaagcggcgtgggtctatctcgaagctccagcttacgccctgccaacgggcagtatggctactccgtgtggagctattgc
gggtgtggctga

SEQ ID NO.: 22

Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Gly Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Asp Leu Asp
Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Val
Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp Tyr Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 23

atggccaagtactccgagctggaagagggcggttatagtgcaggcccttactgggacgtcccagggtggaggaatctgggtggacaccat
caggagcaagataccggagtgtgtagggcggaataccgccatttggattccccggcgagcaagggcatggcgccgctattcgtg
ggctacgaccctacgacttcttgacctcggtagtagtaccaggaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
catgataaacacggcctatgctttagggcctaaaggctatgcggacatgcataaacctcggcagggtggagacctggagtggaacccg
ttcgtgggggactaacctggagccttctaaagggtgctcggcgaatatactgcaactacctcgaactcccccgaagagctatg
cgggcgattccggaaacttggagggtatccgaatatgacacgacaagagctgggaccagtactgggtcgtggccagccaggagagctac
gcggcatatctcaggagcatcggcatcgtgcctggcgcttcgactacgtcaagggtacggagcgtgggtcgtcaaggactggctggactg
gtggggagggtggcgctcggggagtactgggacacaaacgttgatgcactgtcactgggcctactcgagcgtgcaaaagtcttcgactt
cccgtctactacaagatggatgaggccttgacaacaaaaacattccagcgtcgtctctgcccttcagaacggccagactgtgtctcccgcg
acccgtcaaggccgtaacctttgtagcaaacacgacaccgatataatctggaacaagtatccagcctacgcgttcacctcacctacgaggg
ccagccgacaaatattctaccgactacgaggagtggtcaacaaggataagctcaagaacctcatctggatacatgacaacctcggcggag
gaagcatgagcatattctacgacagcgacgagatgcttcgtgaggaaaggctatggaaagcaagcctggccttatactacatcaacctc
ggctcgagcaaggttggaaagggtgtctacgtccgaagttcggggagcgtgcatccacgagtacaccggcaacctcggcggtgggtg
acaagtgggtgactcaagcgggtgggtgtacctcaggcccttggccacgacccggccaacggctattacggctactccgtctggagctatt
gcggtgttggctga

SEQ ID NO.: 24

FIGURE 16I

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Val Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Met Ser Ile Val Tyr
 Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 25

atggccaagtacctggagctcgaagagggcggtcataatgcaggccttctactgggacgtcccatgggaggaatctggtgggacacgat
 agcccaagaatacccgactgggcaagcgccgggatttcggcgatatggattcctcccgagcaagggtatgagcggcggtattcgatgg
 gctacgaccctacgattatttgacctcgggtgagtactaccagaagggaacgggtggaacgaggttcggctcaagcaggagctcataaacat
 gataaacaccgcccacgctatggcatgaaggaatagccgatatgcatcaaccaccgcccggcggtgacctggagtggaaccccttcgt
 gaacgactatacctggacgacttctcaagggtcgctcgggtgtaatacacggccaactacctcgacttccaccgaacgagctccatgcggg
 cgattccggaacatttgaggctatcccgacatagccacgacaagagctgggaccagtacggctctgggcccagcaggagagctacgcgg
 catatctcaggagcagcggcatcgatgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactggtggg
 aggctggcggttgagagtagtgggacaccaacgtcgacgtgttctcaactgggatactcgagcgggtgccaaggcttctgacttcgccctc
 tactacaagatggagcaggccttcgataaacaacattccgcccttggtgggcccctcagatacggtcagacagtggtcagccgcgaccc
 gttaaggctgtgacgtttgtagccaaccaacgataccgataatactggaacaagatccagcctacgcgttcacctcacctacgaggccagc
 cgacaatttctaccgcgactacgaggagtggctcaacaaggataagctcaagaacctcatctggatacatgacaacctgccggaggaagc
 accgacatagctactacgataacgatactatcttcgtaggcacggctacggggacaagccgggcttataacctacatcaacctaggt
 cgagcaaggccggaagggtgggtttacgttcgaagttcgcaggctcgtgcatacacgagtagccggcaatctcggcggttggtgggacaa
 gtgggtgggacaaagcgggtgggtctacctgaggtcctgccacgacccggccaacggcagtagcgtactcgtctggagctattgctg
 gttgggtga

SEQ ID NO.: 26

Met Ala Lys Tyr Leu Glu Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Gly Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp

FIGURE 16J

Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr
 Asp Asn Asp Glu Leu Ile Phe Val Arg His Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 27

atggcaagattatccgagctcgaagagggcggtataatgcaggccttctactgggacgtcccaggtggaggaatctggtgggacaccatc
 aggaagcaagataccggagtggtacgagcgagggaatatccgccatttgattctcccgagcaagggtatgagcgcggtattcgatggg
 ctacgacccctacgattattttgacctcggtgagtactaccagaagggaacggtggaaacgaggttcggctcaaagcaggagctcataaacatg
 ataaacacggcccatgcctacggcataaaggctacagcgacatcgtcataaaccaccgcgcaggcgagacctcgagtgaacccgttcgt
 tggggactacacctggacgggacttctcaagggtggcctcgggcaaatatactgccaactacctgacttcaccccgaacgagctccatcgggg
 cgattccggaacatttgaggctatcccgacatatgccacgacaagagctgggaccagtagctggctcggccagccaggagagctacggg
 catatctcaggagcatcggcacgatgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtaaggactggctgaactgggtggg
 aggctgggcggttgagagtagctgggacaccaacgtcgacgctgttctcaactgggcatactcgagcggtgccaaaggctttgacttcgccc
 tactacaagatggagcgcgcccttgacaacaagaacattcccgcactcgtcgaggccctcaagaacgggggcacagtcgtcagccgcgacc
 cgtttaaggccgtaaccttcgttgcaaacacgacaccgatataatctggaacaaglatccagcctacgcgttcacccacccacgagggccag
 ccgacaatattctacgcgactacgaggagtggtcaacaaggataagctcaagaacctcatctggatacatgacaacctgcgggaggaag
 cactgacatcggttactacgacaacgacgagctgatattctgagaaacggctacggaagcaagccgggactgataacatacatcaacctgc
 gtaagcaagccggaagggtgggttacgttcgaagtcgaggtcgtgcatacagagtagaccggcaatctcggcggctgggtggaca
 agtgggtggactcaagcggtgggtctacctgaggtcctcgccacgacccggccaacggccagtagggctactcgtctggagctactgc
 ggtgtgggtga

SEQ ID NO.: 28

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Ala Ala
 Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr
 Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 29

atggccaagtagctcgaagagggcggtcataatgcaggccttctactgggacgtcccaggtggaggaatctggtgggacacggg
 agcccaagatacccgactgggcaagcgccgggatttcggcgatatggattccccggcgagcaaggcgatggcgccgctattcgatg
 ggctacgzcctacgacttcctgacctcggtgagtacgaccagaagggaacgggtagagacgcgcttggctccaagcaggagctcgtgaa

FIGURE 16K

catgataaacacggcccatgcctacggcataaagggtcatagcggacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccg
 ttctgttggggactacacctggacggacttctcaaagggtggtctcgggcaaatatactgccaactacctcgacttccaccgaacgagctccatgc
 gggcgattccgggaacatttggaggctatcccgacatatgccacgacaagagctgggaccagtagctgtctgggcccagccaggagagctac
 gcggcatatctcaggagcatcggcatcgtgcctggcgcttcgactacgtcaaggctatgtccctgggtcgtcaaggactggctgaactggt
 ggggaggctggcggttggagagtactgggacaccaacgtcgacgtgttctcaactgggcatactcagcgggtccaaggctttgacttcg
 ccctctactacaagatggatgaggcccttgacaacaaaacattccagcgtcgtcgtcgtccctcagaacggccagactgttgcctcccgac
 ccgttcaaggccgttaacctttagcaaacacgacaccgataatctggaacaaagtaccttgccttcatcctcactacgaaggccag
 cccgtcatattctaccgcgactacgaggagtgggtcaacaaggacaggttgaacaacctcatatggatacacgaccacctcgcagggggaag
 caccgacatagctactacgataacgatgaactcatcttcgaggaacggctacggggacaagccggggctataacctacatcaacctaggc
 tcgagcaaggccgggaagggtgggttatgtgccgaagtcgcggcggtgcatccacgagtatactggtaacctcggaggctgggtagacaa
 gtactgtactcaagcggctgggtctatctgaagctccagcttacgacctgccaacgggcagtagtggctactccgttggaagtactcgggt
 gttgggtga

SEQ ID NO.: 30

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met
 Gly Gly Ile Trp Trp Asp Thr Val Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Val Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 31

atggcaagactcagagctggaagaggggcggtataatgcaggcttctactcggacgtccaggtgaggaatctggtgggacacat
 caggagcaggaacatttggaggctatcccgacatatgccacgacaagagctgggaccagtagctgtctgggcccagccaggagagctac
 ggctacgacccctcagacttcttgactcgggtgagtagcagcagaagggaacggtagagacgcgtttggtcccaagcaggagctcgtgaa
 catgataaacacggcccatgcctacggcataaagggtcatagcggacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccg
 ttctgttggggactacacctggacggacttctcaaagggtggtcgggcaaatatactgccaactacctcgacttccaccgaacgagctccatg
 cggcgattccgggaacatttggaggctatcccgacatatgccacgacaagagctgggaccagtagctgtctgggcccagccaggagagctac
 gcggcatatctcaggagcatcggcatcgtgcctggcgcttgactacgtgaagggtacggagcgtgggtcgtcaaggactggctcaactgg
 tggggcggtggcggttggcgagtactgggacaccaacgttgatgcactcctcaactgggcctactcagcggcgccaaggctcttcgacttc
 ccgctctactacaagatggacgaggccttcgataacaacaacattcccgccctgggtggacgcctcagatacggtcagacagtggctagccgc
 gacccgttcaaggctgtgacgttttagccaaccacgataccgatataatctggaaacagatccagcctacgcgttcctcactacgaggg
 ccagccgacaatttctaccgcgactacgaggagtggctcaacaaggataagctcaagaacctcatctggatacatgacaacctggccggag
 gaagcacgagcatagtttactacgacagcagagatgcttctgtaggaccggctatggaagcaagcctggccttataacttacatcaacct
 cgggtcagcaagggttgaagggtgggttatgtgccgaagtcgcggcggtgcatccacgagtatactggtaacctcggaggctgggtaga
 caagtactgtactcaagcggctgggtctatctcgaagctccagcttacgacctgccaacgggcagtagtggctactcgggtggagctatggcg
 gttgggtga

FIGURE 16L

SEQ ID NO.: 32

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Arg Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr
 Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Thr Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 33

atggccaagtactccgagctggaagagggcggggtcataatgcaggcgttctactgggacgtgccttcaggaggaatatgggtggacacaat
 acggcagaagataccggagtggtacgatgccggaatciccgaatatggattctctcccgagcaagggtatgagcggcggctattc gatgg
 gctacgacctacgattatttgacctcggtagtactaccagaagggaacgggtggaacgaggttcggctcaaacgaggagctcataaacat
 gataaacacggcccatgctacggcataaaggctcatagcggacatcgtcataaaccaccgcgcaggcggagacctcaggtggaaccggtc
 gttggggactacacctggacggacttctcaagggtgacctcgggcaataatactgccactacctcgaactccaccggaacgagctccatgcg
 ggcgattcgggaacatttgagggtatcccgacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctacgc
 ggcatatctcaggagcatcggcatcgtatgcttggcgctttgactactgtgaagggtacggagcgtgggtcgttaaggactggctcaactgggtg
 gggcggtctgggcccgttgaggactgggacaccaacgttgatgcactcctcaactgggctactcagcggcgccaaagggtcttcgacttcc
 gctctactacaagatggacgcggcctttgacaacaagaacattcccgactcgtcaggccctcaagaacggggggcacagtcgtcagccgcg
 acccggttaaggccgtaaccttcgttgcaaacacgacaccgataatactggaccaagtaaccttgcttatgcttctacctcactacgaaggcca
 gcccgtcatattctaccgcgactacgaggagtggtcacaacaggacagggtgaacaacctcatatggatacacgaccacctcgcagggtgaag
 caccgacatagtctactacgataacgatgaactcatcttcgtcaggaaacggctacggggacaagccggggcgtataacctacatcaaccttaggc
 tcgagcaaggccggaaggtgggtttacgttcgaagtcgaaggtcgtgataacacgagtaacacgggcaatcaggcggtctgggtggacaa
 ctgggtggtactcaagcggttgggttctcgtgaggtctgtgcacgacccggccaacggcagtaagggtactcogtctgggtgactggc
 ggttggcga

SEQ ID NO.: 34

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala

FIGURE 16M

Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Thr Lys Tyr Leu Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 35

atggccaagtactcggagctggaagaggcggttataatgcaggccttctactgggacgtccagggtgaggaatctggtgggacacccat
 caggagcaagataaccggagtggtacgagcgagggaataatccgccatttgattccccggcgagcaagggcattggcggcctattcgatg
 ggctacgacccctacgactctttgacctcggtgagtagcaccagaagggaacggtagagacgcgtttggctecaagcaggagctcgtgaa
 catgataaacaccgcccacgcctacggcatcaaggctcgcagacatagtaataaccaccgcgcggaggagaccttgagtgaacccct
 tctcaatgactacacctggacggacttctgaaggctcgttcggcaagtacacggccaactacctgacttccaccccaacgaggtcaagt
 ctgtgacgagggcacatttgaggcttccagacatagcccacgagaagagctgggaccagcactggctctggcgagcgatgagagctac
 gccgcctacctaaggagcatcggcgttgatgcctggcgctcgcactacgtcaagggtatgtccctgggtcgaaggactggctgaactggt
 ggggaggctggcggttgagagtagtactgggacaccaacgtcgcagctgttctcaactgggcatactcagcgggtgccaaggctttgacttcg
 ccttactacaagatggacgcggccttgacaacaagaacattcccgcactcgtcagggccctcaagaacgggggcacagctcgcagccge
 gaccggttaaggccgtaaccttctgtgcaaacacgacaccgataataatctggaacaagtatccagcctacgcgttcacctcacctacgagg
 ccagccgacaatatctaccgcgactacgaggagtggtcaacaaggataagctcaagaacctcatctggatacatgacaacgtcgcggag
 gaagcaccgacatagctactacgataacgatgaactcatcttcgcaggaaacggctacggggacaagccggggctataacctacatcaacct
 aggtcgcagcaaggccggaagggtgggtttacgttcgaagttcgcaggctcgtgcatacacgagtagaccggcaatctcggcggtgggtgg
 acaagtgggtggactcaagcggctgggtctacctcagggtcctgccacgacccggccaacggccagtaggctactcgtctggagctac
 tgcgggttggtga

SEQ ID NO.: 36

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
 Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Gln Ser Tyr
 Ala Ala Tyr Leu Asp Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp
 Val Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
 Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp
 Ala Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
 Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
 Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Val Ala Gly Gly Ser Thr Asp Ile Val
 Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His
 Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu
 Ala Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 71

atggccaagtacctggagctcgaaggaggcggttcataatgcaggcgttctactgggacgtccagggtgaggaatctggtgggacacccat

FIGURE 16N

acggcagaagataccggagtggtacgatgccggaatctccgcaatatggattccccggcgagcaaggcagtgggcgccgctattcga
 ggctacgacccctacgacttctttgacctcggtagtacgaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
 catgataaacacggcccatgcctacggcataaagggtacatagcggacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccg
 ttcgttggggactacacctggacggacttctcaaaaggtagcctcgggcgaatatactgccaaactacctcgaactccacccgaacgagctccatg
 cgggcgattccggaacatttggaggctatcccacatagccacgacaagagctgggaccagtactggctcgtggccagccaggagagctac
 gcggcatactcaggagcatcggcatcgaatgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactggt
 ggggaggtcggcggttggagagtactgggacccaacgtcgcagctgttctcaactgggcatactcagcgggtgccaaggctttgacttcg
 cccctactacaagatggatgaggccittgacaacaaaacattccagcgcctcgtctcgtccctcagaacggccaactgttgcctcccgac
 ccgttcaaggccgtaaccttttagcaaacacgacacccgatataatctggaacaagtatccagctacgcgttcatctcactacgaggggcc
 agccgacaatattctaccggaactacgaggagtggctcaacaaggataagctcaagaacctcatctggatacatgacaacctcggcgaggga
 agcactgacatcgtttactacgacaacgacgagctgatatctgagaacggctacggaagcagccgggactgataacacacacacacac
 gcccaagcaaaaggcggaagggtgggttatgtgccgaagttcggggcggtgcatccacgagtatactggtaacctcggaggctgggtaga
 caagtacgtctactcaaggcggtgggtctatctcgaagctccagcttacgacctgccaacgggcagtagtgcctactcgtgtggagctactgc
 ggggtgggctga

SEQ ID NO.: 72

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
 Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
 Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
 Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
 Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 49

gtggttatgacgatgtccgctatgacattatgcctaggaatggcggtgtttatcgtttcacgagctcctgttggagccaaagcgcgtctct
 accgttcccttctggtcgtactcaccgaggtcaaaagaagtogtaggggtcgtagcccatcgaataggcgcgccccatgcccttgcctcggg
 ggaatccatcgcggaaatccccggcgcttcccagtcgggtatcttctgggctatcgtgtcccaccagattcctccatggggacgtcccagta
 gaaggcctgcatatgagccccctcttcgagccccgaatacttggcataagttacctctactagtagaataaaattctgttctgtgtgaaatt
 gtt

SEQ ID NO.: 50

Val Val Tyr Asp Asp Val Arg Tyr Asp Leu Tyr Ala Val Gly Met Gly Arg Val Tyr His Val His Glu
 Leu Leu Leu Gly Ala Lys Ala Arg Leu Tyr Arg Ser Leu Leu Val Val Leu Thr Glu Val Lys Glu Val
 Val Gly Val Val Ala His Arg Ile Gly Ala Ala His Ala Leu Ala Arg Arg Gly Asn Pro Tyr Arg Arg
 Asn Pro Gly Ala Cys Pro Val Gly Tyr Leu Leu Gly Tyr Arg Val Pro Pro Asp Ser Ser His Gly Asp
 Val Pro Val Glu Gly Leu His Tyr Glu Pro Ala Leu Phe Glu Pro Gly Ile Leu Cys His Lys Leu Pro
 Pro Thr Ser Arg Leu Lys Phe Cys Phe Leu Cys Glu Ile Val

FIGURE 160

SEQ ID NO.: 51

ATGGCCAAGTACCTGGAGCTCGAAGAGGGCGGGGTCATAATGCAGGCGTTCTACTGGG
 ACGTGCCTTCAGGAGGAATATGGTGGGACACAATACGGCAGAAGATACCGGAGTGGT
 ACGATGCCGGAATCTCCGCAATATGGATTCCCCGGCGAGCAAGGGCATGGGCGGCGC
 CTATTCGATGGGCTACGACCCCTACGACTTCTTTGACCTCGGTGAGTACGACCAGAAG
 GGAACGGTAGAGACGCGCTTTGGCTCCAAGCAGGAGCTCGTGAACATGATAAACACC
 GCCCACGCCTATGGCATGAAGGTAATAGCCGATATAGTCATCAACCACCGCGCCGGCG
 GTGACCTGGAGTGGAACCCCTTCGTGAACGACTATACCTGGACCGACTTCTCAAAGGT
 CGCGTCGGGTAAATACACGGCCAACCTACCTCGACTTCCACCCCAACGAGGTCAAGTGC
 TGTGACGAGGGCACATTTGGAGGCTTCCAGACATAGCCACGAGAAGAGCTGGGAC
 CAGCACTGGCTCTGGGCGAGCGATGAGAGCTACGCCGCTACCTAAGGAGCATCGGCG
 TTGATGCCTGGCGCTTTGACTACGTGAAGGGCTACGGAGCGTGGGTCTGCAAGGACTG
 GCTCAACTGGTGGGGCGGCTGGGCCGTTGGCGAGTACTGGGACACCAACGTTGATGCA
 CTCCTCAACTGGGCCTACTCGAGCGGCGCCAAGGTCTTCGACTTCCCGCTCTACTACAA
 GATGGATGAGGCCTTTGACAACAAAAACATTCCAGCGCTCGTCTCTGCCCTTCAGAAC
 GGCCAGACTGTTGTCTCCCGCGACCCGTTCAAGGCCGTAACCTTTGTAGCAAACCACG
 ACACCGATATAATCTGGAACAAGTATCCAGCCTACGCGTTCATCCTACCTACGAGGG
 CCAGCCGACAATATTCTACCGCGACTACGAGGAGTGGCTCAACAAGGATAAGCTCAAG
 AACCTCATCTGGATACATGACAACCTCGCCGGAGGAAGCACTGACATCGTTTACTACG
 ACAACGACGAGCTGATATTCGTGAGAAACGGCTACGGAAGCAAGCCGGGACTGATAA
 CATAATCAACCTCGCCTCAAGCAAAGCCGGAAGGTGGGTTTACGTTCCGAAGTTCGC
 AGGCTCGTGCATACACGAGTACACCGGCAATCTCGGCGGCTGGGTGGACAAGTGGGTG
 GACTCAAGCGGCTGGGTCTACCTCGAGGCTCCTGCCACGACCCGGCCAACGGCCAGT
 ACGGCTACTCCGTCTGGAGCTATTGCGGTGTTGGCTGA

SEQ ID NO.: 52

MAKYLELEEGVIMQAFYWDVPSGGIWWDTIRQKIPEWYDAGISAIWIPPASKGMGGAYS
 MGYDPYDFDLGEYDQKGTVETRFSGKQELVNMINTAHAYGMKVIADIVINHRAAGDLE
 WNPFFVNDYTWDTSKVASGKYTANYLDFHPNEVKCCDEGTFGGFPDIAHEKSWDQHWL
 WASDESYAAYLRSIGVDAWRFDYVKGYGAWVVKDWLNWWGGWAVGEYWDTNVDAL
 LNWAYSSGAKVDFPLYYKMDEAFDNKNIPALVSALQNGQTVVSRDPFKA VTFVANHDT
 DIFWNKYPAYAFILTYEGQPTIFYRDYEEWLNKDKLKNLIWIHDNLAGGSTDIVYYDNDELI
 FVRNGYGSKPLITYINLASSKAGRWWYVPKFAGSCIHEYTGNLGGWVDKWVDSSGWVY
 LEAPAHDPANGQYGYSVWSYCGVG

SEQ ID NO.: 37

atggcgaagtacctggagctcgaagagggggggtcataatgcaggcggttctactgggacgtgccttcaggaggaatatgtgtggacacaaat
 acggcagaagataccggagtggtacgatccggaatctccgcaatattgattccccggcgagcaaggcgatggcgggcgctattcgaig
 ggctacgacccctacgacttcttgacctcgggtgagtagcaccagaagggaacggtagagacgcgctttggctccaagcaggagctcgtgaa
 catgataaacacggccacgcctatggcatgaagtaataagccgatatagtcatcaaccaccgcgccggcggtgacctggagtggaacccctt
 cgtgaacgactatacctggaccgacttcaaaaggctcgctcgggtaataacacggccaactacctcgaacttccacccgaacgagctccatgc
 gggcgattccggaaacatttgaggctatcccgacatatgccacgacaagagctgggaccagtactggctcggccagcgaaggagagctac
 gcggcatatctcaggagcctcggcatcgatgcctggcgctttgactacgtgaagggtacggagcgcggtgtcgaaggactggctcaactg
 gtggggcggtcggcgcttggtgcgagtactgggacaccaacgttgatgcactcctcaactgggctactcgaaggcgccaaggcttctcgactt
 cccgctctactacaagatggatgagcgctttgacaacaaaacattccagcgctcgtctctgcccttcagaacggccagactgtgtcctccgcg
 acccgctcaaggccgtaaccttttagcaaacacgacacccgatataatctggaacaagatccagcctacgcgttcaactcctacacagagg
 ccagccgacaataattctatcgcgactacgaggagtggtcacaagaagataagctcaagaacctcatctggatacatgacaacctcggcgagg
 aagcactgacatcgttactacgacaacgacgagctgatatcgtgaagaacggctacgggaagcaaggcggaactgataacatcatcaacct
 cgcttcgaagcaagccgggaagggtgggtttacgttcgaaggttcgaggctcgtgcatacacgagttacacccggcaatctcggcggtcgggtgg

FIGURE 16P

acaagtgggtggactcaagcggctgggtctacctcgaggctcctgccacgacccggccaacggccagtagcggtactccgtctggagctac
tgcgggggtgggtga

SEQ ID NO.: 38

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly
Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala
Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Arg Val
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu
Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 39

atggccaagtacctggagctcgaagagggcggggtcataatgcaggcgttctactgggacgtgccttcaggaggaatatggtgggacacaat
acggcagaagataccggagtgtacgatccggaatcctccgaatatggattcctcccgagcaggggatgagcggcggtattcgtatgg
gctacgacctacgattatattgacctcgggtgagtactaccagaagggaacgggtggaacgaggttcggctcaagcaggagctcataacat
gataaacaccgcccacgctatggcatgaagtaataagccgataatgcatcaaccaccgcgccggcggtgacctggagtggaaccccttctg
gaacgactatacctggaccgacttctcaaggtgcgtcgggtaaatacacggccaactacctgacttccaccgaacgagctccatgcggg
cgattccggaacatttggaggctatcccgacatagccacgacaaagagctgggaccagtactggctctgggccagccaggagagctacgcgg
catactcaggagcatcggtatcgatgcctggcgcttgactactgtaagggtacggagcgtgggtcgtcaaggactggctcaactgggtggg
gcggctggggcgttgccgagtactgggacccaacgttgatgcccctcctccctgggctactcgagcgcccaaggcttctgacttccgc
tctactacaagatgatgaggcctttgacaacaaaacattccagcgctcgtctcgtccctcagaacggccagactgtgtctcccgagcccg
ttcaaggccgtaacctttgtagccaaccacgataccgataatactggaacaagatccagccctacgcgttcatctcacctacgagggccagcc
gacaatttctaccgcgactacgaggagtggctcaacagagataagctcaagaactctctggataaagcaacctcggcggaaggagca
ccgacatgtctactacgataaagaggaactcatttggcaggaaggctacggggaacggcggttataactatcatcaactagagtc
gagcaaggccggaagggtgggtctacgttcgaggttggggagcgtgcatccacgagtaacgggaacctcggcggttgggtggacaa
gtgggtggactcaagcgggtgggtgtacctcgaggccctgccacgacccggccaacggctattacggctactcgtctggagctactgcg
gggtgggtga

SEQ ID NO.: 40

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Arg Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala

FIGURE 16Q

Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Pro Asn Val Asp Ala Leu
 Leu Pro Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr
 Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu Tyr
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 41

atggccaagtacgtgagctcgaagagggcggggtcataatgcaggcggttactgggacgtgccttcaggaggaaatggtgggacacaat
 acggcagaagataccggagtggtacgatgccgaatctccgcaatatggattctcccgcgagcaagggtatgagcggcggtattcgatgg
 gctacgaccctacgattatttgacctcggtagtactaccagaagggaacgggtggaacgaggttcggctcaaagcaggagctcataaacat
 gataaacacggccatgcctacggcataaaggctacagcgacatcgtcataaaccaccgcgcaggcggagacctcagtggaacccgttc
 gttggggactacacctggacggacttctcaaagggtggcctcgggcaaatatactgccaactacctgacttccaccggaacgagctccatgcg
 ggcgattccggaacatttgaggctatcccgacatagccatgacaagagctgggaccagtactggctctggccagccaggagagctacgc
 ggcatatctcaggagcatcggcatcgtgcttggcgttgactactgtaagggtacggagcgtgggtcgtcaaggactggctcaactgggtg
 ggccggctgggcccgttgaggtagtactgggacaccaacgttgatgcactctcaactgggctactcagcggcgccaagggtcttcgacttccc
 gctctactacaagatggacgcggccttgacaacaagaacatctccgcactcgtcaggccctcaagaacgggggcacagctcgtcagccgcg
 acccgittaaggccgttaacctcgttgcaaacacgacaccgataaatctggaacaagtatccagcctacgcgttcactctacctacgagggc
 cagccgacaatattctaccgcgactacgaggagtggtcacaagaaggataagctcaagaacctcatctggatacatgacaacctcgccggagg
 aagcacgagcatagtttactacgacagcgacgagatgatctctgtgaggaacggctatggaaagcaagcctggccttataactacatcaacctc
 ggctcagcaagggtggaagggtggttatgtgccgaagtgcgggcgctgcatccacgagtatactggtaacctcgagggtgggtagac
 aagtacgtctactcaagcggctgggtctatctcgaagctccagcttaccacctgccaacgggcagtagtggtactccgtgtggagctactgcg
 gtgtgggtga

SEQ ID NO.: 42

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr
 Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

FIGURE 16R

SEQ ID NO.: 43

atggccaagtactccgagctggaagagggcggcgttataatgcaggccttctactgggacgtccaggtggaggaaatctggtgggacacat
caggagcaagataccggagtggtacgagggcgggaatatccgccatttgattccccggcgagcaaggcatggcgggcgctattcga
ggctacgaccttacgacttctttgacctcggtagtacgaccagaagggaacggtagagacgcgtttggctccaagcaggagctcgtgaa
catgataaacacggcccatgcctacggcataaaggatagcggacatcgtcataaaccaccgcgagggcgagacctcgaagtgaacccg
ttcgttggggactacacctggacggacttctcaaagggtggcctcgggcaaatatactccaactacctcgaactccaccccaacgaggtcaagt
gctgtgacgagggcacatttggaggcttccagacatagcccacgagaagagctgggaccagcactggctctggggcagcgatgagagcta
cgccgcctacctaaggagcatcggcgttgatgcctggcgcttcgactacgtcaagggtacggagcgtgggtcgtcaaggactggctggact
gggtggggaggtcggcgctcggggagtagtgggacacaaacgttgatgactgctcaactggcctactcgaagcgtgcaaaaagtcttcgac
ttcccgtctactacaagatggatgagggcctttgacaacaaaacattccagcgctcgtctcgtcccttcagaacggccagactgtgtctccgc
gacctgttaaggccgttaaccttttagcaaacacgacaccgataatactggaacaagtatccagcctacgcgttcatctcacctacgagg
gccagccgacaatatctaccgcgactacgaggagtggtcaacaaggataagctcaagaacctatctggatacatgacaacctcgtcggag
gaagcacgagcatagttactacgacagcgacgagatgattcgtgaggaacggctatggaagcaagcctggccttataactacatcaacct
cggctcagagcaaggttgaagggtgggttacgtccgaagttcgcaggctcgtgcatacacgaglacaccggcaatctcggcggtgggtgg
acaagtggtggactcaagcggctgggtctacctcagggtcctgccacgacccggccaacggccagtagcgtactccgctcgtgagctac
tgcgggtgttggtga

SEQ ID NO.: 44

Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
Val Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
Glu Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser
Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Val Gly Gly Ser Thr Ser Ile Val
Tyr Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His
Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu
Ala Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 45

atggccaagtactccgacctggaagagggcggcgttataatgcaggccttctactgggacgtccaggtggaggaaatctggtgggacacat
caggagcaagataccggagtggtacgagggcgggaatatccgccatttgattccccggcgagcaaggcatggcgggcgctattcga
ggctacgaccttacgacttctttgacctcggtagtacgaccagaagggaacggtagagacgcgtttggctccaagcaggagctcgtgaa
catgataaacacggcccatgcctacggcataaaggatagcggacatcgtcataaaccaccgcgagggcgagacctcgaagtgaacccg
ttcgttggggactacacctggacggacttctcaaagggtggcctcgggcaaatatactccaactacctcgaactccaccccaacgaggtcaagt
gctgtgacgagggcacatttggaggcttccagacatagcccacgagaagagctgggaccagcactggctctggggcagcgatgagagcta
cgccgcctacctaaaggagcatcggcgttgatgcctggcgctttgactacgtgaagggtacggagcgtgggtcgtcaaggactggctcaactg
gtggggcggtcggcggttggcgagtactgggacaccaacgttgatgactcctcaactggcctactcgaagcggcgcccaagggtcttcgact
cccgtctactacaagatggatgagggcctttgacaacaaaacattccagcgctcgtctcgtcccttcagaacggccagactgttgtcctccgc
acctgttaaggccgttaaccttttagcaaacacgacaccgataatactggaacaagtatccagcctacgcgttcatctcacctacgagg
ccagccgacaatatctaccgcgactacgaggagtggtcacaagaagataagctcaagaacctcaatcggatcatgacaacctcggcggag

FIGURE 16S

gaagcaccgacatagtctactacgataacgatgaactcatcttcgtcaggaacggctacggggacaagccggggcttataacctacatcaacct
aggctcgagcaaggccggaagggtgggttatgtgccgaagttcgccggcgctgcatccacgagtatacttgtaacctcggaggctgggtag
acaagtacgtctactcaagcggtgggtctatctcgaagctccagcttacgacctgccaacgggcagtatggctactccgtgtggagctattgc
gggtgtgggtga

SEQ ID NO.: 46

Met Ala Lys Tyr Ser Asp Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
Val Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
Glu Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser
Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val
Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His
Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu
Ala Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 47

atggccaagtacaccgagctggaagaggcgccgttataatgcaggcccttactgggacgtcccagggtggaggaatctgggtggacacccat
caggagcaagataccggagtggtacgagggcggaatatccgccatttgattccccggcgagcaagggcagtgggcgccctattcgatg
ggctacgacccctacgactctttgacctcggtgagtagcagaccagaagggaacggtagagacgcgcttggctccaagcaggagctcgtgaa
catgataaacaccgcccacgcctatggcatgaaggtaatagccgatatagtcataaccaccgcgcggcggtgacctggagtggaacccctt
cgtgaacgactatacctggaccgactctcaaaaggctcgcgtcgggtaatacacggccaactacctcgaattccacccaacgaggtcaagtg
ctgtgacgagggcacatttgagggttccagacatagcccacgagaagagctgggaccagcactggctctgggcgagcgatgagagctac
gccgcctacctaaggagcatcgccgttgatgcctggcgctttgactacgtgaagggtcagcgagcggtggctcgaaggactggctcaactgg
tggggcggttggccggttggcgagtagtgggacaccaacggtgagcactcctcaactgggctactcagcgccgccaagggtcttcgacttc
ccgctactacagatggatgaggctttgacacaaaacattccagagctcgtctcgccttcaggaacggcagactgtttgtctcaggsa
ccggttaaggccgtaacgtttggaacacagacacagatataatgggaacaggaactgtttgtttctcctacacacgaaggcc
gccegcatacttaccgcgactacgaggatgggtcaacaaggacaggtgaacaacotatatggatacatgaccacctcgcagggtgaag
cacgagcatagtttactacgacagcgacgagatgatcttcgtgaggaacggctatggaagcaagcctggccctataacttacatcaacctcggt
cgagcaagggttgaagggtgggttacgtccgaagttcgaggcccggtgcatacacgagtagcaccggcaatctcggcggtgggtggacaag
tgggtggactcaagcggtggtctacctcgaggctcctgccacgacccggccaacggccagtagcggtactccgtctggagctactgcgg
tgttgggtgtag

SEQ ID NO.: 48

Met Ala Lys Tyr Thr Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys

FIGURE 16T

Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
 Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
 Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
 Val Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
 Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
 Glu Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu
 Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
 Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Ser Ile Val
 Tyr Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Pro Cys Ile His
 Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu
 Ala Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 53

ATGGCCAAGTACTCCGAGCTGGAAGAGGGCGGCGTTATAATGCAGGCCTTCTACTGGG
 ACGTCCAGGTGGAGGAATCTGGTGGGACACCATCAGGAGCAAGATACCGGAGTGGT
 ACGAGGCGGGAATATCCGCCATTGGATTCCCCCGGCGAGCAAGGGCATGGGCGGCG
 CCTATTCGATGGGCTACGACCCCTACGACTTCTTTGACCTCGGTGAGTACGACCAGAA
 GGAACGGTAGAGACGCGCTTTGGCTCCAAGCAGGAGCTCGTGAACATGATAAACAC
 GGCCCATGCCTACGGCATAAAGGTCATAGCGGACATCGTCATAAACCACCGCACAGGC
 GGAGACCTCGAGTGGAAACCCGTTCTGGTGGGACTACACCTGGACGGACTTCTCAAAGG
 TGGCCTCGGGCAAATATACTGCCAACTACCTCGACTTCCACCCCAACGAGGTCAAGTG
 CTGTGACGAGGGCACATTTGGAGGCTTCCAGACATAGCCACGAGAAGAGCTGGGA
 CCAGCACTGGCTCTGGGCGAGCGATGAGAGCTACGCCGCTACCTAAGGAGCATCGGC
 GTTGATGCCTGGCGCTTCGACTACGTCAAGGGCTACGGAGCGTGGGTCGTCAAGGACT
 GGCTGGACTGGTGGGAGGCTGGGCCGTCGGGGAGTACTGGGACACAAACGTTGATG
 CACTGCTCAACTGGGCCTACTCGAGCGATGCAAAAGTCTTCGACTTCCCGCTCTACTAC
 AAGATGGATGAGGCCTTTGACAACAAAAACATTCCAGCGCTCGTCTCTGCCCTTCAGA
 ACGGCCAGACTGTTGTCTCCCGCGACCCGTTCAAGGCCGTAACCTTTGTAGCAAACCA
 CGACACCGATATAATCTGGAACAAGTATCCAGCCTACGCGTTCATCTCACCTACGAG
 GGCCAGCCGACAATATTCTACCGCGACTACGAGGAGTGGCTCAACAAGGATAAGCTCA
 AGAACCTCATCTGGATACATGACAACCTCGCCGGAGGAAGCACTGACATCGTTTACTA
 CGACAACGACGAGCTGATATTCGTGAGAAACGGCTACGGAAGCAAGCCGGGACTGAT
 AACATACATCAACCTCGCCTCAAGCAAAGCCGGAAGGTGGGTCTACGTTCCGAAGTTC
 GCGGGAGCGTGATCCACGAGTACACCGCAACCTCGGCGGCTGGGTGGACAAGTGG
 GTGGACTCAAGCGGCTGGCTGTAACCTCGAGGCCCTGCCCCAGACCCGCGCAACGGCT
 ATTACGGCTACTCCGCTCTGGAGCTACTGCGGTGTTGGCTGA

SEQ ID NO.: 54

MAKYSELEEGGVIMQAFYWDVPGGGIWWDITRSKIPEWYEAGISAIWIPPASKGMGGAYS
 MGYDPYDFFDLGEYDQKGTVETFRGSKQELVNMINTAHAYGIKVIADIVINHRTGGDLEW
 NPFVGDYTWDFSKVASGKYTANYLDFHPNEVKCCDEGTFGGFPDIAHEKSWDQHWLW
 ASDESYAAYLRSIGVDAWRFDYVKGYGAWVVKDWLDWWGGWAVGEYWDTNVDALL
 NWAYSSDAKVDFPLYKMDFAFDNKNPALVSALQNGQTVVSRDPFKA VTFVANHDTD
 IWNKYPAYAFILTYEGOPTIFYRDYEEWLNKDKLKNLIWHDNLAGGSTDIVYYDNDELIF
 VRNGYGSKPGLITYINLASSKAGRWVYVPKFAGACIHEYTGNLGGWVDKWVDSSGWVY
 LEAPAHDPANGYYGYSVWSYCGVG

SEQ ID NO.: 55

FIGURE 16U

ATGGCCAAGTACCTGGAGCTCGAGGAGGGCGGGGTCATAATGCAGGCGTTCTACTGGG
 ACGTGCCTTCAGGAGGAATATGGTGGGACACAATACGGCAGAAGATACCGGAGTGGT
 ACGATGCCGGAATCTCCGCAATATGGATTCCCCCGGCGAGCAAGGGCATGGGCGGCGC
 CTATTCGATGGGCTACGACCCCTACGACTTCTTTGACCTCGGTGAGTACGACCAGAAG
 GGAACGGTAGAGACGCGCTTTGGCTCCAAGCAGGAGCTCGTGAACATGATAAACACC
 GCCCACGCCTATGGCATGAAGGTAATAGCCGATATAGTCATCAACCACCGCGCCGGCG
 GTGACCTGGAGTGGAACCCCTTCGTGAACGACTATACCTGGACCGACTTCTCAAAGGT
 CGCGTCGGGTAAATACACGGCCAACCTACCTCGACTTCCACCCGAACGAGCTCCATGCG
 GGCGATTCCGGAACATTTGGAGGCTATCCCGACATATGCCACGACAAGAGCTGGGACC
 AGTACTGGCTCTGGGCGAGCCAGGAGAGCTACGCGGCATATCTCAGGAGCATCGGCAT
 CGATGCCTGGCGCTTTGACTACGTGAAGGGCTACGGAGCGTGGGTCTCAAGGACTGG
 CTCAACTGGTGGGCGGCTGGGCGGTTGGCGAGTACTGGGACACCAACGTTGATGCAC
 TCCTCAACTGGGCCTACTCGAGCGGCGCCAAGGTCTTCGACTTCCCGCTCTACTACAAG
 ATGGATGAGGCCTTTGACAACAAAAACATTCCAGCGCTCGTCTCTGCCCTTCAGAACG
 GCCAGACTGTTGTCTCCCGCGACCCGTTCAAGGCCGTAACCTTTGTAGCAAACCACGA
 CACCGATATAATCTGGAACAAGTACCTTGCTTATGCTTTCATCCTCACCTACGAAGGCC
 AGCCCGTCATATTCTACCGCGACTACGAGGAGTGGCTCAACAAGGACAGGTTGAACAA
 CCTCATATGGATACACGACCACCTCGCAGGTGGAAGCACGAGCATAGTTTACTACGAC
 AGCGACGAGATGATCTTCGTGAGGAACGGCTATGGAAGCAAGCCTGGCCTTATAACTT
 ACATCAACCTCGGCTCGAGCAAGGTTGGAAGGTGGGTTTACGTTCCGAAGTTCGCAGG
 CTCGTGCATACACGAGTACACCGGCAATCTCGGCGGCTGGGTGGACAAGTGGGTGGAC
 TCAAGCGGCTGGGTCTACCTCGAGGCTCCTGCCACGACCCGGCCAACGGCCAGTACG
 GCTACTCCGTCTGGAGCTATTGCGGTGTTGGCTGA

SEQ ID NO.: 56

MAKYLELEEGGVIMQAFYWDVPSGGIWWDTIRQKIPEWYDAGISAIWIPPASKGMGGAYS
 MGYDPYDFDLGEYDQKGTVETRFSGSKQELVNMINTAHAYGMKVIADIVINHRAGGDLE
 WNPVFNDYTWTDFSKVASGKYTANYLDFHPNELHAGDSGTFGGYPDICHDKSWDQYWL
 WASQESYAAYLRSIGIDAWRFDYVKGYGAWVVKDWLNWWGGWAVGEYWDTNVDALL
 NWAYSSGAKVFDPLYYKMDEAFDNKNPALVSALQNGQTVVSRDPFKA VTFVANHDTD
 IWNKYLAYAFILTYEGQPVIFYRDYEEWLNKDRLNLIWTHDHLAGGSTSIVYYDSDEMIF
 VRNGYGSKPGLITYINLGSSKVGRWVYVPKFAGSCIHEYTG NLGGWVDKWVDSSGWVYL
 EAPAHDPANGQYGYSVWSYCGVG

SEQ ID NO.: 57

ATGGCCAAGTACCTGGAGCTCGAAGAGAGCGGGGTCATAATGCAGGCGTTCTACTGGG
 ACGTGCCTTCAGGAGGAATATGGTGGGACACAATACGGCAGAAGATACCGGAGTGGT
 ACGATGCCGGAATCTCCGCAATATGGATTCTCCCGGCGAGCAAGGGTATGAGCGGCGG
 CTATTCGATGGGCTACGACCCCTACGATTATTTGACCTCGGTGAGTACTACCAGAAGG
 GAACGGTGGAAACGAGGTTCCGGCTCAAAGCAGGAGCTCATAAACATGATAAACACCG
 CCCACGCCTACGGCATCAAGGTCATCGCAGACATAGTAATCAACCACCGCGCCGGAGG
 AGACCTTGAGTGGAACCCCTTCGTCAATGACTACACCTGGACCGGACTTCTCGAAGGTC
 GCTTCCGGCAAGTACACGGCCAACCTACCTCGACTTCCACCCCAACGAGGTCAAGTGCT
 GTGACGAGGGCACATTTGGAGGCTTCCAGACATAGCCCACGAGAAGAGCTGGGACC
 AGCACTGGCTCTGGGCGAGCGATGAGAGCTACGCCGCTACCTAAGGAGCATCGGCGT
 TGATGCCTGGCGCTTTGACTACGTGAAGGGCTACGGAGCGTGGGTCTCAAGGACTGG
 CTCAACTGGTGGGCTGGCTGGGCGGTCGGGAGTACTGGGACACAAACGTTGATGCAC
 TGCTCAACTGGGCCTACTCGAGCGATGCAAAAGTCTTCGACTTCCCGCTCTACTACAAG
 ATGGACGAGGCCTTCGATAACAACAACATTCCCGCCCTGGTGGACGCCCTCAGATACG
 GTCAGACAGTGGTCAGCCGCGACCCGTTCAAGGCTGTGACGTTTGTAGCCAACCACGA

FIGURE 16V

TACCGATATAATCTGGAACAAGTACCTTGCTTATGCTTTCATCCTCACCTACGAAGGCC
 AGCCCGTCATATTCTACCGCGACTACGAGGAGTGGCTCAACAAGGACAGGTTGAACAA
 CCTCATATGGATACACGACCACCTCGCAGGTGGAAGCACTGACATCGTTTACTACGAC
 AACGACGAGCTGATATTCGTGAGAAACGGCTACGGAAGCAAGCCGGGACTGATAACA
 TACATCAACCTCGCCTCAAGCAAAGCCGGAAGGTGGGTCTACGTTCCGAAGTTCGCGG
 GAGCGTGCATCCACGAGTACACCGGCAACCTCGGCGGCTGGGTGGACAAGTGGGTGG
 ACTCAAGCGGGTGGGTGTACCTCGAGGCCCTGCCACGACCCGGCCAACGGCTATTA
 CGGCTACTCCGTCTGGAGCTATTGCGGTGTTGGCTGA

SEQ ID NO.: 58

MAKYLELEESGVIMQAFYWDVPSGGIWWDTIRQKIPEWYDAGISAIWIPPASKGMSGGYS
 MGYDPYDYFDLGEYYQKGTVETRFGSKQELINMINTAHAYGIKVIADIVINHRAGGDLEW
 NPFVNDYTWTDFSKVASGKYTANYLDFHPNEVKCCDEGTFGGFPDIAHEKSWDQHWLW
 ASDESYAAYLRSIGVDAWRFDYVKGYGAWVVKDWLNWWGGWAVGEYWDTNVDALL
 NWAYSSDAKVDFPLYKMDFAFDNNNIPALVDALRYGQTVVSRDPFKA VTFVANHDTD
 ITWNKYLAYAFILTYEGQPVIFYRDYEEWLNKDRNLNLIWIHDHLAGGSTDIVYYDNDELIF
 VRNGYGSKPGLITYINLASSKAGRWVYVPKFAGACIHEYTGNLGGWVDKWVDSSGWVY
 LEAPAHDPANGYYGYSVWSYCGVG

SEQ ID NO.: 59

atggccaagtacctggagctcgaagagggcggggcataatgcaggcgttctactgggacgtgccttcaggaggaatatgggggacacaa
 acggcagaagataccggagtggtacgatgccggaatctccgcaatgatgattctcccgagcaagggtatgagcggcggtattc
 gctacgaccctacgattatttgacctcggtagtactaccagaagggaacgggtgaaacgaggttcggctcaaagcaggagctcataa
 gataaacaccgccacgcctacggcatcaaggctacgcagacatagtaataaccaccgcggcggaggagaccttgagtggaacccctc
 tcaatgactacacctggagcggacttctgaaggctcgttcggcgaatcacggccaactacctgacttcacccgaacgagctccatgcg
 gcgattccggaacatttggaggctatcccgacatagccacgacaagagctgggaccagtaactggctctggccagccaggagagctacg
 gcatactcaggagcatcgccatcgatgcctggcgcttcgactacgtaagggtatgctccctgggtcgtcaaggactggctgaactgggg
 gaggtggcggttgagagtagtgggacaccaacgctcagcgtgttctcaactgggcatactcgagcgggtccaaggtcttgacttcg
 ctactacaagatggagcaggccttgataacaacaacattcccgccctgggtggagcgcctcagatacgggtcagacagtggtcagccg
 cgttcaaggctgtgacgtttagccaaccacgataccgataaatttgaacaaglacccggcctacgccttcacccacacgagggccag
 ccgacgataattaccgcgactacgaggagtggtcacaaggacagggtcaagaacctcatctggatacacgaccctcggcggtggaag
 cactgacatcgttactacgacaacgacgagctgatatgtgagaacggctcgggaagcaagccgggactgataacatacaaacctgc
 gtaagcaagccggaagggtggttatgtgccaaggtcggggcggtgcacccagagatactgtaacctcgaggttggttagaca
 agtagcttactcaagcggctgggttatctcgaagctccagcttacgacctgccaacgggcagtagtggctactccgtgtggagctattgcggt
 gttgggtga

SEQ ID NO.: 60

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser-Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr

FIGURE 16W

Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr
 Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu Tyr
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 61

atggccaagtactccgagctgaaaaaggcgggggtcataatgcaggcgttctactgggacgtgccttcaggaggaatatgggtggacacaat
 acggcagaagataccggagtggtacgaggcgggaataatcgccatttgattctcccgagcaagggtatgagcggcggtctattcgatgg
 gctacgacccctacgattattttgacctcgggtgagtactaccagaagggaacgggtgaaacagggttcggctcaaacaggagctcataaacat
 gataaacaccgcccacgcctacggcatcaaggatcgcagacatagtaataaccaccgcccggaggagaccttgagtgaaccccttcg
 tcaatgactacacctggacggacttctgaaggtcgttccggcaagtagcacggccaactacccaacttccacccgaacgagctccatgcgg
 gcgattccggaacatttgaggctatcccacatatgccacgacaagagctgggaccagtagctctggccagccaggagagctacgcg
 gcatactcaggagcatcggcatcgatgcctggcgcttcgactacgtcaagggtacggagcgtgggtcgtcaaggactggctggactgggtg
 gggaggtcggccgctggggagtagctgggacacaaacgttgatgcactgctcaactgggcctactcgaagcgtgcaaaagtcttgactccc
 gctctactacaagatggatgagcccttgacacaaaaacattccagcgtcgtctctgcccttcagaacggccagactgttctctcccgacc
 cgttcaaggccgtaaccttttagcaaacatgacaccgatataatctggaacaagtagtcagcctacgcgttcatctacccacgagggccag
 ccgacaatatctaccgcgactacgaggagtggtcacaacaggataagctcaagaacctcatctggatacatgacaacctcggcggaggaaag
 caccgacatagctactacgataacgatgaactcatcttcgtcaggaaacggctacggggacaagccggggcttataacctacatcaacctaggc
 tcgagcaaggccggaaggtgggtctacgttccgaagttcgcgggagcgtgcacccacgagtagaccggcaacctcggcggctgggtggaca
 agtgggtgactcaagcgggtgggtgtacctcgaggccctcgccacgacccggccaacggctattacggctactccgtctggagctactgc
 ggggtgggtga

SEQ ID NO.: 62

Met Ala Lys Tyr Ser Glu Leu Lys Lys Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asn Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
 Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
 Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr
 Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 63

atggccaaglacctggagctcgaagaggcgggggtcataatgcaggcgttctactgggacgtgccttcaggaggaatatgggtggacacaat
 acggcagaagataccggagtggtacgatgccgaatctccgcaatatggattccccggcgagcaagggtatggcgggcgctattcgatg
 ggctacgacccctacgacttctttgacctcgggtgagtacgaccagaagggaacgggtgagacgcgccttggctccaaagcaggagctcgtgaa
 catgataaacacggcccatgcctcggcataaaggccatagcggacatcgtcatzaaccacccggcgaggcggagacctcgtggaacccg

FIGURE 16X

ttcgttggggactacacctggacggacttctcaaagggtggcctcgggcaataatactgccaactacctcgaacttccacccaacgaggtcaagt
gctgtgacgagggcacatttgaggcttccagacatagcccacgagaagagctgggaccagcactggctctgggcgagcgatgagagcta
cgccgcctacctaaggagcatcggttgatgcttggcgcttgactacgtgaagggtacggagcgtgggtcgtcaaggactggctcaactg
gtggggcggtcggcggttggtgagtgactgggacaccaacgttgatgcactcctcaactgggctactcgaaggcgccaaggtcttcgactt
cccgtctactacaagatggacgcggccttgacaacaagaacattcccgaactcgtcgaaggccctcaagaacgggggcacagtcgtcagcc
gcgaccggttaaggccgtaaccttcgttgcaaacacgacaccgataatctggaacaagtatccagcctacgcgttcacctcacctacgag
ggccagccgacaatattctaccgcgactacgaggagtggtgcaacaaggataagctcaagaacctcatctggatacatgacaacctcggcg
aggaagcaccgacatagctactacgataacgatgaactcatctcgtcaggaaaggctacggggacaagccggggcttataacctacatcaa
cctaggtggagcaaggccggaagggtgggttatgtgccgaagttcggcggtcgtgcatccacgagtatactgtaacctcggagggtggg
tagacaagtacgtctactaagcggtgggtctatctgaagctccagcttacgacctgccaacgggcagtatggctactccgtgtggagcta
ctgcgggggtggggtga

SEQ ID NO.: 64

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp
Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn
Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Ala Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly
Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr
Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp
Val Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp
Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp
Ala Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser
Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro
Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu
Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val
Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
Ile Asn Leu Gly Trp Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His
Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu
Ala Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 65

atggccaagtactccgagctggaagaaggcggcggtataatgcaggccttacttgggacgtcccagggtggaggaatctggtggggcaccat
caggagcaagataccggagtgtacgaggggggaatatccgcattttgattactcgcgcgagcaagggtatagcggcggtattcgaagg
gctacgacccctacgattatgacgggggaggtacacgaagaagggaagggtggaacgaggttggctcaagaaggaggtcctataaacat
gataaacaccgcccacgctatggatgaaggtaataggcgatatagtcataaccacgcgcggcggtgacctggagtggaaaccttctgt
gaacgactatacctggaccgacttctcaaaggctcgcgtcgggtaaatacacggccaactacctcgaacttccaccgaacgagctccatgcggg
cgattccggaacatttggaggctatcccgacatagccacgacaagagctgggaccagtaactggctctgggccagccaggagctacgcgg
catactcaggagcatcgcatcgaatgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactgggtggg
aggctggcggttggagagtactgggacccaacgtcgcgctgttctcaactgggcatactcgaagggtgccaaggcttctgacttgcctc
tactacaagatggacgagggccttgataacaacaacattcccgccttgggtggacggcctcagatacggtcagacagtggtcagccgcgaccc
gttcaaggctgtgacgtttgtgccaaccacgataccgatataatttggacaagtaacccggcctacgccttcatctcacctacgagggccagc
cgacgatattctaccgcgactacgaggagtggctcaacaaggacaggtcgaagaacctcatctggatacacgaccacctcggcggtggaagc
acgagcatagttactacgacgcgacgagatgatcttctgtgaggaacggctatggaagcaagcctggcctataacttcatcaacctcggctc
gagcaagggttggaaagggtgggttacgttccgaagttcgcaggctcgtgcatacacgagtacaccggcaatctcggcggttgggtggacaagt
gggtggactcaagcggtgggtctacctcgaaggctcctgccacgaccggccaacggccagttacggctactccgtctggagctatttgcggt
gttggctga

FIGURE 16Y

SEQ ID NO.: 66

Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly
 Gly Gly Ile Trp Trp Gly Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp Leu
 Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile
 Asn Thr Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr
 Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val
 Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr
 Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr
 Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 67

atggccaagtacctggagctcgaagagggcggggtcataatgcaggcggttctactgggacgtgccttcgggaggaatatggtgggacacaat
 acggcgagaagataccggagtggtacgatgccggaatctccgcaatatggattctcccgagcaagggtatgagcggcggtattcgaatg
 gctacgacccctacgattatttgacctcggtagtactaccagaagggaacgggtggaacgaggttcgggtcaaaagcaggagctcataaacat
 gataaacacggcccatgcctacggcataaaggatagcgacatcgtcataaaccacgcgagggcggagacctcgagtgaacccgttc
 gttggggactacacctggacggactctcaaaagggtggcctcgggcaaatatactgccaactacctgacttccaccccaacgaggtcaagtgt
 gtgacgagggcacatttgagggttccagacatagcccacgagaagagctgggaccagcactggctctgggcgagcgatgagagctacg
 ccgctacctaaggagcatcggcgttgatgcctggcgttcgactacgtcaagggtactcgagcgtgggtcgtcaaggactggctggactgtg
 ggggaggtcggcggtcggggagtgactgggacacaaacgttgatgcactgctcaactgggctactcagcagatgcaaaagtcttcgacttc
 ccgctctactacaagatggagcaggccttcgataacaacaacattcccgcctgggtggacgcctcagatagcgtcagacagtggtcagccgc
 gaccggtcaaggctgtgacgtttgtagccaaccacgataccgataataatctggaacaagtatccagccctacgcgttcacctacacaggg
 ccagccgacaaatttaccgcgactacgaggagtggtcacaagaagataagctcaagaacctcatctggatatacgaacctcggcggag
 gaagcagcagcatggttactacgacagcgacgatgatctcgtgaggaacggctatggaagcaagcctggcctataacttatcatcaacct
 cggtcgcagcaaggttggaaggtgggtctacgttccgaagttcggggagcgtgcatccacgagtacaccggcaacctcggcggctgggtg
 gacaaagggtggaactcaagcgggtgggttaactcgaagccctgcgaagacccggcgaacggctattacggctactcgtctggagcta
 ctggcgttgggtgga

SEQ ID NO.: 68

Met Ala Lys Tyr Leu Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
 Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met
 Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe Gly Gly
 Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser Tyr Ala
 Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu

FIGURE 16Z

Ala Phe Asp Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala
 Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn
 Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr
 Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile
 Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala
 Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Val Val Gly

SEQ ID NO.: 73

atggctctggaagaggggcgttataatgcaggccttactgggacgtcccaggagggaatctggtgggacaccatagcccagaagat
 acccgactggggcagcgccgggatttcggcaatatggattcctccgcgagtaaggcatgagcggcggtattcgatgggctacgaccct
 acgatttcttgacctcggtagtactaccagaagggaagcgttgagacccgcttcggatcaaaaggagagcttgtaacatgataaacaccgc
 ccatgctcacaacatgaaggtcatagcggacatagtcacaaaccgcgcggcgacgtgagtggaatccttcaccaacagctacac
 ctggaccgatttctcgaaggctcgcgtcgggcaagtacacggccaactacctgacttccaccgaacgagcttcacgcggcggtaccgga
 catttgagggtatcccgacatatgccacgacaagagctgggaccagcactggctcgggccaacgaaagctacgccgctacctccgg
 agcatcggcatcgacgcctggcgcttcgactacgtcaagggtacgctccctgggtcgttaagaactggctgaaccgggtgggcggctgggc
 gggtggagagtactgggacaccaacgtcgtgactcctgagctgggctacgacagcggtgctaaagcttcgacttccgcttactacaag
 atggacgaggccttcgataacaacaacatccccgccctcgtggacgccctcaagaacggagggcacggctgcagccgcgacccgttcaag
 ccgtgaccttcgtgccaaccacgataccaacataatctggaacaagtatccggcctacgccttcatcctacatgaggagacggcgaat
 attctaccgagctacgaggagtggtcacaaggacaggtcaggaacctcatctggatacacgaccacctcggggaggaagcacagac
 atcatctactacgacagcgagcttatctcgtgagaacgggtacggggacaagccgggactgataacctacatcaacctcggtcgaagc
 aaggccggaaaggtgggtctacgttcgaagttcgcaggctcgtgcatacacgagtagaccggcaacctcgcggtggtgattgacaagtgggt
 tgactcaagcggtcgggtctacctgaggccccgccacgacccggccaacggccagtacggctactcctgtaggtactcgtggtg
 ggtga

SEQ ID NO.: 74

Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp
 Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala
 Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr
 Tyr Gln Lys Gly Ser Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala
 His Ala His Asn Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp
 Asn Pro Phe Thr Asn Ser Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn
 Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro Asp Ile
 Cys His Asp Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala Tyr Leu Arg
 Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Lys Asn Trp
 Leu Asn Arg Trp Gly Gly Trp Ala Val Gly Gln Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Ser Trp
 Ala Tyr Asp Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp
 Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg Asp Pro Phe
 Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asn Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 Leu Thr Tyr Glu Gly Gln Pro Ala Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Arg Leu
 Arg Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Ile Tyr Tyr Asp Ser Asp Glu
 Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser Ser
 Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu
 Gly Gly Trp Ile Asp Lys Trp Val Asp Ser Ser Gly Arg Val Tyr Leu Glu Ala Pro Ala His Asp Pro
 Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 75

atggctctggaagaggggcgttataatgcaggcatttactgggacgtcccaggagggaatctggtgggacacgatagcccagaagat
 acccgactgggcaagcgccgggatttcggcgatatggattccccgcgagcaagggtatgagcggcggtattcgatgggctacgacccct

FIGURE 16AA

acgattattttgacctcggtagtactaccagaagggaacgggtggaacaagattcggctcaaagcaggagctcataaacatgataaacaccg
 cccacgcctatggcatgaaggtaatagccgatatagtcatacaaccaccgcgccggcgatctggagtggaaccccttcgtgaacgactata
 cctggaccgacttctcgaaggtagcgtcgggtgaaatacacggccaactacctcgactccaccgaacgagctccacgcggcgattccgga
 acatttggaggctatcccacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctacgcggcctatctcag
 gagcatcggcatcgacgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaggagactggctgaactggtggggaggctggg
 cagttggagagtactgggacccaacgtcgacgctgttctcaactgggcatactcgagcgggtccaaggctttgacttcgcccttactacaag
 atggacgaggcccttcgataacaacaacattcccgcctggtagacgccctcagatacggccagacagtggtcagccgcgacccgttcaaggc
 tgtgacgttttagccaaccacgataccgacataatctggaacaagtatccagcctacgcgttcacctacgagggccagccgacaatat
 tctaccgcgactacgaggagtggtcacaaggacaagctcaagaacctcatctggatacatgacaacctcggcgaggaggagcactgacatc
 gtttactacgacaacgacgagctgatatctgagaacggctacggaagcaagccgggacgataacatacatcaacctcggctcaagcaa
 gccggaagggtgggttacgttccgaagttcgaggctcgtgcatacacgagtacaccggcaacctcggcggtgggagcaagtggtgga
 ctcaagcggctgggttacctcgaggctcgtcccacgacccggccaacggccagtacggctactcgttggagctattgcggtgtgggtga

SEQ ID NO.: 76

Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp
 Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala
 Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr
 Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr Ala
 His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp
 Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn
 Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro Asp Ile
 Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg
 Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Arg Asp Trp
 Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp
 Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn
 Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp Pro Phe Lys
 Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu
 Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp
 Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser Ser
 Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu
 Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp Pro
 Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 77

atggctctggaagaggcgggctcataatgcaggccttctactgggacgtcccatgggaggaatcgggtgggacacgatagccagaagat
 acctgagctgggaaggcggcggttctggcgatatgataccctccgcgaggaagggtatgagcggcggttatcgatggctacgacct
 acgattattttgacctcggtagtactaccagaagggaacgggtggaacaagaggttcggctcaaagcaggagctcataaacatgataaacaccg
 cccacgcctatggcatgaaggtaatagccgatatagtcatacaaccaccgcgccggcggtgacctggagtggaaccccttcgtgaacgactata
 cctggaccgacttctcaaaggtagcgtcgggtgaaatacacggccaactacctcgactccaccgaacgagctccatgcggcgattccgga
 catttggaggctatcccacatatgccacgacaagagctgggaccagtactggctctgggccagccaggagagctacgcggcctatctcagg
 agcatcggcatcgatgcctggcgcttcgactacgtcaagggtatgctccctgggtcgtcaaggactggctgaactggtggggaggctgggc
 ggttggagagtactgggacccaacgtcgacgctgttctcaactgggcatactcgagcgggtccaaggctttgacttcgcccttactacaaga
 tggacgaggcccttcgataacaacaacattcccgcctggtagacgccctcagatacgggtcagacagtggtcagccgcgacccgttcaaggct
 gtgacgttttagccaaccacgataccgacataatctggaacaagtatccagcctacgcgttcacctacctacgagggccagccgacaatat
 ctaccgcgactacgaggagtggtcaacaaggataagctcaagaacctcatctggatacatgacaacctcggcgaggaggagcactgacatcg
 ttactacgacaacgacgagctgatatctgagaacggctacggaagcaagccgggactgataacatacatcaacctcgcctcaagcaang
 ccggaagggtgggttacgttccgaagttcgaggctcgtgcatacacgagtacaccggcaatctcggcggtgggtggacaagtgggtggac
 tcaagcggctgggttacctcgaggctcgtcccacgacccggccaacggccagtacggctactcgttggagctactcgggtgttgggtga

FIGURE 16BB

SEO ID NO.: 78

Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp
Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala
Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr
Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr Ala
His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp
Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn
Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro Asp Ile
Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg
Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Lys Asp Trp
Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp
Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn
Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp Pro Phe Lys
Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu
Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp
Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Ala Ser Ser
Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu
Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp Pro
Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEO ID NO.: 79

atgaagcctgcgaactcctcgtctttgtgctcgtagtctctatcctcgcggggctctacgccagccgcgggggcggcgaagtacctggagc
tcgaagaggcgcggtcataatgcaggcgttctacttgggagctgacctcaggaggaatatgtgtgggacacaatacggcagaagataccgga
gtgtgtacgatgccggaatctccgcaatatggattccccggcgagcaagggcattggcgccctattcgtatgggtacgacccctacgactt
ctttgacctcgggtgagtacgaccagaagggaacggtagagacgcgcttggctccaagcaggagctcgtgaacatgataaacaccgccacg
cctacggcatcaaggctatcgcagacatagtaataccaccgcgccggaggagacattgagtgaaccccttcgtaatgactacacctgga
cggacttctcgaaggctcgttcggcaagtacacggccaactacctgacttccaccccaacgaggtcaagtgctgcgacgaggggcacctttg
gagggttccccggacatagccacgagaagagctgggaccagtactggctctgggcgagcaacgagagctacgccgcctacctcaggagca
tcggcggttgacgatggcgcttcgactacgtcaagggtctacggagcgtgggtcgtcaaggactggctggactgggtggggaggctgggcccgt
cgggggagtactgggacacaaacgttgatgactgtcactgggcttactcgaagcgtgcaaaaagcttcgacttccccgtctactacaagatg
gacgcggcccttgacaacaagaacattcccgactcgtcgaaggccctcaagaacgggggcacagtcgtcagccgogacccttaagggcgt
aaccttcgttgcaaacacgacacggacataattggaaacagtaccggccctacgcttcatctcactacgtacgagggccagccgacgatattc
taccgcgactacgaggagtgtggtcacaaggacaggctcaagaacctcatctggaacacgaccacctcgcgggtggaagcaccgacatag
ctactacgataacgatgaactcatcttcgcaggaaacggctacggggacaagccggggcctataacctacatcaacctaggctcagcaagg
ccgggaggtgggttacgttcgaaggtggggggaggtgcatcacaggtacacggcaacctcggcgggggggtggacaagtggtggga
ctcaagggggtgggtgactctcagggtcgtggcagcagccggcaacgggtattagggtactccgttggaggtactcgggggtgggt
ga

SEO ID NO.: 80

Met Lys Pro Ala Lys Leu Leu Val Phe Val Leu Val Val Ser Ile Leu Ala Gly Leu Tyr Ala Gln Pro
 Ala Gly Ala Ala Lys Tyr Leu Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val
 Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu. Trp Tyr Asp Ala Gly. Ile Ser Ala
 Ile Trp Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe
 Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val
 Asn Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly
 Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr. Asp Phe Ser Lys Val Ala Ser Gly
 Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Asn Glu Ser
 Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala

FIGURE 16CC

Trp Val Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val
 Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met
 Asp Ala Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val
 Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr
 Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp
 Leu Asn Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile
 Val Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr
 Tyr Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile
 His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu
 Glu Ala Pro Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 81

atgaagaagttgtcgcctgttcataacatgttttcgtagtgcagtcgttgcacagccagctagcgcgcaaagtattccgagctc
 gaagaaggcggcggtataatgcaggccttctactgggacgtccagggtggaggaatctggtgggacaccatcaggagcaagataccggagt
 ggtacgaggcgggaatatccgccatttgattccgccagccagcaaggggatgagcggcggttactcgatgggtacgatccctacgatttctt
 tgacctcggcgagtacaaccagaagggaaccatcgaaacgcgtttggctctaaccaggagctcatcaatatgataaacacggccatgccta
 cggcataaagggtacatagcggacatcgtaataaccaccgcgcaggcggagacctcgagtggaaacccgttcgttggggactacacctggacg
 gacttctcaaaagggtggcctcgggcaaatatactgccaactacctgacttccacccaacgaggtcaagtgtgtgacgagggcacatttggag
 gcttccagacatagcccacgagaagagctgggaccagcactggctctgggcgagcgatgagagctacgccgcctaccttaaggagcatcg
 gcgttgatgcctggcgctttgactacgtgaagggtacggagcgtgggtcgtcaaggactggctcaactgggtggggcggtggcggttggc
 ggtactgggacaccaacgttgatgcactcctcaactgggcctactcgagcggcgccaaggtcttcgactcccgctctactacaagatggatg
 aggccttgacaacaaaaacattccagcgctcgtctcgtccctcagaacggccagactgttctccgcgacccgttcaaggccgtaacctt
 gtgcaaacccacgacacgatataatctggaacaagtaccttgcattatgcttcatctcaccacgaaggccagcccgcatattctaccgcgac
 tacgaggagtggctcaacaaggacaggttgacaacctcatatggatacacgaccacctcgaggttggaagcacgagcatagtctactacga
 cagcgacgagatgatctcgtgaggaacggctatggaagcaagcctggccttataacttacatcaacctggctcgagcaaggttggaaggtg
 ggttatgtccgaagttcggggcgctgcatccacgagtatactgtaacctggaggctgggtgtagacaagtagctactacaagcggtg
 ggtctatctcgaaggtccagcttacgacctgccaacgggcagtatggctactccgtgtggagctattgcggtgttgggtga

SEQ ID NO.: 82

Met Lys Lys Phe Val Ala Leu Phe Ile Thr Met Phe Phe Val Val Ser Met Ala Val Val Ala Gln Pro
 Ala Ser Ala Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val
 Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala
 Ile Trp Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe
 Phe Asp Leu Gly Glu Tyr Asn Gln Lys Gly Thr Ile Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile
 Asn Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly
 Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly
 Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Gln Gly Thr Phe
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser
 Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala
 Trp Val Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val
 Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met
 Asp Glu Ala Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln Thr Val Val
 Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr
 Leu Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp
 Leu Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Ser Ile
 Val Tyr Tyr Asp Ser Asp Glu Met Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr
 Tyr Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile
 His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu
 Glu Ala Pro Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

FIGURE 16DD

SEQ ID NO.: 83

atggctctggaagacggcgggctcataatgcaggccttctactgggatgttccctggaggaggaatctgggtggacacaaatagctcaaaagata
cccgaatgggcaagtgcaggaaatctcagcgatatggattccaccagcgagtaaggggcatgagcgggtgttatccatgggctacgatccctac
gatttcttgacctcggcgagtactatcagaaggggacagttgagacgcgcttcggctcaagggaagaactggtaacatgataaacaccgca
cactcctacggcataaaggatagcagacatagtcataaaccaccgcgcgggtggagaccttgagtggaaacccctcgtgaacgactatacct
ggacagacttctcaaaagtcgctccggtaaatatacggccaactaccttgacttccaccaaacgagcttcaactgtgtgatgaaggtaaccttg
gaggataccctgatatgtcacgacaaaagctgggaccagctactggctctgggcgagcagcgaaagctacgctgcctacctcaggagcata
gggggtgacgcctggcgttcgactacgtcaagggtacggagcattgggtgttaacgactggctcagctgggtgggaggctgggccgttgga
gagtactgggacacgaacgttgatgcactcctcaactgggcatacagcagcgccgccaaggcttctgacttcccgtctactacaagatggacg
aagccttcgacaacaccaacatcccggcattagtgatgcactcagatacggccagacagtggtcagccgcgacccctcaaggcggtaacttt
cgttgccaaccacgatacagataatctggaacaagatccggcttatgcattcacttacctatgaggggacagcctgttatctaccgcgac
tacgaggagtggctcaacaaggataagcttaacaacctatctggatacagatcaccttctggaggagtagtactgacattgtttactacgacag
cgacgagcttatcttgtgagaacggctatggcaccacacaggaactgataacctatacaacctcggctcaagcaaagtggaaagggtgggc
tacgttccaaagttcggcggttcgacccacgagtacaccggcaacctcggcggttggaagacaagtagcttctccagcggctgggtct
atcttgaggccccagcccacgacccggcgaacggctactacggctactccgtatggagctactgcggggttggtga

SEQ ID NO.: 84

Met Ala Leu Glu Asp Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp
Trp Asp Thr Ile Ala Gln Lys Ile Pro Glu Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala
Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr
Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala
His Ser Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp Asn
Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
Leu Asp Phe His Pro Asn Glu Leu His Cys Cys Asp Glu Gly Thr Phe Gly Gly Tyr Pro Asp Ile Cys
His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Ser Glu Ser Tyr Ala Ala Tyr Leu Arg Ser
Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Asn Asp Trp Leu
Ser Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala
Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Thr
Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp Pro Phe Lys Ala
Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu
Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu Asn
Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Ser Asp Glu
Leu Ile Phe Val Arg Asn Gly Tyr Gly Thr Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser Ser
Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu
Gly Gly Trp Ile Asp Lys Tyr Val Ser Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp Pro
Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO.: 85

atggctctggaagaggcgggcttataatgcaggcattctatgggacgtcccagggtggaggaatctgggtggacaccatagcccagaagata
cccgaatgggcaagtgcaggaaatctcagcgatatggattccaccagcgagtaagggaatgagogggtgttatccatgggctacgatccctac
gatttcttgacctcggcgagtactatcagaaggggacagttgagacgcgcttcggctcaagggaagaactggtaacatgataaacaccgca
cactcctacggcataaaggatagcggacatagtcataaaccaccgcgcgggtggagggcctcagtggaacccctcgtgaacgactatacc
tgacagacttctcaaaagtcgctccggtaaatatacagccaactaccttgacttccaccaaacgagcttcaactgtgtgatgaaggtaaccttg
gaggataccctgatatgtcacgacaaaagctgggaccagctactggctctgggcgagcagcgaaagctacgctgcctacctcaggagcata
gggggtgacgcctgggtgttcgactacgtcaagggtacggagcctgggtgttaacgactggctcagctgggtgggaggctgggccgttgga
gagtactgggacactaacgttgatgcactcctcaactgggcatacaacagcgccgccaaggcttctgacttcccgtctactacaagaatggacg
aagccttcgacaataccaacatcccgcgtttgttacgcctcaagaatggcgggacagtggtcagccgcgacccattcaaggcggtaacttt
cgttgccaaccacgatacagataatctggaacaagatccggcttatgcattcacttacctatgaggggacagcctgttatctaccgcgac
tacgaggagtggctcaacaaggataagcttaacaacctatctggatacagatcaccttctggaggaggtagtactgacattgtttactacgacag
cgacgagcttatcttgtgagaacggctatggcaccacacaggaactgataacctatacaacctcggctcaagcaaagctggaaagggtgggc

FIGURE 16EE

tacgttccaaagtgcgcggttcacatccacgagtiacaccggcagcctcggcggttgatagacaagtacgtctcctccagcggctgggct
accttgaggccccggccacgacccggccaatggccagtatggctactcgtctggagctattgcggggttgggtga

SEQ ID NO.: 86

Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp
Trp Asp Thr Ile Ala Gln Lys Ile Pro Glu Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala
Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr
Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala
His Ser Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Gly Leu Glu Trp Asn
Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
Leu Asp Phe His Pro Asn Glu Leu His Cys Cys Asp Glu Gly Thr Phe Gly Gly Tyr Pro Asp Ile Cys
His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Ser Glu Ser Tyr Ala Ala Tyr Leu Arg Ser
Ile Gly Val Asp Ala Trp Cys Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Asn Asp Trp Leu
Ser Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala
Tyr Asn Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn
Thr Asn Ile Pro Ala Leu Val Tyr Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg Asp Pro Phe Lys
Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu
Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Ser Asp
Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Thr Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser
Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Ser
Leu Gly Gly Trp Ile Asp Lys Tyr Val Ser Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp
Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO: 87

atgttctgtctcgcgtttttgtcactgcctcgtgttctgcccacaggacagccgccaaaggctgccgcaccgttaacggcaccatgatgca
gtattttgaatggtacttgcgggatgatggcacgttatggaccaagtggccaatgaagccaacaactatccagccttggcatcaccgctcttg
gctgccgccgcttacaanggaacaagccgcagcgacgttagggtagcggagtatacgaactgtatgacctcggcgaaatcaatcaaaaaggga
ccgtccgcacaaaatacgaacaaaagctcaatatttcaagccaattcaagccgccacgccgctggaatgcaagtgtacgcgcgatgtcgtgtt
cgaccataaaggcggcgctgacggcacggaatgggtggacgccgtcgaagtcaatccgtccgaccgcaaccaagaatctcgggcacatt
caaatccaagcatggacgaaatttgatttcccggcggggcaacacctactccagctttaagtggcgctgtgtaccatttgacggcggttgattg
gacgaaagccgaaatgagccgcatttacaattccgcggtcgcgcaagcgctgggattgggaagtagacacggaaacggaactatg
actacttaattgtatgccgacctgatatggatcaccgaagtcgtgaccgagctgaaaaactgggggaaatggatgtcaacacacgaacatt
gatgggttcggcttgatgccgtcaagcatattaagttcagtttttcttgattggtgtcgtatgtcgttctcagactggcaagccgctattaccg
tcggggaatattggaactatgacatcaacaggtgcacaaattacattacgaaacagacggaacgatgtcttittgtatgcccggttacacaaca
aattttacogottcaaatcaggggcggttggatggcgcgttaattgacaaatctcattgaaagatcaacggacattggcgcacctt
cgttgataatcattacacggcggcgaagcggaagtcgtgacacatgggtgaaacgttggcttgcctttatccttgaatccttgaatccttga
aggaaagataccgtgcgtttttatggtgactatattgcaatccacataacattcctcgtgaaaagcaaaatcgatccgctcctcctatgc
gcgcagggtattatgctacggaacgcaacatgattatctgatcactccgacatcatcgggtggacaagggaagggtgactgaaaaaccagg
atccgggctggccgactgatcaccgatggcgccggagggaagcaaatggatgtactgttggcaacaacacgctggaaaagtgtctatga

SEQ ID NO: 88

Met Phe Leu Leu Ala Phe Leu Leu Thr Ala Ser Leu Phe Cys Pro Thr Gly Gln Pro Ala Lys Ala Ala
Ala Pro Phe Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr
Lys Val Ala Asn Glu Ala Asn Asn Leu Ser Ser Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr
Lys Gly Thr Ser Arg Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln
Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Leu Gln Ala Ile Gln Ala Ala His Ala
Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asp His Lys Gly Gly Ala Asp Gly Thr Glu Trp Val
Asp Ala Val Glu Val Asn Pro Ser Asp Arg Asn Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala Trp
Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp

FIGURE 16FF

Gly Val Asp Trp Asp Glu Ser Arg Lys Leu Ser Arg Ile Tyr Lys Phe Arg Gly Ile Gly Lys Ala Trp
 Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp
 His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly Lys Trp Tyr Val Asn Thr Thr Asn Ile Asp Gly
 Phe Arg Leu Asp Ala Val Lys His Ile Lys Phe Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Ser
 Gln Thr Gly Lys Pro Leu Phe Thr Val Gly Glu Tyr Trp Ser Tyr Asp Ile Asn Lys Leu His Asn Tyr
 Ile Thr Lys Thr Asp Gly Thr Met Ser Leu Phe Asp Ala Pro Leu His Asn Lys Phe Tyr Thr Ala Ser
 Lys Ser Gly Gly Ala Phe Asp Met Arg Thr Leu Met Thr Asn Thr Leu Met Lys Asp Gln Pro Thr
 Leu Ala Val Thr Phe Val Asp Asn His Asp Thr Glu Pro Gly Gln Ala Leu Gln Ser Trp Val Asp Pro
 Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly
 Asp Tyr Tyr Gly Ile Pro Gln Tyr Asn Ile Pro Ser Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile Ala Arg
 Arg Asp Tyr Ala Tyr Gly Thr Gln His Asp Tyr Leu Asp His Ser Asp Ile Ile Gly Trp Thr Arg Glu
 Gly Val Thr Glu Lys Pro Gly Ser Gly Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp
 Met Tyr Cys Trp Gln Thr Thr Arg Trp Lys Ser Val Leu

SEQ ID NO: 89

atgaaagaagcgggtgtgtatcaaattttcccgatcgggttttaattggcaaccttcaaatgataacgcaagcagcaggcacgcgggggcgc
 agccgattgagcatcgcgattggcggatttcccgaataatccgcctgaaagggacgagcggctacgatggcgacgggtgaatggtcgaat
 gacttttccggcgagacatcgccggaattgaacaaagtggattattgagtcgcttggagtgaacacgatttacttaaatccgatcgccaatg
 cgccatcgaaaccataaataatgatgcgagcaattacaagaattggatccgatgttcggttccccggaagaattccaatcggttggcaggcgcttg
 cgaaccgggggatgcattcatcttagacgggggtgtcaaccacgtatccgacgattcgaattttagaccgctaccaccgctatccgaccgto
 ggtgcgatgaatttgggaagcgggttacgattgatgaatgaaaaggattgagcgagggaagaagcgcgggaacaagtggaagagaagttc
 aaacaagagggacagacgttcagcccgtatgggttcatcttggttcaattatgaaaacaaaaagtcgaatggccattatcaataccaatcatggt
 ggggctatgacagtcgtccggagtttaagtcggtgacgggggaaaaagtgcccatccgagtgaaatgaacaacgatgcgtcgcgaattac
 attttccgtgaatcggttcgggtggcgaagaagctggatggcctcggcgctccggctggcgggttgatgtggccaatgaggtggtatccggcgt
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 ttgggatgacgcatcgaataatttttaggcgaccagtagcttcgtatgaactaccggttccgcggggcggtgcttacttttgaaaaacg
 gaaatgcagaagagggcggaacaagcggctgacggccataagggaagactaccacaagtgaaagcgtttatgcgtgatgaacttaacggttcg
 catgacacggcgggcggttcttctgttgggaacggaacggattcatccgagcggcggaagcttgatccgaattataatgaggaacttggg
 aaaaagcggcctcaagctggcggtgatttgcagatgggataccgggagcggcagcatttattacggcgatgaagcgggagtaacaggctc
 aaaagaccagacaaccggcgacgtatccggtgggcaagaagatcaaaatctgttgccttattatcagaagaatggggcacattcgccagc
 accatcaatcgttgggtggccatggcgacatcaagacgggtgatgcgaaggggatgtatcgtatttggccccaatcggcggtgaagcgg
 cgctcatgcatcaaccggcgcaatgaggacaagacgggtggcggtgacgtcgttctgttgcctcgaacggcaccgtgcttaccgatgagtt
 gcatgatggcggggaagctacggctcgtggcgaacggtgacggtcagcattccggccctggatggacggatgatgttgggacgggtgacg
 gcggaatgcggcgagcagtcagcaatttgcagcgagcgcttcggatggtcgtgacgttaacgtgggaaggaatgcatcgagatacc
 gaattttagatccacgttaaaaggtgcccgttatcagatggtgcaagagacggaacaacttcggccacgacgtggttcgttgcgaacggaa
 cagcctattacttgcgttggcggttgcgtatgaagaagggaatgataccgaagtggaacgaatcgttgccttacttaccgctgac
 gagcgaaatgctcagttggtggaacggttggatgacaaacttgaattgcaagcggagcgaatggttgccttacttaccgctgac
 gtgacaagcaaggagcagctgatgggtgcaagcgggttgcgaatgaaggcgcgaatgacgaacatggaagaatcagagcggctt
 accaaggacaagacggcgacgccaacgtgtccgagctgcttcactccgctcgcgcagggagcgtatcgtatgcgtgacgacc
 aaccttggcgaggagtggtgtatcagaaagagaagcaagtgacgttggcgagacaacagcgaccaaatagcgccagcagacgcatcg
 agctgcggcgagcctgcgggtgaatcgggacaagtgaatttatcatggacgttgggtggaaaaagatggggatgcttattgttagcatcgag
 cgcaacgggtgatcgtgcatacaaccacttcgatcggcgattcatttacagactacgatgtcgaaaacggcaccgagtcacgtatgtgtcaa
 gttgtatgaccgcccggcaatgttggcggtcaaacacggtaagtgacggcgacattgtatgtgaaagtattttaaagtgaagcgg
 ccggattacacaccggttgatggccgaattacgattccgaacagcttgaaaggcgtggaacacaggggcctgggagatgtcgcgaacgggtgc
 ggtgacggccgattggcaattaccgctgaggtgacgaaggggaaacgatacctataagtatgtgaaaggcggtatcgtggatcaagagg
 ggttggccgaccatacggctgaggacgacaacgatgatgacgtgagctactacggctatgggacgattggcaccgacttgaagtacggctc
 cacaatgaaggaaacaatacgtatgtgcaagaccgatttgcgtggatcgatatggcggtcgtatcgaaagggtgcaaaaacagga
 agtcaagtgacgatcaagggaatgccattaaacgggtgtttgacgatcaatggcgagcgggtgccgattgatggcggatggcattctcgt
 acacgtttgcggcgccagccatcaaaaagaggtgttgcataatgaaccatcgccggaagcaaaacagccattttcaaacacgacggcg

FIGURE 16GG

gagcgattgcgaaaaacacaaaagattacgtgctgaatttagaagcaagcaattcaaaaagcttctcgagagtacttctagagcgccgcgg
gcccatcgattttccaccgggtgggtaccaggta

SEQ ID NO: 90

Met Lys Glu Ala Val Val Tyr Gln Ile Phe Pro Asp Arg Phe Phe Asn Gly Asn Pro Ser Asn Asp Asn
Ser Lys Gln Gln Ala Arg Gly Ala Gln Pro Ile Glu His Arg Asp Trp Ser Asp Leu Pro Asp Asn Pro
Arg Leu Lys Gly Thr Ser Gly Tyr Asp Gly Asp Gly Glu Trp Ser Asn Asp Phe Phe Gly Gly Asp Ile
Ala Gly Ile Glu Gln Lys Leu Asp Tyr Leu Gln Ser Leu Gly Val Asn Thr Ile Tyr Leu Asn Pro Ile
Ala Asn Ala Pro Ser Asn His Lys Tyr Asp Ala Ser Asn Tyr Lys Glu Leu Asp Pro Met Phe Gly Ser
Pro Glu Glu Phe Gln Ser Phe Val Gln Ala Leu Ala Asn Arg Gly Met His Leu Ile Leu Asp Gly Val
Phe Asn His Val Ser Asp Asp Ser Ile Tyr Phe Asp Arg Tyr His Arg Tyr Pro Thr Val Gly Ala Tyr
Glu Tyr Trp Glu Ala Val Tyr Asp Leu Met Asn Glu Lys Gly Leu Ser Glu Glu Glu Ala Arg Lys Gln
Val Glu Glu Lys Phe Lys Gln Glu Gly Gln Thr Phe Ser Pro Tyr Gly Phe His Leu Trp Phe Asn Ile
Glu Asn Lys Lys Val Asn Gly His Tyr Gln Tyr Gln Ser Trp Trp Gly Tyr Asp Ser Leu Pro Glu Phe
Lys Ser Val Thr Gly Glu Lys Val Pro His Pro Ser Glu Leu Asn Asn Asp Ala Leu Ala Asn Tyr Ile
Phe Arg Glu Ser Asp Ser Val Ala Lys Ser Trp Ile Ala Leu Gly Ala Ser Gly Trp Arg Leu Asp Val
Ala Asn Glu Val Asp Pro Ala Phe Trp Arg Glu Phe Arg Gln Glu Leu Leu Gln Gly Ser Tyr Gly Arg
Gly Pro Thr Leu Lys Glu Gly Glu Gln Pro Leu Ile Leu Gly Glu Ile Trp Asp Asp Ala Ser Lys Tyr
Phe Leu Gly Asp Gln Tyr Asp Ser Val Met Asn Tyr Arg Phe Arg Gly Ala Val Leu Asp Phe Leu
Lys Asn Gly Asn Ala Glu Glu Ala Asp Lys Arg Leu Thr Ala Ile Arg Glu Asp Tyr Pro Ser Glu Ala
Phe Tyr Ala Leu Met Asn Leu Ile Gly Ser His Asp Thr Ala Arg Ala Val Phe Leu Leu Gly Asn Gly
Thr Asp Ser Ser Glu Arg Ala Glu Leu Asp Pro Asn Tyr Asn Glu Glu Leu Gly Lys Lys Arg Leu
Lys Leu Ala Val Ile Leu Gln Met Gly Tyr Pro Gly Ala Pro Thr Ile Tyr Tyr Gly Asp Glu Ala Gly
Val Thr Gly Ser Lys Asp Pro Asp Asn Arg Arg Thr Tyr Pro Trp Gly Lys Glu Asp Gln Asn Leu
Leu Ser His Tyr Gln Lys Val Gly His Ile Arg Gln His His Gln Ser Leu Leu Ala His Gly Asp Ile
Lys Thr Val Tyr Ala Gln Gly Asp Val Tyr Val Phe Ala Arg Gln Tyr Gly Arg Glu Ala Ala Leu Ile
Ala Ile Asn Arg Gly Asn Glu Asp Lys Thr Val Ala Leu Asp Val Ala Ser Leu Leu Pro Asn Gly Thr
Val Leu Thr Asp Glu Leu His Asp Gly Gly Glu Ala Thr Val Ala Gly Gly Thr Leu Thr Val Thr Ile
Pro Ala Leu Asp Gly Arg Met Met Phe Gly Thr Val Thr Ala Glu Met Pro Ala Ala Val Ser Asn Leu
Gln Ala Ser Ala Ser Asp Gly Cys Val Thr Leu Thr Trp Glu Gly Asn Ala Ser Arg Tyr Arg Ile Tyr
Glu Ser Thr Leu Lys Gly Ala Gly Tyr Thr Met Val Gln Glu Thr Glu Thr Thr Ser Ala Thr Ile Gly
Ser Leu Thr Asn Gly Thr Ala Tyr Tyr Phe Ala Val Ala Val Asp Glu Asn Gly Asn Glu Ser Pro
Lys Val Glu Thr Asn Arg Val Val Pro His Tyr Pro Leu Thr Ser Asp Asn Val Gln Phe Val Thr Thr
Leu Ser Asp Ala Thr Leu Asp Leu Ser Lys Pro Gln Gln Val Asp Val His Val Asn Ile Asp Asn Val
Thr Ser Lys Gly Ala Ala Asp Gly Leu Gln Ala Val Leu Gln Val Lys Gly Pro His Asp Glu Thr Trp
Lys Glu Tyr Arg Ala Ala Tyr Gln Gly Gln Asp Gly Asp Ala Asn Val Phe Arg Ala Ala Phe Thr Pro
Leu Ala Ala Gly Thr Tyr Thr Tyr Arg Tyr Ala Leu Thr Thr Asn Leu Gly Glu Glu Trp Met Tyr Thr
Glu Glu Lys Gln Val Thr Phe Ala Ala Asp Asn Ser Asp Gln Ile Ala Pro Ala Asp Ala Ile Glu Leu
Arg Gln Pro Ala Val Glu Ser Gly Gln Val Asn Leu Ser Trp Thr Phe Val Gly Lys Lys Asp Gly Asp
Ala Tyr Leu Leu Ala Ile Glu Arg Asn Gly Asp Ile Val His Thr Thr Thr Ser Ile Gly Asp Ser Phe Thr
Asp Tyr Asp Val Glu Asn Gly Thr Glu Tyr Thr Tyr Val Val Lys Leu Tyr Asp Arg Ala Gly Asn
Val Val Ala Ser Asn Thr Val Lys Val Thr Pro Asp Ile Val Met Val Lys Val Ile Phe Lys Val Arg
Ala Pro Asp Tyr Thr Pro Leu Asp Ala Arg Ile Thr Ile Pro Asn Ser Leu Asn Gly Trp Asn Thr Gly
Ala Trp Glu Met Ser Arg Asn Gly Ala Val Thr Pro Asp Trp Gln Phe Thr Val Glu Val Gln Glu Gly
Glu Thr Ile Thr Tyr Lys Tyr Val Lys Gly Gly Ser Trp Asp Gln Glu Gly Leu Ala Asp His Thr Arg
Glu Asp Asp Asn Asp Asp Asp Val Ser Tyr Tyr Gly Tyr Gly Thr Ile Gly Thr Asp Leu Lys Val Thr
Val His Asn Glu Gly Asn Asn Thr Met Ile Val Gln Asp Arg Ile Leu Arg Trp Ile Asp Met Pro Val
Val Ile Glu Glu Val Gln Lys Gln Gly Ser Gln Val Thr Ile Lys Gly Asn Ala Ile Lys Asn Gly Val
Leu Thr Ile Asn Gly Glu Arg Val Pro Ile Asp Gly Arg Met Ala Phe Ser Tyr Thr Phe Ala Pro Ala
Ser His Gln Lys Glu Val Leu Ile His Ile Glu Pro Ser Ala Glu Ser Lys Thr Ala Ile Phe Asn Asn Asp

FIGURE 16HH

Gly Gly Ala Ile Ala Lys Asn Thr Lys Asp Tyr Val Leu Asn Leu Glu Thr Lys Gln Phe Lys Lys Leu
Leu Glu Ser Thr Ser Arg Ala Ala Ala Gly Pro Ser Ile Phe His Pro Gly Gly Val Pro Gly

SEQ ID NO: 91

gtgctaaccgtttaccgcacattcgaaggatggatgttcctgctcgcgtttttgctcactgcctcgtctgttcgcccaacaggacagcccgcca
aggtgcccgcaccgttaacggcaccatgatgcagtaatttgatgttgcggatgatggcacgttatggaccaaagtggccaatgaagc
caacaacttatccagccttgcatcaccgctctttggctgccgccgttataaaggaaacagccgcagcgacgtagggtacggagtatacga
ctgtatgacctcggcgaaatcaatcaaaaaggaccgtccgcacaaaatacggacaaaagctcaatcttcaagccattcaagccgccac
gccgctggaatgcaagtgtacgccgatgtcgtttcaccataaaggcggcgccgacggcacggaaatgggtggacgccgtcgaagtcaatc
cgctccgaccgcaaccaagaaatctcgggcacctatcaaatccaagcatggacgaaatttgatttccggggcggggaacacactactccagctt
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ctcatgaaagataaccgacattggcgtcaccttcgttgataatcatgacaccgaaccggccaagcgctgcagtcattggtgcacccatggt
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cgctgaaaagcaaaatcgatccgctcctcatcgccgcagggaattatgcttacggaacgcaacatgattatcttgactccgacatcaggg
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agtcaatggcggttcggttcggtttggttctagaaaacgaccgtctctaccatcgcttggccgacacaccegaccgtggactggtgaatt
cgctcgttgaccgaaccacggttggtggcatggccttga

SEQ ID NO: 92

Val Leu Thr Phe His Arg Ile Ile Arg Lys Gly Trp Met Phe Leu Leu Ala Phe Leu Leu Thr Ala Ser
Leu Phe Cys Pro Thr Gly Gln Pro Ala Lys Ala Ala Ala Pro Phe Asn Gly Thr Met Met Gln Tyr Phe
Glu Trp Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr Lys Val Ala Asn Glu Ala Asn Asn Leu Ser Ser
Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Thr Ser Arg Ser Asp Val Gly Tyr Gly
Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr
Lys Ala Gln Tyr Leu Gln Ala Ile Gln Ala Ala His Ala Ala Gly Met Gln Val Tyr Ala Asp Val Val
Phe Asp His Lys Gly Gly Ala Asp Gly Thr Glu Trp Val Asp Ala Val Glu Val Asn Pro Ser Asp Arg
Asn Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn
Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Val Asp Trp Asp Glu Ser Arg Lys Leu
Ser Arg Ile Tyr Lys Phe Arg Gly Ile Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn
Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn
Trp Gly Lys Trp Tyr Val Asn Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys
Phe Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Ser Gln Thr Gly Lys Pro Leu Phe Thr Val Gly
Glu Tyr Trp Ser Tyr Asp Ile Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Thr Met Ser Leu
Phe Asp Ala Pro Leu His Asn Lys Phe Tyr Thr Ala Ser Lys Ser Gly Gly Ala Phe Asp Met Arg Thr
Leu Met Thr Asn Thr Leu Met Lys Asp Gln Pro Thr Leu Ala Val Thr Phe Val Asp Asn His Asp
Thr Glu Pro Gly Gln Ala Leu Gln Ser Trp Val Asp Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile
Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Gln Tyr Asn Ile
Pro Ser Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln His
Asp Tyr Leu Asp His Ser Asp Ile Ile Gly Trp Thr Arg Glu Gly Val Thr Glu Lys Pro Gly Ser Gly
Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Gln His Ala Gly
Lys Val Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ser Asp Gly Trp Gly Glu
Phe Lys Val Asn Gly Gly Ser Val Ser Val Trp Val Pro Arg Lys Thr Thr Val Ser Thr Ile Ala Trp
Pro Ile Thr Thr Arg Pro Trp Thr Gly Glu Phe Val Arg Trp Thr Glu Pro Arg Leu Val Ala Trp Pro

FIGURE 16II

SEQ ID NO: 93

atgaaatcgtttgcaatcatgcctatccctttttatgcaaacgatttcatcagtgaaagggaggaggaggaatggggagaagaatgagaaga
 agattcagctattttcaatcttcttattgttcgttcagctgttttcatgttgcaaccgctagcgccaatggaacggtagaacagtagtccgttggttaa
 tggaaacgaagtcacgtttctatatggaggaaacaggaaacgagcagctctgttactggcaggctccttaatgattggcagaaagatggtagaca
 agaagattgcactaacaanaaggcgacaataacgtctgtctgtcacgcaaacacttcaagatgggacatatatgataagttgttgtagatggc
 aatgggtggcggatccgcttaacccgaatcaagtagacgacgggtacggcgccgtaatagtgctgtgtgctgggacaccgggtgcaacaag
 aacggacagtgacgctgttggttaacttacaagacgaattaggtcatagcagcgaatgggatccgaaagcgacagctacagtgatgaaaagg
 aagggaacgggttatatacgtttacaggtacacttccagccgggaacgtacgagtataaaatggcgaatattggcagctgggacgaaactatggt
 gtcggcggccgcgatggcggaatattaaagctgtatataaagaacaaacgggttactttattacaacgacagaacgcatgcgattgcgg
 attcgacttggtatgaccaattctaaaaaagcagccgcggctcgttggaacgattttaccagctattggtatgaacagacgtgaacgggt
 ggacgccgaaacatcaacggcgtgtgtcagatgatgatttgcattatcaggttaaggcgcgtgtgcaaaaggacatatgaatataa
 agtagttcttgggaatgattggacatatgaaattatccacaagataatgccaaataaattgtgtgaagaagacaaattaccttttttcaacgc
 gaaacgaagtagtgataaccgattacaatccaagcgggtcggatggtatcgtccaaaaagaccgttgaagcataatacgtgggattcgttgta
 tcgccaaaccgtttggtgcggtgaaagctgggacagaagtgaccttcgtttatcagcgaaaaaagggtgattgacaaaagcggatgtatatgtaa
 aaaatcagacaaccggcagcagaaactatctcagtaaaaaagcgggtgtcttggcgaagaagaatttgggaagcgacattcacaccgg
 atgtgaaaggagtatacgggtataaattattgcggtagatgctggaacgaaagcagaatacggggagatacacaagaaggcgagtgggga
 aaagcagtagataaaaatgcagagctgttccaattaacgggtgacgaccatcctaccaaacaccggattggatgaaagaagcagttgtatatca
 aatttccctgatccaaag

SEQ ID NO: 94

Met Lys Ser Phe Ala Phe Met Pro Ile Leu Phe Tyr Ala Asn Asp Phe Ile Ser Glu Arg Glu Gly Gly
 Gly Lys Met Gly Lys Asn Met Arg Arg Arg Phe Thr Tyr Phe Ser Ile Phe Leu Leu Phe Val Gln Leu
 Phe Ser Phe Ser Ala Thr Ala Ser Ala Asn Gly Thr Val Asn Ser Ser Pro Val Val Asn Gly Asn Glu
 Val Thr Phe Leu Tyr Gly Gly Thr Gly Asn Glu Gln Ser Val Leu Leu Ala Gly Ser Phe Asn Asp Trp
 Gln Lys Asp Gly Asp Lys Lys Ile Ala Leu Thr Lys Gly Asp Asn Asn Val Trp Ser Val Thr Gln Thr
 Leu Gln Asp Gly Thr Tyr Thr Tyr Lys Phe Val Val Asp Gly Gln Trp Val Ala Asp Pro Leu Asn Pro
 Asn Gln Val Asp Asp Gly Tyr Gly Gly Arg Asn Ser Val Val Val Gly Thr Pro Val Gln Gln Glu
 Arg Thr Val Thr Leu Val Gly Asn Leu Gln Asp Glu Leu Gly His Thr Ser Glu Trp Asp Pro Lys Ala
 Thr Ala Thr Val Met Lys Lys Glu Gly Asn Gly Leu Tyr Thr Phe Thr Gly Thr Leu Pro Ala Gly Thr
 Tyr Glu Tyr Lys Ile Ala Ile Asn Gly Ser Trp Asp Glu Asn Tyr Gly Val Gly Gly Arg Asp Gly Gly
 Asn Ile Lys Leu Leu Leu Asn Glu Gln Thr Thr Val Thr Phe Tyr Tyr Asn Asp Arg Thr His Ala Ile
 Ala Asp Ser Thr Trp Tyr Ala Pro Ile Leu Lys Glu Lys Gln Pro Arg Leu Val Gly Thr Ile Leu Pro
 Ala Ile Gly Tyr Glu Thr Asp Val Asn Gly Trp Thr Pro Gln Thr Ser Thr Ala Leu Leu Ser Asp Asp
 Asp Phe Asp Ser Ile Tyr Thr Phe Lys Ala Arg Val Pro Lys Gly Thr Tyr Glu Tyr Lys Val Val Leu
 Gly Asn Asp Trp Thr Tyr Glu Asn Tyr Pro Gln Asp Asn Ala Lys Leu Asn Val Leu Glu Glu Thr
 Thr Ile Thr Phe Phe Phe Asn Ala Lys Thr Lys Val Val Tyr Thr Asp Tyr Asn Pro Ser Gly Ser Asp
 Gly Ile Val Gln Lys Asp Arg Leu Lys His Asn Thr Trp Asp Ser Leu Tyr Arg Gln Pro Phe Gly Ala
 Val Lys Ala Gly Thr Glu Val Thr Leu Arg Leu Ser Ala Lys Lys Gly Asp Leu Thr Lys Ala Asp Val
 Tyr Val Lys Asn Thr Thr Thr Gly Thr Ala Lys Leu Tyr Ser Met Lys Lys Ala Gly Val Leu Gly Glu
 Glu Glu Tyr Trp Glu Ala Thr Phe Thr Pro Asp Val Lys Gly Val Tyr Gly Tyr Lys Phe Ile Ala Val
 Asp Ala Gly Thr Lys Ala Glu Tyr Gly Glu Asp Thr Gln Glu Gly Gln Trp Gly Lys Ala Val Asp Lys
 Asn Ala Glu Leu Phe Gln Leu Thr Val Tyr Asp Pro Ser Tyr Gln Thr Pro Asp Trp Met Lys Glu Ala
 Val Val Tyr Gln Ile Phe Pro Asp Pro Lys

SEQ ID NO: 95

atgtatacactatcatcgttcataatttgatactgatgggtgatgggttaggagactttagtggagttgctgaaaaggtagattatctaaatctcttg
 gagtagatacagctcgtttttaccatttaataaaagtaaatcttcatcagatgatgttggaagattactatgatgtaaccagattatggaacact
 acaagatcttgataatataaagttctaaatgaaatgaataaaggtagtaatggatcttgttgtaatcatacgtcggatacacatccatgggt
 tcttgatgcagttgaaaatactactaattctccatattggaactattacattatgagcttggatgagcctcaaaataagaatcatggcattataagggt
 aaticaaaaggacaaactgtgtgtgatttggattgtttgattcaatgccggaccttaattacgacaaacctaaagtaattggatgaagtgaanaa

FIGURE 16JJ

aataatagatttttgggcagatatgggagtagatggatttagatgacgcaaaacattatttgatttgactggagcgatggaattgaacag
tcagcaagcgttgcaaaagagatagaagactatataaaagataaactaggggaaaaatgcaatagttgtgagtgagggttacgatggagattcaa
atgtcttttaaaattgtcctcaatgcctgtgttaatttagtttatgtacaatttgagaggaaatttgaggagagataaactaattcagactctatt
agttgggttgattcctcgtgtataatttaaattgttttcattttccattatgtatgcatgactcttgacagatttattctgagctgtagatagtaaatatc
agggagatgtaatactgccacaaaacaataattgtctagtttaagctttactactctcattaacagccatgccaaactatttactatggtgatgaatag
gacttaggggatggaagtggcattcagaaccatgggatatcctgtgcgtgagccaatgcaatgggataaggatcaaaaagggaacgggtcaaa
cttattggacaaaagagttttacgaaggtaattactgaagggaagtgctaatgaagatggagcaatatacagatgatccagatgatggagtagtctgtag
aagaacaagaaaatggaattctattttaaaccttttaaaagaattatcaacttacgaaaagattatccggcactgtcttttgaagtactacgatga
gagagattggaaaaactgtatgttttgaagaaagctgtataactccaggatgtctgttattaataaccttgatccaacgtattcaaatatacagaa
gttcagaaagggtataaatgggtgtggtatgcaattttgatgggtgacaactatgaatttgagcaaaagatgaatgattttacagaatacaagtgtg
gacgataaatccaaggcaatttatattgttaaagtaa

SEQ ID NO: 96

Met Tyr Thr Leu Phe Ile Arg Ser Tyr Phe Asp Thr Asp Gly Asp Gly Val Gly Asp Phe Ser Gly Val
Ala Glu Lys Val Asp Tyr Leu Lys Ser Leu Gly Val Asp Thr Val Trp Phe Leu Pro Phe Asn Lys Ser
Lys Ser Tyr His Gly Tyr Asp Val Glu Asp Tyr Tyr Asp Val Glu Pro Asp Tyr Gly Thr Leu Gln Asp
Leu Asp Asn Met Ile Lys Val Leu Asn Glu Asn Gly Ile Lys Val Val Met Asp Leu Val Val Asn His
Thr Ser Asp Thr His Pro Trp Phe Leu Asp Ala Val Glu Asn Thr Thr Asn Ser Pro Tyr Trp Asn Tyr
Tyr Ile Met Ser Leu Asp Glu Pro Gln Asn Lys Asn His Trp His Tyr Lys Val Asn Ser Lys Gly Gln
Thr Val Trp Tyr Phe Gly Leu Phe Asp Ser Ser Met Pro Asp Leu Asn Tyr Asp Asn Pro Lys Val Met
Asp Glu Val Lys Lys Ile Ile Asp Phe Trp Ala Asp Met Gly Val Asp Gly Phe Arg Leu Asp Ala Ala
Lys His Tyr Tyr Gly Phe Asp Trp Ser Asp Gly Ile Glu Gln Ser Ala Ser Val Ala Lys Glu Ile Glu
Asp Tyr Ile Lys Asp Lys Leu Gly Glu Asn Ala Ile Val Val Ser Glu Val Tyr Asp Gly Asp Ser Asn
Val Leu Leu Lys Phe Ala Pro Met Pro Val Phe Asn Phe Ser Phe Met Tyr Asn Leu Arg Gly Asn
Phe Glu Gly Arg Asp Asn Leu Ile Ser Asp Ser Ile Ser Trp Val Asp Ser Ser Leu Tyr Asn Leu Asn
Val Phe His Phe Pro Phe Ile Asp Ser His Asp Leu Asp Arg Phe Ile Ser Glu Leu Val Asp Ser Lys
Tyr Gln Gly Asp Val Ile Ser Ala Thr Lys Gln Tyr Leu Leu Val Asn Ala Leu Leu Leu Ser Leu Thr
Gly Met Pro Thr Ile Tyr Tyr Gly Asp Glu Ile Gly Leu Arg Gly Trp Lys Trp His Ser Glu Pro Trp
Asp Ile Pro Val Arg Glu Pro Met Gln Trp Tyr Lys Asp Gln Lys Gly Asn Gly Gln Thr Tyr Trp Thr
Lys Glu Phe Tyr Glu Gly Ile Thr Glu Gly Ser Ala Asn Glu Asp Gly Ala Ile Tyr Asp Asp Pro Asp
Asp Gly Val Ser Val Glu Glu Gln Glu Asn Gly Tyr Ser Ile Leu Asn Phe Phe Lys Glu Phe Ile Asn
Leu Arg Lys Asp Tyr Pro Ala Leu Ala Phe Gly Ser Thr Thr Ile Glu Arg Asp Trp Lys Asn Leu Tyr
Val Leu Lys Lys Ser Tyr Asn Phe Gln Asp Val Leu Val Leu Ile Asn Leu Asp Pro Thr Tyr Ser Asn
Thr Tyr Glu Val Pro Glu Gly Tyr Lys Trp Val Trp Tyr Ala Phe Phe Asp Gly Asp Asn Tyr Glu Phe
Gly Ala Lys Asp Glu Met Ile Leu Gln Asn Thr Ser Trp Thr Ile Asn Pro Arg Gln Ile Tyr Ile Phe Val
Lys

SEQ ID NO: 97

atgaggaagaagatgtcgcattcaagatttactttttttgactttagcacttttttttctccgggtgtatttcagaagtttaaaagcgaagccag
ctactaaattcaaaagcgaaggtccttgtaaaagtaaatgttaatacgcatttattgagaatgctactactaatacgtggagtgttcaaaagaatct
tttattgattatcttagtaaaagtatttactgttaaggatgtaaatgatcagattgtatttactaaggaaacaacgaacaaacaaatattttttgaa
attgaacttcttctggaactttacatttgaggtaaaaggatagaggaagatttagttaattttcaggggaaaaagtttaacagatcatagatgag
aaaaataatattgttaattgtcgaactttttttgttaattggaalatgttaggacaaatgaagttgacgataattttataaaaaattatgataattacatgg
caacgttgatcttcaaaaaagatacagcacaagaagattatgaagaggtacctgtacacttacagggtacttccactttaattaataaagaattatat
cctggtagtggaactgttaaaattgaagttgatcttaaatcaaggatgcaagatgttaccagaaaaagttcatcttgaaaaatgaatttagcataga
agtgtccagcaaaagacaaaaagtttaacatttaattgtatgtctttgatacagaggttaataaccgaaatttagtagttgtatttccgcaattgagtt
gcctttgtggatcctgtacaaatttaagtggaagataaatgaattgaagggaatcttcaatgaattgggactattcagatccaatgcagaat
tttatgtgtataaagaattagaggaacaaggagaatattgtatgaatttggtaaaacacgcgagaaaagttatacaatagaattttccaag
caagaattcgataaatttagtggaatcgcttaattgtttatgccaacggtaaaagagagtgattagttgttctaaaaaaagaaattattaactata
gatttagaaggtgttgacagtaaaagtgctactataacgttgatcgaatgagcttaagttggattggaattataccaattcaaggttacttttgaag

FIGURE 16KK

tttgaaaaaagggtataaatagcaatgaatacgaataatttctcaactaacacaaaattcttttcaacagaattcacaggcaggcaattttgggac
 ttgagaaaattgcgattagagtagttgctaattggatttgaaagtaagattaatgagatttcaagagatgataactataacatcattgaatcttctct
 tacatcgctactatgtatacactattcaccgttcataatttgatactgatggatggtaggagacttttagtgaggttgctgaaaaggtagattatc
 taaaaictcttgaggtagatagacgtctggtttttaccatttaataaaagtaaatcttatcatggatatgatgtgaagattactatgtagaaccagat
 tatggaacactacaagatcttgataatgatataaaagttcnaatgaaaatggaataaaggtagaattggatcttggtttaatcatacgtcggatac
 acatccatggtttcttgatgcagttgaaaatactactaaattccatattggaactattacattatgagcttggatgagcctcaaaaataagaatcattgg
 cattataagggttaattcaaaaggacaaactgtgtggtattttggattgtttgattcaatgcccggacccttaattacgacaaccctaaagtaattggat
 gaagtgaaaaaataatagatttttgggcagatatgggagtagatggattagattatgcagcaaaacattattatggatttgactggagcgatg
 gaattgaacagtcagcaagcgttgcaaaagagatagaagactatataaagataaactaggggaaaatgcaatggttgagtgagggtttacga
 tggagattcaaatgttcttttaaaatttgcctcaatgctgtgttaattttatgttatacaatttgagaggaaaatttgaaggagagataacttaatt
 tcagactctattatggttggttgcctctgtgtataatttaaatgttttccatttccatttattgatgcatgcttgacagattatttctgagctttag
 atagtaaatatcagggagatgtaatactgccacaaaacaatttgcctgttaattgcttactactctcattaacaggcatgccaaactatttactatgg
 tgatgaataggacttaggggatggaagtggcattcagaaccatgggataacctgtgcgtgagccaatgcaatggtataaggatcaaaaagg
 gaacgggtcaaacctattggacaaaagagttttcgaaggtattactgaaggaggtgctaatgaagatggagcaatatacagatgccagatgatg
 gagtatctgtagaagaacaagaaaatggatattctatttaaaacttttttaagaattatcaacttacgaaaagattatccggcacttgcctttggaagt
 actacgattgagagagattggaaaaactgtatgtttgaaaaagtcgtataacttcaggatgttcttgatttaattaaccttgatccaacgtattcaa
 atacatacgaagttccagaagggtataaatgggtgtggtatgcatttttgatggtgacaactatgaatttggagcaaaagatgaaatgattttacag
 aatacaagttggacgataaatccaaggcaatttatataatttgaagtaa

SEQ ID NO: 98

Met Arg Lys Lys Met Ser His Ser Arg Phe Thr Phe Leu Leu Ile Leu Ala Leu Phe Ile Phe Phe Ser
 Gly Cys Ile Ser Glu Val Lys Ser Glu Ser Gln Leu Leu Asn Ser Lys Gln Lys Val Leu Val Lys Val
 Asn Val Asn Thr Pro Phe Ile Glu Asn Ala Thr Thr Asn Thr Trp Ser Val Ser Lys Glu Ser Phe Ile
 Asp Tyr Leu Ser Lys Val Ile Ile Thr Val Lys Asp Val Asn Asp Gln Ile Val Phe Thr Lys Glu Thr
 Thr Asn Lys Thr Asn Ile Tyr Phe Glu Ile Glu Leu Leu Pro Gly Thr Tyr Thr Phe Glu Val Lys Gly
 Tyr Glu Glu Asp Leu Val Ile Phe Ser Gly Glu Lys Val Asn Gln Ile Ile Asp Glu Lys Asn Asn Ile
 Val Asn Val Glu Thr Phe Phe Val Asn Gly Ile Val Arg Thr Ile Ile Glu Val Asp Asp Ile Ile Tyr Lys
 Asn Tyr Asp Ile Thr Ser Ala Thr Leu Ile Phe Lys Lys Asp Thr Ala Gln Glu Asp Tyr Glu Glu Val
 Pro Val Thr Leu Thr Gly Thr Ser Thr Leu Ile Asn Lys Glu Leu Tyr Pro Gly Met Trp Thr Val Lys
 Phe Glu Val Asp Leu Lys Ser Lys Asp Ala Ser Met Leu Pro Glu Lys Val His Leu Glu Asn Glu Phe
 Ser Ile Glu Val Leu Pro Ala Lys Thr Lys Ser Leu Thr Phe Asn Val Val Phe Asp Thr Glu Val Asn
 Glu Pro Lys Leu Val Val Val Phe Pro Gln Ile Glu Leu Pro Phe Val Asp Pro Val Thr Asn Leu Ser
 Gly Glu Ile Asn Glu Leu Glu Gly Asn Leu Ser Met Asn Trp Asp Tyr Ser Asp Pro Asn Ala Glu Phe
 Tyr Val Tyr Lys Glu Leu Glu Glu Gln Gly Glu Tyr Leu Tyr Glu Phe Val Gly Lys Thr Arg Glu Lys
 Ser Tyr Thr Ile Glu Asn Phe Thr Lys Gln Glu Phe Asp Lys Phe Ser Gly Ile Ala Ile Asn Val Tyr
 Ala Asn Gly Lys Glu Ser Gly Leu Val Val Leu Lys Lys Glu Asn Ile Lys Leu Ile Asp Leu Glu Ser
 Val Asp Ser Ile Ser Ala Thr Tyr Asn Val Asp Thr Asn Glu Leu Lys Leu Asp Trp Asn Tyr Thr Asn
 Ser Ser Val Thr Phe Glu Val Leu Lys Lys Gly Ile Asn Ser Asn Glu Tyr Glu Ile Ile Ser Gln Leu Thr
 Gln Asn Ser Phe Ser Thr Glu Phe Thr Gly Arg Gln Phe Trp Asp Leu Glu Lys Ile Ala Ile Arg Val
 Val Ala Asn Gly Phe Glu Ser Lys Ile Asn Glu Ile Ser Arg Asp Asp Ile Thr Ile Thr Ser Leu Asn Leu
 Pro Leu Thr Ser Ser Thr Met Tyr Thr Leu Phe Ile Arg Ser Tyr Phe Asp Thr Asp Gly Asp Gly Val
 Gly Asp Phe Ser Gly Val Ala Glu Lys Val Asp Tyr Leu Lys Ser Leu Gly Val Asp Thr Val Trp Phe
 Leu Pro Phe Asn Lys Ser Lys Ser Tyr His Gly Tyr Asp Val Glu Asp Tyr Tyr Asp Val Glu Pro Asp
 Tyr Gly Thr Leu Gln Asp Leu Asp Asn Met Ile Lys Val Leu Asn Glu Asn Gly Ile Lys Val Val Met
 Asp Leu Val Val Asn His Thr Ser Asp Thr His Pro Trp Phe Leu Asp Ala Val Glu Asn Thr Thr Asn
 Ser Pro Tyr Trp Asn Tyr Tyr Ile Met Ser Leu Asp Glu Pro Gln Asn Lys Asn His Trp His Tyr Lys
 Val Asn Ser Lys Gly Gln Thr Val Trp Tyr Phe Gly Leu Phe Asp Ser Ser Met Pro Asp Leu Asn Tyr
 Asp Asn Pro Lys Val Met Asp Glu Val Lys Lys Ile Ile Asp Phe Trp Ala Asp Met Gly Val Asp Gly
 Phe Arg Leu Asp Ala Ala Lys His Tyr Tyr Gly Phe Asp Trp Ser Asp Gly Ile Glu Gln Ser Ala Ser
 Val Ala Lys Glu Ile Glu Asp Tyr Ile Lys Asp Lys Leu Gly Glu Asn Ala Ile Val Val Ser Glu Val

FIGURE 16LL

Tyr Asp Gly Asp Ser Asn Val Leu Leu Lys Phe Ala Pro Met Pro Val Phe Asn Phe Ser Phe Met Tyr
 Asn Leu Arg Gly Asn Phe Glu Gly Arg Asp Asn Leu Ile Ser Asp Ser Ile Ser Trp Val Asp Ser Ser
 Leu Tyr Asn Leu Asn Val Phe His Phe Pro Phe Ile Asp Ser His Asp Leu Asp Arg Phe Ile Ser Glu
 Leu Val Asp Ser Lys Tyr Gln Gly Asp Val Ile Ser Ala Thr Lys Gln Tyr Leu Leu Val Asn Ala Leu
 Leu Leu Ser Leu Thr Gly Met Pro Thr Ile Tyr Tyr Gly Asp Glu Ile Gly Leu Arg Gly Trp Lys Trp
 His Ser Glu Pro Trp Asp Ile Pro Val Arg Glu Pro Met Gln Trp Tyr Lys Asp Gln Lys Gly Asn Gly
 Gln Thr Tyr Trp Thr Lys Glu Phe Tyr Glu Gly Ile Thr Glu Gly Ser Ala Asn Glu Asp Gly Ala Ile
 Tyr Asp Asp Pro Asp Asp Gly Val Ser Val Glu Glu Gln Glu Asn Gly Tyr Ser Ile Leu Asn Phe Phe
 Lys Glu Phe Ile Asn Leu Arg Lys Asp Tyr Pro Ala Leu Ala Phe Gly Ser Thr Thr Ile Glu Arg Asp
 Trp Lys Asn Leu Tyr Val Leu Lys Lys Ser Tyr Asn Phe Gln Asp Val Leu Val Leu Ile Asn Leu Asp
 Pro Thr Tyr Ser Asn Thr Tyr Glu Val Pro Glu Gly Tyr Lys Trp Val Trp Tyr Ala Phe Phe Asp Gly
 Asp Asn Tyr Glu Phe Gly Ala Lys Asp Glu Met Ile Leu Gln Asn Thr Ser Trp Thr Ile Asn Pro Arg
 Gln Ile Tyr Ile Phe Val Lys

SEQ ID NO: 99

atgtacacactcttcacccgctctttttacgatacaaaacacgacgggtgtaggtgactacaacgggttgcccaaaaagtagactatctcaaaacg
 ctggagtggtacacagtttggttctgccgttcaacaaagcaaaatcgaccacgggtacgatgtgaagactactacgatgtagaacctgactatg
 gaacatacgcacaacttgaaaatagataaagacactcaatcagaacggaattcgtgtgtatggacttggttgtaaccacactccgatacac
 actcgtggtttctggatgccgttgagaacacaacgaattcgaaatattggagctactacataatgacacttgaaaatagagacgggtggaatcact
 ggcatggagaataaactcaaaaggcaaaaagtttactacttcggactggttgactcatcaatgcccgaatttgacaatccacaagtgtat
 gaacgaaatcaagagaataaactgatttctggataacagttgggtggtggttgcagacttgatgcaccaaagcactacaaggctgggattggg
 acgacggcatttcaggttcagcagcaatcgcgagggaatagaaggtacacaggagcaagtaggaacgatgcgtagtgtgctggggaa
 gtgtacgatggaaatccatcggttcttcaaatggcaccgatgccggcgttcaattcacattcatgtatggaataacaggcaacctgagggg
 aaagataacctgctgggagaacaatttcattggttaattggagcgagttattatcacaacgtaaaacattccccgttcatagacaatcacgattga
 acagatggatcgcatacttaccgacaaaagtatagtggaacacacaagttggtacgaagcagtagtatatttaacaaatgcgctcttcttcctta
 aacggtagtgcctgtatttattatgggaatgaataggcttgagaggatggaatggggacaagaccctgggattgcccgggtgagagagccga
 tgcagtgtgacgcaagtcgaagtgagctgggcagacatggggacaaaagccctgtctaccagcaaaaaggaatcacatttggaatgcaaac
 gtcgatgtgtcgcgatgtacgatgaccaaagtgatggggttcagtagaagagcagatgaatggttacacgataataaacttcttaacaattcataa
 ccctgaggaagacatatccggctctatcgaagggttcgataacgatagaacgcgactggaagaacctgtacgttaacaaacgagctacggaa
 atcaggaagtgtgtattgataaacttagcccaacttgccggaacaattacacgttaccaggttgatacaggtgggtcgtgtatgcgttcttaa
 tgggagttgttgatttgcaataaaacgaatcaccactgagcgaagataccaactggacagtcgaatccaaggcaagtgatgtgtgttgaa
 ggactaa

SEQ ID NO: 100

Met Tyr Thr Leu Phe Ile Arg Ser Phe Tyr Asp Thr Asn Asn Asp Gly Val Gly Asp Tyr Asn Gly Val
 Ala Gln Lys Val Asp Tyr Leu Lys Thr Leu Gly Val Asp Thr Val Trp Phe Leu Pro Phe Asn Lys Ala
 Lys Ser Tyr His Gly Tyr Asp Val Glu Asp Tyr Thr Asp Val Glu Pro Asp Tyr Gly Thr Tyr Ala Gln
 Leu Glu Asn Met Ile Lys Thr Leu Asn Gln Asn Gly Ile Arg Val Val Met Asp Leu Val Val Asn His
 Thr Ser Asp Thr His Ser Trp Phe Leu Asp Ala Val Glu Asn Thr Thr Asn Ser Lys Tyr Trp Ser Tyr
 Tyr Ile Met Thr Leu Glu Asn Arg Asp Gly Trp Asn His Trp His Trp Lys Ile Asn Ser Lys Gly Gln
 Lys Val Tyr Tyr Phe Gly Leu Phe Asp Ser Ser Met Pro Asp Leu Asn Phe Asp Asn Pro Gln Val
 Met Asn Glu Ile Lys Arg Ile Ile Asp Phe Trp Ile Thr Val Gly Val Asp Gly Phe Arg Leu Asp Ala
 Pro Lys His Tyr Lys Gly Trp Asp Trp Asp Asp Gly Ile Ser Gly Ser Ala Ala Ile Ala Arg Glu Ile Glu
 Ser Tyr Ile Arg Ser Lys Leu Gly Asn Asp Ala Ile Val Val Gly Glu Val Tyr Asp Gly Asn Pro Ser
 Val Leu Ser Gln Phe Ala Pro Met Pro Ala Phe Asn Phe Thr Phe Met Tyr Gly Ile Thr Gly Asn His
 Glu Gly Lys Asp Asn Leu Leu Gly Glu Thr Ile Ser Trp Val Asn Gly Ala Ser Tyr Tyr Leu Asn Val
 Lys His Phe Pro Phe Ile Asp Asn His Asp Leu Asn Arg Trp Ile Ser Ile Leu Ile Asp Gln Lys Tyr Ser
 Gly Asn Thr Gln Val Gly Thr Lys Gln Tyr Ile Leu Thr Asn Ala Leu Leu Leu Ser Leu Asn Gly Met
 Pro Val Ile Tyr Tyr Gly Asn Glu Ile Gly Leu Arg Gly Trp Lys Trp Gly Gln Asp Pro Trp Asp Leu
 Pro Val Arg Glu Pro Met Gln Trp Tyr Ala Ser Gln Ser Gly Ala Gly Gln Thr Trp Trp Thr Lys Pro

FIGURE 16MM

Val Tyr Gln Gln Lys Gly Ile Thr Phe Gly Asn Ala Asn Val Asp Gly Ala Met Tyr Asp Asp Pro Asn
 Asp Gly Val Ser Val Glu Glu Gln Met Asn Gly Tyr Thr Ile Asn Asn Phe Phe Lys Gln Phe Ile Thr
 Leu Arg Lys Thr Tyr Pro Ala Leu Ser Lys Gly Ser Ile Thr Ile Glu Arg Asp Trp Lys Asn Leu Tyr
 Val Ile Lys Arg Val Tyr Gly Asn Gln Glu Val Leu Val Leu Ile Asn Leu Asp Pro Thr Trp Pro Asn
 Asn Tyr Thr Leu Pro Gly Gly Tyr Arg Trp Val Trp Tyr Ala Phe Phe Asn Gly Ser Leu Phe Glu Phe
 Gly Asn Lys Asn Glu Ser Pro Leu Ser Gln Asp Thr Asn Trp Thr Val Asn Pro Arg Gln Val Tyr Val
 Phe Val Lys Asp

SEQ ID NO: 101

ttgcgattcttccaaagtaataatccccctttccgcaaaacaccagagagtgccagcgaagcgcagtaicaagagacactgaacaattacaag
 gaaagtaataatgatcaatttgaaaaaacaccattagcgccctggcgaggtatggtattaggctttgcatccaacgcaatggcgggtccctag
 aaccgctttgtacacctcttgaaatggaaatgggaagatgttgacaggagtgtaaacatttctcgacctaaggctttgccgagtgcaagt
 ctctccgccaactaaatctcacaacacggatgcatgtggggccggtatcaacccgttagttatgctttgaaggacgcagcggtaatgcagcc
 aatttaaaataggtgcaacgttgtaagcigtaggcgtcgaatatactagatgcagtgataaaccacatggcagcctacgacagaatttcc
 ctgatgtaccctatagcagtaatgactttaactctgtacaggagatattgactataataaccgttgcaaacacagcattgtgatttagtcggctta
 atgatctaaaaacaggatctgactacgtccgcaaaaaatagcggatataatgaacgacgcaatcagtagtggtgtagctggttccgtattgatg
 cagccaaacataataccagcaggtgatatagtgccattaaaggtaaaatgaatccatacatcttcaagaggtaattggtgcatccggcg
 aacctgttcgaccgactgaatacaccittatcggtggtgtcacggaattcaatttgctcgaattgggtccagcctccgcaatagtaattgctt
 gggttaaaagacattggcagtcgaatgaattatccagtgtgatgccgaacattgtaacgaatcatgatgaagcgcataacccgaatggtc
 ctatttggcacggcgticaaggtaattggtatgcattagcaaatattttaccttagcttaccctacggctatccaaaatcatgacagatactt
 ccacgggtgactttaacgcagctccaccaagcagtggtatacacacaggaatgcgtgtggtttgatggcggagactgggtatgcgaacacaa
 atggcgcggtattgtaacatggttgccttccgcaactatacagaacgcaatggcgatcagtaattggtggcaaacagtaacgaccaatg
 ctttggcgcgggtggttaggtttgtgttattataaacgtgctaattggtgacataacaaatgttgatacgggaatgctgtgccaatctgt
 aacataatagaagctaactttgatgaagcaccggccaatgtagtgcagctacagattccaacggtaagccgttattaccgtagtggtggca
 agctaactttaatgtagcaggcgatcatgtgtgcaattcatgttgccgcaaaatgggtatcaatgtagtggtgatgattgccatgtacagga
 tccgattgtaataatgatccfaaacctgattttgagtagcagcaacatcaattgtacatcagaaaattacctaagctatattactggggagcaca
 gccacagatagcttagcgaatgcagcttgccaggtgtcgaatgcaaacaaatggcgacttaagtgtcatgatttaggtgtcgaactaacca
 aaattaacgccatcttagtgacaatgggtgcaataaaacagctgatctaactgttactggtgcaggtgtgataaagacgggacttggagcacctt
 acaaaattgtgcttgaattaccgggtgcacaaccaatccagctgggtggcgacgaagcttggtacttccgaggtactgtaataactggtgta
 aagcacaattagattatgacgcaactagcgggtgtattacacaatacaaaagcttaattggtgaagagcacctgcccgttttaaaatgataatggt
 agttggactgaagctatccaacagctgattaccaagttacagataacaattacacgcattaacttaatagcgatagcgaagcgattacagtaa
 accgcacaataa

SEQ ID NO: 102

Met Arg Phe Phe Pro Lys Leu Ile Ser Pro Phe Pro Gln Asn Thr Arg Glu Trp Gln Arg Ser Ala Val
 Ser Arg Asp Thr Glu Gln Leu Gln Arg Lys Val Ile Met Ile Asn Leu Lys Lys Asn Thr Ile Ser Ala
 Leu Val Ala Gly Met Val Leu Gly Phe Ala Ser Asn Ala Met Ala Val Pro Arg Thr Ala Phe Val His
 Leu Phe Glu Trp Lys Trp Glu Asp Val Ala Gln Glu Cys Glu Thr Phe Leu Gly Pro Lys Gly Phe Ala
 Ala Val Gln Val Ser Pro Pro Thr Lys Ser His Asn Thr Asp Ala Trp Trp Gly Arg Tyr Gln Pro Val
 Ser Tyr Ala Phe Glu Gly Arg Ser Gly Asn Arg Ser Gln Phe Lys Asn Met Val Gln Arg Cys Lys Ala
 Val Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn His Met Ala Ala Tyr Asp Arg Asn Phe Pro Asp
 Val Pro Tyr Ser Ser Asn Asp Phe Asn Ser Cys Thr Gly Asp Ile Asp Tyr Asn Asn Arg Trp Gln Thr
 Gln His Cys Asp Leu Val Gly Leu Asn Asp Leu Lys Thr Gly Ser Asp Tyr Val Arg Gln Lys Ile Ala
 Asp Tyr Met Asn Asp Ala Ile Ser Met Gly Val Ala Gly Phe Arg Ile Asp Ala Ala Lys His Ile Pro
 Ala Gly Asp Ile Ala Ala Ile Lys Gly Lys Leu Asn Gly Asn Pro Tyr Ile Phe Gln Glu Val Ile Gly Ala
 Ser Gly Glu Pro Val Arg Pro Thr Glu Tyr Thr Phe Ile Gly Gly Val Thr Glu Phe Gln Phe Ala Arg
 Lys Leu Gly Pro Ala Phe Arg Asn Ser Asn Ile Ala Trp Leu Lys Asp Ile Gly Ser Gln Met Glu Leu
 Ser Ser Ala Asp Ala Val Thr Phe Val Thr Asn His Asp Glu Glu Arg His Asn Pro Asn Gly Pro Ile
 Trp His Gly Val Gln Gly Asn Gly Tyr Ala Leu Ala Asn Ile Phe Thr Leu Ala Tyr Pro Tyr Gly Tyr
 Pro Lys Ile Met Ser Gly Tyr Phe Phe His Gly Asp Phe Asn Ala Ala Pro Pro Ser Ser Gly Ile His Thr

FIGURE 16NN

Gly Asn Ala Cys Gly Phe Asp Gly Gly Asp Trp Val Cys Glu His Lys Trp Arg Gly Ile Ala Asn Met
 Val Ala Phe Arg Asn Tyr Thr Ala Ser Glu Trp Arg Ile Ser Asn Trp Trp Gln Asn Ser Asn Asp Gln
 Ile Ala Phe Gly Arg Gly Gly Leu Gly Phe Val Val Ile Asn Lys Arg Ala Asn Gly Ser Ile Asn Gln
 Ser Phe Asp Thr Gly Met Pro Asp Gly Gln Tyr Cys Asn Ile Ile Glu Ala Asn Phe Asp Glu Ser Thr
 Gly Gln Cys Ser Ala Ala Thr Asp Ser Asn Gly Gln Ala Val Ile Thr Val Ser Gly Gly Gln Ala Asn
 Phe Asn Val Ala Gly Asp His Ala Ala Ala Ile His Val Gly Ala Lys Ile Gly Asp Gln Cys Ser Gly
 Asp Asp Cys Pro Cys Thr Gly Ser Asp Cys Asn Asn Asp Pro Lys Pro Asp Phe Ala Val Pro Ala
 Thr Ser Ile Cys Thr Ser Glu Asn Leu Pro Thr Leu Tyr Tyr Trp Gly Ala Gln Pro Thr Asp Ser Leu
 Ala Asn Ala Ala Trp Pro Gly Val Ala Met Gln Thr Asn Gly Asp Phe Lys Cys His Asp Leu Gly Val
 Glu Leu Thr Lys Ile Asn Ala Ile Phe Ser Asp Asn Gly Ala Asn Lys Thr Ala Asp Leu Thr Val Thr
 Gly Ala Gly Cys Tyr Lys Asp Gly Thr Trp Ser Thr Leu Gln Asn Cys Gly Phe Glu Ile Thr Gly Ala
 Gln Thr Asn Pro Val Gly Gly Asp Glu Val Trp Tyr Phe Arg Gly Thr Ala Asn Asp Trp Gly Lys Ala
 Gln Leu Asp Tyr Asp Ala Thr Ser Gly Leu Tyr Tyr Thr Ile Gln Ser Phe Asn Gly Glu Glu Ala Pro
 Ala Arg Phe Lys Ile Asp Asn Gly Ser Trp Thr Glu Ala Tyr Pro Thr Ala Asp Tyr Gln Val Thr Asp
 Asn Asn Ser Tyr Arg Ile Asn Phe Asn Ser Asp Ser Lys Ala Ile Thr Val Asn Ala Gln

SEQ ID NO: 103

gtgctaacggttcaccgcacatcgcgaaaggatggatgttctgctgcggttttgcactgcctgcgtgttctgcccaacaggacagcccgcca
 aggcgtcccgaccgtttaacggcaccatgatgcagttttgaatggctacttgcggatgatggcagcttatggaccaaagtggccaatgaagc
 caacaactatccagccttggcatcaccgctcttggctgcgcccgttacaaaggacaagccgcagcgacgtagggtacggagtatacga
 ctgtatgacctggcggaattcaatcaaaaaggacgcgcacaaaatacggacaacaaagctcaatatctcaagccattcaagccggccac
 gccgctggaatgcaagtgtacgcgatgtcgtgttcgaccataaaggcgccgacggcacggcaatgggtggacgccgtcgagtaac
 cgccgacgcgaaccaagaatcgcggcacctatcaaatcaagcatggacgaaattgatttccggcggggcaaacctactccagctt
 taagtggcgctggtaccattttgacggcgttgattgggacgaaagccgaaattgagccgattacaattccggcgatcggaagcggtg
 gattgggaagtacacggaacggaactatgactacttaattgatgacccgtgatgatgatccgaagtcgtgacccgagctgaaa
 actggggggaatggtatgtcaacacacgaacattgaggttccggctgatgcctgaagcatattaagttcagtttttctgattggtgctg
 atgtgcgttctcagactggcaagccgctattaccgctgggaatattggagctatgacatcaacaagttgcacaattacattacgaaacaaacg
 gaacgatgtcttggatgacccgttacacacaaatttataccgcttcacaaatcaggggcgccatttgatgacgcaggttaattgaccaatct
 ctcatgaaagatcaaccgacattggccgtcacccttctgataatcatgacaccgaaccggccaagcgctgcagtcacgtgacccatggt
 tcaaacggttggttacgccttatttaactcggcaggaaggatacccgctgcgttttatggtgactattatggcatccacaatataacatccttc
 gctgaaaagcaaaatcgatccgctcctcatcgcgcgaggggattatgcttacggaacgaacatgattatcttgatcactccgacatcatcggt
 ggacaagggaagggtcactgaaaacacaggatccgggctggccgactgatcacogattggccgggaggaagcaaatggatgtacgttg
 gcaaacacacgctggaagaagtgtctatgacctaccggcaaccggagtgacaccgtcacatcaacagtgatggatggggggaattcaaa
 gtcaatggcggttcggttgcgttgcctagaaaacgacccgttctaccatcgctcgccgatcacaaccggaccgtgactggtgaattc
 gtccgttgaccgaaccacggttggtggcatggcctga

SEQ ID NO: 104

Val Leu Thr Phe His Arg Ile Ile Arg Lys Gly Trp Met Phe Leu Leu Ala Phe Leu Leu Thr Ala Ser
 Leu Phe Cys Pro Thr Gly Gln Pro Ala Lys Ala Ala Ala Pro Phe Asn Gly Thr Met Met Gln Tyr Phe
 Glu Trp Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr Lys Val Ala Asn Glu Ala Asn Asn Leu Ser Ser
 Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Thr Ser Arg Ser Asp Val Gly Tyr Gly
 Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr
 Lys Ala Gln Tyr Leu Gln Ala Ile Gln Ala Ala His Ala Ala Gly Met Gln Val Tyr Ala Asp Val Val
 Phe Asp His Lys Gly Gly Ala Asp Gly Thr Glu Trp Val Asp Ala Val Glu Val Asn Pro Ser Asp Arg
 Asn Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn
 Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Val Asp Trp Asp Glu Ser Arg Lys Leu
 Ser Arg Ile Tyr Lys Phe Arg Gly Ile Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn
 Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn
 Trp Gly Glu Trp Tyr Val Asn Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys
 Phe Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Ser Gln Thr Gly Lys Pro Leu Phe Thr Val Gly

FIGURE 1600

Glu Tyr Trp Ser Tyr Asp Ile Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Thr Met Ser Leu
Phe Asp Ala Pro Leu His Asn Lys Phe Tyr Thr Ala Ser Lys Ser Gly Gly Ala Phe Asp Met Arg Thr
Leu Met Thr Asn Thr Leu Met Lys Asp Gln Pro Thr Leu Ala Val Thr Phe Val Asp Asn His Asp
Thr Glu Pro Gly Gln Ala Leu Gln Ser Trp Val Asp Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile
Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Gln Tyr Asn Ile
Pro Ser Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln His
Asp Tyr Leu Asp His Ser Asp Ile Ile Gly Trp Thr Arg Glu Gly Val Thr Glu Lys Pro Gly Ser Gly
Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Gln His Ala Gly
Lys Val Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ser Asp Gly Trp Gly Glu
Phe Lys Val Asn Gly Gly Ser Val Ser Val Trp Val Pro Arg Lys Thr Thr Val Ser Thr Ile Ala Arg
Pro Ile Thr Thr Arg Pro Trp Thr Gly Glu Phe Val Arg Trp Thr Glu Pro Arg Leu Val Ala Trp Pro

SEQ ID NO: 105

atgtccctatttcaaaaaatctttccgtggattgtatctctacttctttgttttcgtttatgtctctttttccattcaaacgaaaaagtcgcgctggaa
gtgttccagtgatggaacgatgatgcaaatattcgaatggtaacctccagacgaatggaacactatggacgaaagtagcaaataacgcccaatct
ttaagcaatcttggcattactgccctttggcttccccctgcctataaaggaaacaaagcagcagtgacgttggatatggcgttatgattatagacct
aggagagttaatacaaaaaggaactgtccgaacaaaatacggaaacaaaacacaatatatccaagcaatccaagcggcgcatacagcaggaa
tgcaagtatatgcagatgtcgtcttaaccataaaggcgggtgcagatgggacagaactagtggatgcagtagaagtaaacccttctgaccgcaat
caagaaatatcaggaacataatcaaatccaagcgtggacaaaattgtatttcttggctgtggaaacacatttctagttaaaatggcgttggtatca
ttcgttggaacggactgggatgagagtagaaaactaaatcgtatttacaattccgcggcacgggaaaagcatgggattgggaagttagataca
gaaatgggaattatgactatctcatgtatgcagatttggatatggatcatccagaggttgaatcgaactaaaaaattggggaaaagtggatatgtaa
ccacaaccaatatcgacggattccgcttggatgcagtgagcaataataatagctttttcccgacttggctatcgtatgacgaacccaaacac
aaaagcctcttttggcgttggcgaatttggagcattgacattaacaagctacacaactatattacaaagacgaacggctctatgtccctattcga
gccccgctgcataacaattttatatagcatcgaatacaggtggctatttggatatgcgacattactcaacaacacattgatgaaatgaacccaa
cactatcgggtacattagtagacaatcacgatactgagccagggaacttttcagtcgtgggtcgaagccgttggttaaaccgttagcttacgcat
ttatcttgaccgccaagaaggttatccgttgcatcttttaaggagattactatggtatttccaaaatacaacattcctgcgctgaaaagcgaacttgatc
cgctgttaattgtcgaagagattatgcttacggaacacagcagactattgacaaatgcagataattatcggttggacgcgggaaggagtagct
gaaaagcgaattcgggacttgcctactattaccgacggacctggcggaagcgaatggatgtatgttggcaacaacacgctggcaaac
gtttatgactaaccggcaatcgaagtgtacagtgacaatcaacgcgatgttgatggggaagaatttaaagtcgaatggaggtctgtatccatg
ggttccaaaaacatcaaccacttcccaatcacaatttactgtaaataatgccacaaccgttgggggacaaaatgtatacgttgcgggaattttcg
cagctgggcaac

SEQ ID NO: 106

Met Ser Leu Phe Lys Lys Ile Phe Pro Trp Ile Val Ser Leu Leu Leu Leu Phe Ser Phe Ile Ala Pro Phe
Ser Ile Gln Thr Glu Lys Val Arg Ala Gly Ser Val Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu
Trp Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr Lys Val Ala Asn Asn Ala Gln Ser Leu Ala Asn Leu
Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Thr Ser Ser Ser Asp Val Gly Tyr Gly Val
Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys
Thr Gln Tyr Ile Gln Ala Ile Gln Ala Ala His Thr Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe
Asn His Lys Ala Gly Ala Asp Gly Thr Glu Leu Val Asp Ala Val Glu Val Asn Pro Ser Asp Arg Asn
Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr
Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn
Arg Ile Tyr Lys Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr
Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp His Pro Glu Val Val Ser Glu Leu Lys Asn Trp
Gly Lys Trp Tyr Val Thr Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr
Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Thr Gln Thr Gln Lys Pro Leu Phe Ala Val Gly Glu
Phe Trp Ser Tyr Asp Ile Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu Phe
Asp Ala Pro Leu His Asn Asn Phe Tyr Ile Ala Ser Lys Ser Gly Gly Tyr Phe Asp Met Arg Thr Leu
Leu Asn Asn Thr Leu Met Lys Asp Gln Pro Thr Leu Ser Val Thr Leu Val Asp Asn His Asp Thr

FIGURE 16PP

Glu Pro Gly Gln Ser Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu
 Thr Arg Gln Glu Gly Tyr Pro Cys Ile Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro
 Ala Leu Lys Ser Lys Leu Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln His Asp
 Tyr Ile Asp Asn Ala Asp Ile Ile Gly Trp Thr Arg Glu Gly Val Ala Glu Lys Ala Asn Ser Gly Leu
 Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Gln His Ala Gly Lys
 Thr Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ala Asp Gly Trp Gly Glu Phe
 Lys Val Asn Gly Gly Ser Val Ser Ile Trp Val Pro Lys Thr Ser Thr Thr Ser Gln Ile Thr Phe Thr Val
 Asn Asn Ala Thr Thr Val Trp Gly Gln Asn Val Tyr Val Val Gly Asn Ile Ser Gln Leu Gly Asn

SEQ ID NO: 107

atggacagcctcgacgcgcccggagcagaagccctgggtgaaggatggcaggctctccgctacctggatacaggacaggaccgtggc
 gctcccaggacacctgcgccccgccgccccggccgaggaagtcggcccggtggacaagtggaaaaacgatatcatctattcgtctcac
 cgaccgtttccaggatggcgacaagaccaacaacatggacgtggtcccgacggacatgaaaaatatcatggcggcgacatccaggggctc
 atcgacaagctcgactatatcaaggagacgggttcgacggccatctggtctacgccccctatgaaggggcagacccacttcttcgagaccgac
 aattaccatggttactggcccattgacttctatgacacggacccccatgtgggacacatgcagaaatttgaggagcttatcgagaaagcccatga
 gaaagggctgaagatcgctgctgatatccccctgaaccacacggcctgggagcatcccttctacaaggacgacagcaagaaggactggttcc
 accatataggagatgtgaaggactgggaagatccctactgggtgaaacggctccatattcgttctcctgacctggcgaggaaaaccctg
 ccgtgaaaaagtacctcatcgacgtggccaagttctgggtagacaagggtattgacggcttcaggcttgacggcgtgaagaacgtgccccctca
 acttctgggcgaagttgaccggcgattcacgattatggggcaaggacttctctcgtcggggaatactttgacggaaaccggcgaaagt
 cgcgaactaccagagagaggacatgagctcactcttcgattaccgcctactggacctgaaggacaccttcgccaaggacgggagcatgc
 gcaacctggcggcgaagcttgatgagtgacaggaattatcccgacccgggctcatgtcggtttcttctgataaccacgacacggcgaggtt
 cctcaccgaggccaacggcaacaaggataagctcaaaactggccctcgccctcgcgatgacctcaaccgcacgtgctaccatttattatggcacc
 gaggttgccatggaaggcaactgcgatacatggcgccgtagataaccggaggacatgcagtgggacaaggatcctgacatgttcaata
 cttaagactctcaccactgccgcaatgagcatgaatccctcagggaaggaaagaagctcgagatgtggcaggatgacaagctacgcgta
 cgggaggcagaccccgaaggacgagtcctatcggtgcttaacaacggctatgatacgaggaacgggacataccgctccgcccgagag
 cggcatcaagaacggcagcggtgctgaaggatgtcatcaccggcgaaaccgtgacggtacagaacggaaaaatccatgcgaaatgcggcgg
 caaacaggcgcggatctacgtgccccgctag

SEQ ID NO: 108

Met Asp Ser Leu Asp Ala Pro Glu Gln Lys Pro Trp Val Lys Asp Gly Arg Leu Ser Ala Tyr Leu Asp
 Thr Gly Thr Gly Thr Val Val Ala Pro Glu Ala Pro Ala Pro Pro Pro Ala Glu Glu Val Arg
 Pro Val Asp Lys Trp Lys Asn Asp Ile Ile Tyr Phe Val Leu Thr Asp Arg Phe Gln Asp Gly Asp Lys
 Thr Asn Asn Met Asp Val Val Pro Thr Asp Met Lys Lys Tyr His Gly Gly Asp Ile Gln Gly Leu Ile
 Asp Lys Leu Asp Tyr Ile Lys Glu Thr Gly Ser Thr Ala Ile Trp Leu Thr Pro Pro Met Lys Gly Gln
 Thr His Phe Phe Glu Thr Asp Asn Tyr His Gly Tyr Trp Pro Ile Asp Phe Tyr Asp Thr Asp Pro His
 Val Gly Thr Met Gln Lys Phe Glu Glu Leu Ile Glu Lys Ala His Glu Lys Gly Leu Lys Ile Val Leu
 Asp Ile Pro Leu Asn His Thr Ala Trp Glu His Pro Phe Tyr Lys Asp Asp Ser Lys Lys Asp Trp Phe
 His His Ile Gly Asp Val Lys Asp Trp Glu Asp Pro Tyr Trp Ala Glu Asn Gly Ser Ile Phe Gly Leu
 Pro Asp Leu Ala Gln Glu Asn Pro Ala Val Glu Lys Tyr Leu Ile Asp Val Ala Lys Phe Trp Val Asp
 Lys Gly Ile Asp Gly Phe Arg Leu Asp Ala Val Lys Asn Val Pro Leu Asn Phe Trp Ala Lys Phe Asp
 Arg Ala Ile His Asp Tyr Ala Gly Lys Asp Phe Leu Leu Val Gly Glu Tyr Phe Asp Gly Asn Pro Ala
 Lys Val Ala Asn Tyr Gln Arg Glu Asp Met Ser Ser Leu Phe Asp Tyr Pro Leu Tyr Trp Thr Leu Lys
 Asp Thr Phe Ala Lys Asp Gly Ser Met Arg Asn Leu Ala Ala Lys Leu Asp Glu Cys Asp Arg Asn
 Tyr Pro Asp Pro Gly Leu Met Ser Val Phe Leu Asp Asn His Asp Thr Pro Arg Phe Leu Thr Glu Ala
 Asn Gly Asn Lys Asp Lys Leu Lys Leu Ala Leu Ala Phe Ala Met Thr Ile Asn Arg Met Pro Thr Ile
 Tyr Tyr Gly Thr Glu Val Ala Met Glu Gly Asn Cys Asp Ile Met Gly Ala Val Asp Asn Arg Arg
 Asp Met Gln Trp Asp Lys Asp Pro Asp Met Phe Lys Tyr Phe Lys Thr Leu Thr Thr Ala Arg Asn
 Glu His Glu Ser Leu Arg Glu Gly Lys Lys Leu Glu Met Trp Gln Asp Asp Lys Val Tyr Ala Tyr Gly
 Arg Gln Thr Pro Lys Asp Glu Ser Ile Val Val Leu Asn Asn Gly Tyr Asp Thr Gln Glu Arg Asp Ile

FIGURE 16QQ

Pro Leu Arg Pro Glu Ser Gly Ile Lys Asn Gly Thr Val Leu Lys Asp Val Ile Thr Gly Glu Thr Val
Thr Val Gln Asn Gly Lys Ile His Ala Lys Cys Gly Gly Lys Gln Ala Arg Ile Tyr Val Pro Ala

SEQ ID NO: 109

atggcaagaaaacgctggccataatcttctgacttctagtgcttcttagtctctcggcagttccggcaaggcagaaactctagagaatggtgga
gttataatgcaggcttcttattgggaatgttctggaggaggaatctggtgggacacaaatagctcaaaagataccccaatgggcaagtgcaggaa
ctcagcgatatgattccaccagcgagtaaggcatgagcgggtgttatccatgggctacgatccctacgatttcttgacctcggcgagtacta
tcagaaggggacagttgagacgcgttcggctcaaagggaagaactggtgaacatgataaacaccgcacactcctacggcataaaggatgag
cggacatagtcataaaccaccgcgcgggtggagacgttgagtggaacccctcgtgaacgactataacctggacagacttctcaaaagtgcctc
cggtaaatatacggccaactaccttgacttccacccaacgagcttccactgtgtgatgaagggtacgttggaggataacctgatataatgtcacga
caaaagctgggaccagtagtggctctgggcgagcagcgaagctacgctgctacccaggaagcagatagggtgacgcctggcgttctgact
acgtcaagggtacggagcatgggttgaacgactggctcagctggtgggagggctgggctgtggagagtagtgggacacgaacgtgat
gcactctcaactgggcatacagcagcggcgccaagggtcttgacttccgccttactacaagatggacgaagccttcgacaacaccaatcc
cggcattagtgatgcactcagatacggccagacagtggtcagccgcgatccctcaaggcgtaacttctgccaaccacgatacagatat
aatctggaacaagtatccggttatgcattcattacctatgaggacagcctgttatatttaccgcgactacgaggagtggtcacaacagga
taagcttaacaacctcattctggatacagatcaccttgcggaggagtagtactgacattgttactacgacagcagcagcttatcttggagaac
ggctatggcaccacaacaggactgataacctatataacccggctcaagcaaagtggaggtgggtctacgttccaaagttcgccgggtcat
gcattccagagtacaccggcaacctcggcggttgatagacaagtagcttccctccagcggctgggtctatcttgaggccccagccacgac
ccggcgaaacggctactacggctactctgctggagctactgcgggtggtgga

SEQ ID NO: 110

Met Ala Arg Lys Thr Leu Ala Ile Phe Phe Val Leu Leu Val Leu Leu Ser Leu Ser Ala Val Pro Ala
Lys Ala Glu Thr Leu Glu Asn Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Gly
Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro Glu Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro
Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu
Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr
Ala His Ser Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp
Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn
Tyr Leu Asp Phe His Pro Asn Glu Leu His Cys Cys Asp Glu Gly Thr Phe Gly Gly Tyr Pro Asp Ile
Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Ser Glu Ser Tyr Ala Ala Tyr Leu Arg
Ser Ile Gly Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Asn Asp Trp
Leu Ser Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp
Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn
Thr Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser Arg Asp Pro Phe Lys
Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu
Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Ser Asp
Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Thr Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser
Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn
Leu Gly Gly Trp Ile Asp Lys Tyr Val Ser Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp
Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

SEQ ID NO: 111

atgcccgcttcaaatcaaggatgcacatgaagtggaagtagcttcttagtttggctgtggcttcgataggcctcctcgcactccagt
gggtgctgccaaagtactccgaactcgaagaggcggtgtataatgcaggccttctactgggacgtccctaccgggtgggatctgggtggacac
cataagacagaaaaatcccgagtggtacgacgctggaatcgcgcgataggaattcctccagctagcaaaaggtatgggtggtcattccatg
ggttatgaccttaccgatttcttgacctcggcgagtactatcagaagggaacagttgagacgcgcttcggctcaaaggaggaaactggtgaaca
tgaataacaccgcacactcctatggcataaaggatagcggacatagtcataaacaccgcgcggcgccggcgacgttgagtggaacccctttg
taaacaactaacttggacagacttctccagggtgcctccggtaatacacggccaactaccttgacttccaccaaacgagggtcaagtgctgc
gatgaggggtacattggtagcttccggacatcgccacgagaagagctgggacagtagtggctcggcgaagcaatgagagctacgcgcgc

FIGURE 16RR

tatctccggagcatagggatcgaatggcgtttcgactacgtcaaaaggttacggagcgtgggttgtaacgactggctcagcttggtggggag
gttgggccgttgagagtactgggacaccaacgttgatgcactccttaactgggcatacaacagcgggtgccaaagctcttgactcccgtctac
tacaagatggacgaagccttgacaacaccaacatccccgtttggtttacgccctccagaacggaggaaacagtcgttccgcgatccctcaa
ggcagtaactttcgttgcaaacacgataccgataatctggaacaagtaaccggcttatgcgttcaacctatgagggacagcctgttatat
ctactacgataacgatgagctaattctatgagggagggtctacgggagcaagccgggcccataacctacataaacctcggaacgactggg
ccgagcgtgggtgaacgtcggtcctaaagtgtgccggctacacaatccatgaatacacaggcaatcctggtggctgggttgacaggtgggttc
agtacgacggatgggttaaactgacggcaccctcacgatccagccaacggalatatcggctactcagcttgaggctacgcaggcgtcggat
ga

SEQ ID NO: 112

Met Pro Ala Phe Lys Ser Lys Val Met Lys Leu Lys Tyr Leu Ala Leu Val Leu Leu Ala Val
Ala Ser Ile Gly Leu Ser Thr Pro Val Gly Ala Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile
Met Gln Ala Phe Tyr Trp Asp Val Pro Thr Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro
Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser
Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr
Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala His Ser Tyr Gly Ile Lys Val Ile
Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asn Tyr Thr
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu
Val Lys Cys Cys Asp Glu Gly Thr Phe Gly Asp Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln
Tyr Trp Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg
Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Asn Asp Trp Leu Ser Trp Trp Gly Gly Trp Ala
Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Asn Ser Gly Ala Lys Val
Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Thr Asn Ile Pro Ala Leu Val Tyr
Ala Leu Gln Asn Gly Gly Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His
Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile
Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu Asn Asn Leu Ile Trp Ile His Glu His
Leu Ala Gly Gly Ser Thr Lys Ile Leu Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Met Arg Glu Gly Tyr
Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Asn Asp Trp Ala Glu Arg Trp Val Asn Val
Gly Ser Lys Phe Ala Gly Tyr Thr Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Arg Trp
Val Gln Tyr Asp Gly Trp Val Lys Leu Thr Ala Pro Pro His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr
Ser Val Trp Ser Tyr Ala Gly Val Gly

SEQ ID NO: 113

atgaaacaacaaaaacgctttacgcccgattgctgacgctgttatttgcgcatacttcttgcgcctcatttcgcagcagcggcgccaacttta
atgggacgctgatgagatatttgaatggtacatgcgcgaatgacggcgaacattggaagcgtitgcaaaacgactcggcataattggctgaacac
ggatattcgtgatgctggaattcccggaatataagggaactagcgaagggaatgctggcctacgggtgcttaagaccttatgattatagggaatt
catcaaaaagggaacggttcggacaangtaoggcacaaaaggtagcgtgcaactgcgcatcaaaagtcttctcattccgcgcacattaacgtttacg
gggatgtggtcatcaaccacaaaaggcggcgctgatgcgaccgaagatgtaacccgcggttgaaagtcgacccgctgaccgcaaccgcgtaatt
tcaggagaacaccgaattaaagcctggacacatttcatittccggggcgcggcagcacatacagcgatttaaatggcattggtaccattttgac
ggaaaccgattgggacgagtcgccgaaagctgaaccgcactataagtticaaggaaaggcctgggattgggaagttccaatgaaaacggcaac
tatgattatttgatgatgccgacatcgattatgacatccgatgtcgcagcagaattaaagatggggcacttggatgccaatgaactgcaat
ggacggtttccgcttgatgctgtcaaacacattaaatttctttttcgggattggggttaatcatgtcagggaaaaaacggggaaggaaatgttta
cggtagctgaatatggcagaatgacttgggcgcgctggaaaactatttgaacaaaaaaatttaatacttcagtgtttgacgtgccgcttcattat
cagttccatgctgcatcgacacaggggagcgcgctatgatatgaggaaattgctgaacggtagcggttcgaagcatccgttgaagcggtta
catttgcgataaccatgatacacagccgggggcaatcgcttgatgcgactgtccaaacatggttttaagccgcttgcttacgctttcatttcacaag
ggaaactggataccctcaggttttctacggggatagtacgggacgaaaggagactccagcgcgaaattcctgccttgaaacacaaaatgaa
ccgactttaaagcgagaaaaacagtagtcgtacggagcagcatgattatttcgaccaccatgacattgtcggctggacaaagggaaggcgac
agctcgggttgcaaatcaggtttggcggcattataacagacggaccgggtggggcgaaggcaatgatgttcggccggcgaaacgccgggtga

FIGURE 16SS

gacatggcatgacattaccggaaacggttcggagccgggtgtcatcaattcggaaaggctggggagagttcacgtaaaccggcgggtcgggttca
attatgttcaaagatag

SEQ ID NO: 114

Met Lys Gln Gln Lys Arg Leu Tyr Ala Arg Leu Leu Thr Leu Leu Phe Ala Leu Ile Phe Leu Leu Pro
His Ser Ala Ala Ala Ala Asn Leu Asn Gly Thr Leu Met Gln Tyr Phe Glu Trp Tyr Met Pro Asn
Asp Gly Gln His Trp Lys Arg Leu Gln Asn Asp Ser Ala Tyr Leu Ala Glu His Gly Ile Thr Ala Val
Trp Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Ala Asp Val Gly Tyr Gly Ala Tyr Asp Leu Tyr Asp
Leu Gly Glu Phe His Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Gly Glu Leu Gln Ser Ala
Ile Lys Ser Leu His Ser Arg Asp Ile Asn Val Tyr Gly Asp Val Val Ile Asn His Lys Gly Gly Ala
Asp Ala Thr Glu Asp Val Thr Ala Val Glu Val Asp Pro Ala Asp Arg Asn Arg Val Ile Ser Gly Glu
His Arg Ile Lys Ala Trp Thr His Phe His Phe Pro Gly Arg Gly Ser Thr Tyr Ser Asp Phe Lys Trp
His Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile Tyr Lys Phe Gln
Gly Lys Ala Trp Asp Trp Glu Val Ser Asn Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Ile
Asp Tyr Asp His Pro Asp Val Ala Ala Glu Ile Lys Arg Trp Gly Thr Trp Tyr Ala Asn Glu Leu Gln
Leu Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Phe Ser Phe Leu Arg Asp Trp Val Asn His
Val Arg Glu Lys Thr Gly Lys Glu Met Phe Thr Val Ala Glu Tyr Trp Gln Asn Asp Leu Gly Ala
Leu Glu Asn Tyr Leu Asn Lys Thr Asn Phe Asn His Ser Val Phe Asp Val Pro Leu His Tyr Gln Phe
His Ala Ala Ser Thr Gln Gly Gly Gly Tyr Asp Met Arg Lys Leu Leu Asn Gly Thr Val Val Ser Lys
His Pro Leu Lys Ala Val Thr Phe Val Asp Asn His Asp Thr Gln Pro Gly Gln Ser Leu Glu Ser Thr
Val Gln Thr Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Glu Ser Gly Tyr Pro Gln Val
Phe Tyr Gly Asp Met Tyr Gly Thr Lys Gly Asp Ser Gln Arg Glu Ile Pro Ala Leu Lys His Lys Ile
Glu Pro Ile Leu Lys Ala Arg Lys Gln Tyr Ala Tyr Gly Ala Gln His Asp Tyr Phe Asp His His Asp
Ile Val Gly Trp Thr Arg Glu Gly Asp Ser Ser Val Ala Asn Ser Gly Leu Ala Ala Leu Ile Thr Asp
Gly Pro Gly Gly Ala Lys Arg Met Tyr Val Gly Arg Gln Asn Ala Gly Glu Thr Trp His Asp Ile Thr
Gly Asn Arg Ser Glu Pro Val Val Ile Asn Ser Glu Gly Trp Gly Glu Phe His Val Asn Gly Gly Ser
Val Ser Ile Tyr Val Gln Arg

SEQ ID NO: 115

atggcgaagtactccgagctgggacgaggcgagtcataatgcaggcccttacttgggacgttccggagggaagaaatctggtgggacacaat
acggcagaagatccctgaatggtacgatgcaggcatalccgccatctggataccccggcgagcaaggccatgggcggggcctactcga
ggctacgacccctacgattacttcgatctgggcgagtttaccagaagggaaccgttgagaccgcgttcggctccaaaggaaagagctcgtcaaca
tgatctccacggcccaccagtacggcatcaagggttatagcggacatagtataaaccaccgcgcaggtggagaccctcgaatggaaccatac
gtcggcgactatacctggacggacttttctaaggctgccctcgggaataacaaggccactacatggacitccatccaaacaactacagcacct
cagacgaggggaaccttcggttggttccagacatigatcacctcgtgcccttcaaccagtaactggtctgtggcgagcaacgagagctacgccg
cctactcagggcgaataggatcgatgcgtggcggtttgactacgttaagggtacggcgcggtgggtgttcaaggactggctgagtcagtggg
gcggcggggcggtggggaagctcgaacacacgttcgaatgggtctcaatgggtctcaggaacggggcgaagggttctgaatcc
gtctatacagatggagggcggtgacaaaggaatctccgccctgtttacgccatccagaacgggtgaacccgtcgtcagcaggat
cccttcaaggccgttaccttcgtggctaaccagatacgaacataatctggaaacagtacctgcctatgccttcatcctgacctacgaaggtcag
cccgtcatcttctaccgcgactacgaggatgggtcaacaaggacaaactcaaacctcatatggattcacgagcacctggcagggggaag
caccaagatactctactacgacgacgatgagctatcttcatgagggaaggctacggcgacaggccgggttataacctacatcaacctcgg
agcgactgggcggagagatgggtgaacgttggctcaaaagtgcgggctatacaatccagaaatacaccggaaacctcggcggttggtcg
acaggtagctccagtacgacggctgggtcaagcttaccgtctccacacgatccggcaaacggctattacggctactcggcttggagctacg
ccggagtgtggaagatctcatcaccatcaccatcactaa

SEQ ID NO: 116

Met Ala Lys Tyr Ser Glu Leu Glu Gln Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Glu
Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp
Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
Leu Gly Glu Phe Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met

FIGURE 16TT

Ile Ser Thr Ala His Gln Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
 Leu Glu Trp Asn Pro Tyr Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr
 Lys Ala His Tyr Met Asp Phe His Pro Asn Asn Tyr Ser Thr Ser Asp Glu Gly Thr Phe Gly Gly Phe
 Pro Asp Ile Asp His Leu Val Pro Phe Asn Gln Tyr Trp Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala
 Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val
 Lys Asp Trp Leu Ser Gln Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu
 Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala
 Phe Asp Asn Lys Asn Ile Pro Ala Leu Val Tyr Ala Ile Gln Asn Gly Glu Thr Val Val Ser Arg Asp
 Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asn Ile Ile Trp Asn Lys Tyr Pro Ala Tyr
 Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 Asp Lys Leu Asn Asn Leu Ile Trp Ile His Glu His Leu Ala Gly Gly Ser Thr Lys Ile Leu Tyr Tyr
 Asp Asp Asp Glu Leu Ile Phe Met Arg Glu Gly Tyr Gly Asp Arg Pro Gly Leu Ile Thr Tyr Ile Asn
 Leu Gly Ser Asp Trp Ala Glu Arg Trp Val Asn Val Gly Ser Lys Phe Ala Gly Tyr Thr Ile His Glu
 Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Arg Tyr Val Gln Tyr Asp Gly Trp Val Lys Leu Thr Ala
 Pro Pro His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Ala Gly Val Gly Arg Ser
 His His His His His His

SEQ ID NO: 117

ttgcgagtgttctgtgtgtgccaagctgagccgccatttcaggcagagtcacaacaacagacaggacataacaatgaacacacagcg
 ggaatgtctggcgatcgaggtatgtctgatcgcccccttggcgcatgccgatgtcactatgcacgccttcaactggaaatacagtgaaatcaccg
 ccaaggccgatctcatcaaggtgcccgttacaagcaggtgtctatctaccgccttgaagtctcgggcaacgagtggtgggctcgttacc
 agccccaggatctgcccgtgtgacaccccccttggcaacaagcaggtatctggagcagctgatcgccgcgatgcagacccgggcatg
 cgtctacgcggagctgtgtgtctcaaccacatggccaacgaaagctggaagcgcagcgacctcaactacccggcagcgagctgtgcaag
 ctacgcccggcaatcggcctactttgaacgccagaagcttttggcgatctggggcagaacttctcgcggccaggttttcatccggagggg
 tgcatacccgactggcaaatccgggccatgtccagttactggcgactgtgcccggggcggtgacaaggggctgccggatctggaccca
 acaactgggtgtgtaaccageaacaggttacctgcaggcgctcaaggggatggggatcaagggttttcgggtcgatcggtcaagcatg
 agcgattaccagatcaacgccgtgttcaccccgagatcaaacaggggatgcacgtctttggcgaggtgatcacccacggggggcgccggca
 acagcgactatgagaacttctcaaacctacctgcacagcagcgccagggggcctacgacttcccgtcttcgctccctgcgtggagcgc
 tgggtctacggcgagcatgaacctgtgtggcgatcccggtgctatgttcaggcgctgcccgggtagccgcgcccacatttcgcatcacc
 cagcacatccccaccaacgacggtttccgctaccagatcctcaaccagaccgacgagagactggcctatgctacgtctcggctcgatggc
 ggttcgctctgtgtactccgatcacggtgaaaccagggaaggacggatgctgctggcaggactactatctgcaccgatctcaagggg
 atgatccgcttcataacacagtgagggtaaccgatgcagctcatggcagtaacgactgctcgtgtgttcaagcgtggcaagcagggc
 gtggtcgcatcaacaagtgactacgagcaggagtagctgctcgtacccgagattcgagatgaactggtatcgcaactacgggatgtg
 ctcgaccagaatgccgtgtgtaacgtgcagagccagtggttaaggctgaccatcccggcccggcgccagaaatgtggtgcaggagtgga

SEQ ID NO: 118

Met Arg Val Phe Leu Val Val Pro Lys Leu Ser Arg Pro Phe Gln Ala Glu Ser Gln Gln Gln Asp Arg
 Asp Ile Thr Met Lys His Thr Ala Gly Met Leu Ala Ile Ala Gly Met Leu Ile Ala Pro Leu Ala His
 Ala Asp Val Ile Leu His Ala Phe Asn Trp Lys Tyr Ser Glu Val Thr Ala Lys Ala Asp Leu Ile Lys
 Ala Ala Gly Tyr Lys Gln Val Leu Ile Ser Pro Pro Leu Lys Ser Ser Gly Asn Glu Trp Trp Ala Arg
 Tyr Gln Pro Gln Asp Leu Arg Leu Val Asp Thr Pro Leu Gly Asn Lys Gln Asp Leu Glu Gln Leu Ile
 Ala Ala Met Gln Thr Arg Gly Ile Ala Val Tyr Ala Asp Val Val Leu Asn His Met Ala Asn Glu Ser
 Trp Lys Arg Ser Asp Leu Asn Tyr Pro Gly Ser Glu Leu Leu Gln Ser Tyr Ala Gly Asn Pro Ala Tyr
 Phe Glu Arg Gln Lys Leu Phe Gly Asp Leu Gly Gln Asn Phe Leu Ala Gly Gln Asp Phe His Pro
 Glu Gly Cys Ile Thr Asp Trp Asn Asn Pro Gly His Val Gln Tyr Trp Arg Leu Cys Gly Gly Ala Gly
 Asp Lys Gly Leu Pro Asp Leu Asp Pro Asn Asn Trp Val Val Asn Gln Gln Gln Ala Tyr Leu Gln
 Ala Leu Lys Gly Met Gly Ile Lys Gly Phe Arg Val Asp Ala Val Lys His Met Ser Asp Tyr Gln Ile
 Asn Ala Val Phe Thr Pro Glu Ile Lys Gln Gly Met His Val Phe Gly Glu Val Ile Thr Thr Gly Gly
 Ala Gly Asn Ser Asp Tyr Glu Asn Phe Leu Lys Pro Tyr Leu Asp Ser Ser Gly Gln Gly Ala Tyr Asp
 Phe Pro Leu Phe Ala Ser Leu Arg Gly Ala Leu Gly Tyr Gly Gly Ser Met Asn Leu Leu Ala Asp Pro

FIGURE 16UU

Gly Ala Tyr Gly Gln Ala Leu Pro Gly Ser Arg Ala Val Thr Phe Ala Ile Thr His Asp Ile Pro Thr
 Asn Asp Gly Phe Arg Tyr Gln Ile Leu Asn Gln Thr Asp Glu Arg Leu Ala Tyr Ala Tyr Leu Leu Gly
 Arg Asp Gly Gly Ser Pro Leu Val Tyr Ser Asp His Gly Glu Thr Arg Asp Lys Asp Gly Leu Arg Trp
 Gln Asp Tyr Tyr Leu Arg Thr Asp Leu Lys Gly Met Ile Arg Phe His Asn Thr Val Gln Gly Gln Pro
 Met Gln Leu Ile Gly Ser Asn Asp Cys Phe Val Leu Phe Lys Arg Gly Lys Gln Gly Val Val Gly Ile
 Asn Lys Cys Asp Tyr Glu Gln Glu Tyr Trp Leu Asp Thr Ala Arg Phe Glu Met Asn Trp Tyr Arg
 Asn Tyr Arg Asp Val Leu Asp Gln Asn Ala Val Val Asn Val Gln Ser Gln Trp Val Arg Leu Thr Ile
 Pro Ala Arg Gly Ala Arg Met Trp Leu Gln Glu

SEQ ID NO: 119

atgcaaacgtttgcattcttatttactcaagaagggatgggtgcatgaattattgaaaaagtggtgtattacgctatcgctacccitaa
 tcatttccctttacaccccttttcaacagcacaagctaactgcacctgtaacggaacaatgatgcaatatttcgaatgggacttacctaagtatgg
 gacgctttggacgaaagtaaaaaatgaagctaccaatcttttctactaggtatcacagcactatggctccctccagcatataaagggaacgagcc
 aaagcgatgctggatagcgtgtttacgatttatgaccttgggaatttaatacaaaaaggacgatccgaacgaaatacggaaacaaaacaca
 atatattcaagccattcaaaactgcccaagccgcagggaigcaagtatatgcggatgttatttaatacgaagcaggggctgacagtacagaatt
 tgcgatgcagttgaggtaaaccccttcaatcgaatacaagaacatctggcacatatcaaatcaagcatggacaaaatttgatttctgtcgtg
 gaaacacatactccagcttcaaatggcgctgttaccattttgatggtacggatgggacgaaagtcgtaaaattaatcgtattacaattccgcgg
 tacaggaaaagcgtgggactgggaagtcgatacagaaaacggaactatgattatttaagtcgctgatttagatggatcacccctgaggtgt
 gacagaattaaaaaacgtgggaacgtgtacgtcaatactacaataatcgatggatccgcttagatgccgtaaacatattaaatcacgcttttc
 cctgactggctaataatgtacgtaatacaacaggaaaaatttatttgcggtgggaattttggagctatgacgtcaataagctgcataattacat
 tacaacaaacaaatgggtcgtatgatttattgatgcacccctgcatacaacattttataccgcttccaaatcgagtggaattttgacatgcgttattat
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 aaccttggtttaaacagcttgccttattttaacaagacaagaagggtatccctgctattttacgggtattattatggaatccctaataacaat
 atcccggggttaaaaagtaaaatcgacccgcttttaattgctcgtcgtgattacgcttattggaacacaacgtgattacattgatcatcaagacattat
 cggatggacacgagaaggcattgatgcaaaaccgaactctggactggcggcttaattaccgacggctcgtgggaagtaaatggatgtatgc
 ggtaaaaagcatgccgggaaagtattttatgatttaactggaaatcgaagtacacagtaacgattaatgoggatgggtgggagaattaaagta
 aacggaggatccgtctcaatttgggtggctaaacgtcaaacgtcacattacgtcaataacgccacaacaacagcggacaaaacgtatag
 ttgctggcaacattccagagctaggcaattgtcgacgggtaa

SEQ ID NO: 120

Met Gln Thr Phe Ala Phe Leu Phe Tyr Ser Lys Lys Gly Trp Val Cys Met Asn Tyr Leu Lys Lys Val
 Trp Leu Tyr Tyr Ala Ile Val Ala Thr Leu Ile Ile Ser Phe Leu Thr Pro Phe Ser Thr Ala Gln Ala Asn
 Thr Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp Leu Pro Asn Asp Gly Thr Leu
 Trp Thr Lys Val Lys Asn Glu Ala Thr Asn Leu Ser Ser Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro
 Ala Tyr Lys Gly Thr Ser Gln Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe
 Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly Thr Lys Thr Gln Tyr Ile Gln Ala Ile Gln Thr Ala
 Gln Ala Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala Asp Ser Thr Glu
 Phe Val Asp Ala Val Gln Val Asn Pro Ser Asn Arg Asn Gln Glu Thr Ser Gly Thr Tyr Gln Ile Gln
 Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His
 Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile Tyr Lys Phe Arg Gly Thr Gly Lys
 Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Phe Ala Asp Leu Asp
 Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly Thr Trp Tyr Val Asn Thr Thr Asn Ile
 Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr Ser Phe Phe Pro Asp Trp Leu Thr Tyr Val
 Arg Asn Gln Thr Gly Lys Asn Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Val Asn Lys Leu His
 Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu Phe Asp Ala Pro Leu His Asn Asn Phe Tyr Thr
 Ala Ser Lys Ser Ser Gly Tyr Phe Asp Met Arg Tyr Leu Leu Asn Asn Thr Leu Met Lys Asp Gln
 Pro Ser Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln Pro Gly Gln Ser Leu Gln Ser Trp Val
 Glu Pro Trp Phe Lys Gln Leu Ala Tyr Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe
 Tyr Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Gly Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile
 Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln Arg Asp Tyr Ile Asp His Gln Asp Ile Ile Gly Trp Thr

FIGURE 16VV

Arg Glu Gly Ile Asp Ala Lys Pro Asn Ser Gly Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser
 Lys Trp Met Tyr Val Gly Lys Lys His Ala Gly Lys Val Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp
 Thr Val Thr Ile Asn Ala Asp Gly Trp Gly Glu Phe Lys Val Asn Gly Gly Ser Val Ser Ile Trp Val
 Ala Lys Thr Ser Asn Val Thr Phe Thr Val Asn Asn Ala Thr Thr Thr Ser Gly Gln Asn Val Tyr Val
 Val Gly Asn Ile Pro Glu Leu Gly Asn Cys Arg Thr Gly

SEQ ID NO: 121

atgctcgccctgtcgtcggcgggtgcggcatcgacgaggcccgacaggccctcgcgtcgtggagccgctgccgcagcgccccacgcttc
 cgcaggagtaccgcgccagcggccacgcggccgcccggcgacgtgttcgtgacactgttcgagtggaagtggccggacatcgccggaggat
 gcgagaacgtgctggggccggcgggtacgaggggtgcagggtgcggccgcaggagcacctggcgagcagggggcgccgtggtg
 gcagcgggtaccagccgggtgagctactcgggtggcgtgagccgcagcggcgacggcggtggagttcagcaacatgatcagccgggtgcaaggc
 cgccggcgtggacatctacgtggacggcgtcatcaaccacatgacggcgggtgcggggacggggagcaacggcaccgctacaccaagta
 caactaccccgccgtgacgcgcaggcggactttcacccgacgtgcgggtggcggaactacaccagcgccccaacgtgcaggactgcga
 actgctgggggtggtgacgtgaacaccggcgccggcggtgcagcagaagatcgcggaactacctggtctcgttgcgcggctggcggt
 ggccgggttttcgacgcgcgcggcccaagcacatccagccgggtggaactggacgccatcgtggaccgctgaaccagacgctggcgccgga
 gggggcgcccgcttccctactggttcgccgaggtgatcgacaacggcgccgaggggtgcggcgcgagcactactacggcctgggatacgg
 caccggcgcccgccggacatcacggagttccgctacaaggcggtggcgacaagtcttctgggcagcgccggccagcgccgtggtggacc
 tgaagaacttctcggcggtgacgtggaacctgatcgctcggaagggcgctgctttctggagaaccacgatacgagcgccggcgccggc
 atcggtaccgcgatggcacggcggttcggctggccaacgtgtggtatgctggcgacggcgatcggtatccgtcgggtgatgtccagctacgc
 ctttgaccgcacctccccctttggccgcgacggcgcccgccctccgaggacggcgcgacgaaggacgtgacgtgcgcgccacgctgga
 gacggcggtgctgggcacctgggtgtgcgagcaccgcgaccccgtaactcagcggtggtgggttttcgccgcgcatggcgggcacgga
 cctgaaccgctggtgggacaacggcggaacggcattgcttttcgcgggggaccggggcttcgtcgccatcagccgcgagccgaaggtg
 accatggcgccggtgcccagcggtgttccccggcactactcgacgtgctgacggcggaaggtgggaacgcctgcgcgggaac
 cagcgtgacggtcactctcaggcggtggtgcagctgagcatcgtcgagaactcggtctggtgatccacctggggccaagctgtaacggc
 gcgctggcggtatgtcgaggagg

SEQ ID NO: 122

Met Leu Ala Leu Ser Leu Gly Gly Cys Gly Ile Asp Ala Gly Pro Thr Gly Pro Arg Val Val Glu Pro
 Leu Pro Gln Arg Pro Thr Leu Pro Gln Glu Tyr Arg Ala Ser Gly His Ala Ala Ala Gly Asp Val Phe
 Val His Leu Phe Glu Trp Lys Trp Pro Asp Ile Ala Glu Glu Cys Glu Asn Val Leu Gly Pro Ala Gly
 Tyr Glu Ala Val Gln Val Ser Pro Pro Gln Glu His Leu Val Gln Gln Gly Ala Pro Trp Trp Gln Arg
 Tyr Gln Pro Val Ser Tyr Ser Val Ala Leu Ser Arg Ser Gly Thr Gly Val Glu Phe Ser Asn Met Ile
 Ser Arg Cys Lys Ala Ala Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn His Met Thr Ala Gly Ala
 Gly Thr Gly Ser Asn Gly Thr Ala Tyr Thr Lys Tyr Asn Tyr Pro Gly Leu Tyr Ala Gln Ala Asp Phe
 His Pro Gln Cys Ala Val Gly Asp Tyr Thr Ser Ala Ala Asn Val Gln Asp Cys Glu Leu Leu Gly Leu
 Ala Asp Leu Asn Thr Gly Ala Ala Gly Val Gln Gln Lys Ile Ala Asp Tyr Leu Val Ser Leu Ala Arg
 Leu Gly Val Ala Gly Phe Arg Ile Asp Ala Ala Lys His Ile Gln Pro Val Glu Leu Asp Ala Ile Val
 Asp Arg Val Asn Gln Thr Leu Ala Ala Gln Gly Arg Pro Leu Pro Tyr Trp Phe Ala Glu Val Ile Asp
 Asn Gly Gly Glu Gly Val Arg Arg Glu His Tyr Tyr Gly Leu Gly Tyr Gly Thr Gly Gly Ala Ala Asp
 Ile Thr Glu Phe Arg Tyr Lys Gly Val Gly Asp Lys Phe Leu Gly Ser Gly Gly Gln Arg Leu Val Asp
 Leu Lys Asn Phe Ser Ala Val Thr Trp Asn Leu Met Pro Ser Asp Lys Ala Val Val Phe Leu Glu Asn
 His Asp Thr Gln Arg Gly Gly Gly Ile Gly Tyr Arg Asp Gly Thr Ala Phe Arg Leu Ala Asn Val Trp
 Met Leu Ala Gln Pro Tyr Gly Tyr Pro Ser Val Met Ser Ser Tyr Ala Phe Asp Arg Thr Ser Pro Phe
 Gly Arg Asp Ala Gly Pro Pro Ser Glu Asp Gly Ala Thr Lys Asp Val Thr Cys Ala Pro Thr Leu Glu
 Thr Ala Val Leu Gly Thr Trp Val Cys Glu His Arg Asp Pro Val Ile Gln Arg Met Val Gly Phe Arg
 Arg Ala Met Ala Gly Thr Asp Leu Asn Arg Trp Trp Asp Asn Gly Gly Asn Ala Ile Ala Phe Ser Arg
 Gly Asp Arg Gly Phe Val Ala Ile Ser Arg Glu Pro Lys Val Thr Met Ala Ala Val Pro Ser Gly Leu
 Ser Pro Gly Thr Tyr Cys Asp Val Leu Thr Gly Gly Lys Val Gly Asn Ala Cys Ala Gly Thr Ser Val
 Thr Val Asp Ser Gln Gly Val Val Gln Leu Ser Ile Val Glu Asn Ser Ala Leu Val Ile His Leu Gly
 Ala Lys Leu Arg Arg Ala Gly Gly Cys Ala Glu

FIGURE 16WW

SEQ ID NO: 123

atgccccaggccattcgcactttttcacgttggacgttggtggttaacggcggttttctgcttgggtctcgtcttttctgcccaccccgggcaatcc
 agggccagacaaccccgggccgtaccgttatggttaccctcttcgagtggaaatggaccgacatcgctaaagaatgcgagaatttccctggac
 cgaaaggctttgcccgaatccagggtatcgccgcccaggagcatgtccagggggtcgcaatgggtggaccgctatcagccggtcagctacaag
 atcgagagccgctccggcaccggggcgagttcgccaatattggtctcgcgctgcaaaagccgtcggggctgatactatgtcgatgccgtgatc
 aaccatatgacgactgtcggctcggcactggatggctggatcgacctaccagctacacctatccggggctgtatcagacccaggacttcc
 accactgcgggcgcaatggcaacgatgatatcagcagctacggcgatcgctgggaagtacaaaactgcgaactgctcaacctagccgacctc
 aacaccggcgctgagtatgtccgggtaaacctgccgcctatatgaacgatctgcggcgctggggctcggcggttccggatcgatcgccgc
 aagcacatggataccaacgacatcaacaatactgttggcgccgtgcccgaacgcgccctacatctaccaggaagtgtaccaggggcgccga
 gccattaccggcggaatacttcagaatggcgatgtgaccgagttcaagtacagccgcgagatctcgcgatgttcaaaacggccagct
 gaccatatgagccagttcggcactgcctggggctcatgtccagcgacctggcagtagtttaccgataaccacgacaaccagcgcggtca
 cggcgggcgccggcgatgtcttgacctacaagatggccagctgtacacctgggcaatatcttcgagctagcctggcgatggctaccaca
 ggctatgtcgagctacacgttcagcaacggcgaccaggggcgccatcgaccaatgtgtacgcaaccacaacgcctgattgtggcaacggcc
 gctgggtctgtgagcaccgctggcgaggaaatcgccaatgtgtcggttcgcaactacaccgccccgaccttcagcaccagcaactgggtg
 agcaacggcaacaaccagatcgctttagccgcgggaccctgggcttggcgatcaatcggaagggtggcagcctgaaccgcaccttcca
 aaccggcctgcccgtcgccacactactgcgatgtcattcacggcgatttcaatgccagcgccggcaccctgttcggcccaactatcgctgtaac
 ggctccggcacaggcaacacacgggtcaacgcgatggacgggtggcgatctacggcgagccaggtcgccactccggccagtgtaac
 gtgacattcaacgaaaacggcacgaccacctggggcgagaatgtgtatatcgtcgcaacgtcgccgcccctgggcagctggaacgcaggca
 gcggggtcttactctcctccgctaactacccaatctggagcaagaccatcgccctgccagccaacaccgcccattgagtacaagtacatcaaaa
 ggatggcgcgggcaatgtgtgtgggaaagcgcgccaaccgcgtctttaccacccccggcagcgggcagtgccacgcgaacgatacctg
 gaaatag

SEQ ID NO: 124

Met Pro Gln Ala Ile Arg Thr Phe Ser Arg Trp Thr Leu Phe Gly Leu Ile Gly Val Phe Leu Leu Gly
 Leu Val Phe Ser Val Pro Pro Arg Ala Ile Gln Ala Gln Thr Thr Pro Ala Arg Thr Val Met Val His
 Leu Phe Glu Trp Lys Trp Thr Asp Ile Ala Lys Glu Cys Glu Asn Phe Leu Gly Pro Lys Gly Phe Ala
 Ala Ile Gln Val Ser Pro Pro Gln Glu His Val Gln Gly Ser Gln Trp Trp Thr Arg Tyr Gln Pro Val Ser
 Tyr Lys Ile Glu Ser Arg Ser Gly Thr Arg Ala Glu Phe Ala Asn Met Val Ser Arg Cys Lys Ala Val
 Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn His Met Thr Thr Val Gly Ser Gly Thr Gly Met Ala
 Gly Ser Thr Tyr Thr Ser Tyr Thr Tyr Pro Gly Leu Tyr Gln Thr Gln Asp Phe His His Cys Gly Arg
 Asn Gly Asn Asp Asp Ile Ser Ser Tyr Gly Asp Arg Trp Glu Val Gln Asn Cys Glu Leu Leu Asn
 Leu Ala Asp Leu Asn Thr Gly Ala Glu Tyr Val Arg Gly Lys Leu Ala Ala Tyr Met Asn Asp Leu
 Arg Gly Leu Gly Val Ala Gly Phe Arg Ile Asp Ala Ala Lys His Met Asp Thr Asn Asp Ile Asn Asn
 Ile Val Gly Arg Leu Pro Asn Ala Pro Tyr Ile Tyr Gln Glu Val Ile Asp Gln Gly Gly Glu Pro Ile Thr
 Ala Gly Gly Tyr Phe Gln Asn Gly Asp Val Thr Glu Phe Lys Tyr Ser Arg Glu Ile Ser Arg Met Phe
 Lys Thr Gly Gln Leu Thr His Met Ser Gln Phe Gly Thr Ala Trp Gly Phe Met Ser Ser Asp Leu Ala
 Val Val Phe Thr Asp Asn His Asp Asn Gln Arg Gly His Gly Gly Ala Gly Asp Val Leu Thr Tyr
 Lys Asp Gly Gln Leu Tyr Thr Leu Gly Asn Ile Phe Glu Leu Ala Trp Pro Tyr Gly Tyr Pro Gln Val
 Met Ser Ser Tyr Thr Phe Ser Asn Gly Asp Gln Gly Pro Pro Ser Thr Asn Val Tyr Ala Thr Thr Thr
 Pro Asp Cys Gly Asn Gly Arg Trp Val Cys Glu His Arg Trp Arg Gly Ile Ala Asn Met Val Ala Phe
 Arg Asn Tyr Thr Ala Pro Thr Phe Ser Thr Ser Asn Trp Trp Ser Asn Gly Asn Asn Gln Ile Ala Phe
 Ser Arg Gly Thr Leu Gly Phe Val Ala Ile Asn Arg Glu Gly Gly Ser Leu Asn Arg Thr Phe Gln Thr
 Gly Leu Pro Val Gly Thr Tyr Cys Asp Val Ile His Gly Asp Phe Asn Ala Ser Ala Gly Thr Cys Ser
 Gly Pro Thr Ile Ala Val Asn Gly Ser Gly Gln Ala Thr Ile Thr Val Asn Ala Met Asp Ala Val Ala Ile
 Tyr Gly Gly Ala Arg Leu Ala Thr Pro Ala Ser Val Asn Val Thr Phe Asn Glu Asn Ala Thr Thr Thr
 Trp Gly Gln Asn Val Tyr Ile Val Gly Asn Val Ala Ala Leu Gly Ser Trp Asn Ala Gly Ser Ala Val
 Leu Leu Ser Ser Ala Asn Tyr Pro Ile Trp Ser Lys Thr Ile Ala Leu Pro Ala Asn Thr Ala Ile Glu Tyr
 Lys Tyr Ile Lys Lys Asp Gly Ala Gly Asn Val Val Trp Glu Ser Gly Ala Asn Arg Val Phe Thr Thr
 Pro Gly Ser Gly Ser Ala Thr Arg Asn Asp Thr Trp Lys

FIGURE 16XX

SEQ ID NO: 125

gtggtgcacatgaagttgaagtaccttgcccttagtttggctgtggcttcgataggcctactctgactccagtggtgctgccaaagtactccg
 aactcgaagagggcggtgtataatgcaggcccttacttggtatgttcccgagggggaatctggtgggacaccataagacagaaaatccc
 gagtggtagcagcgtggaatctcggcgatgtgattctccagctagcaaaaggatggcggtgttaltccatgggtacgatccctacgattt
 ctttgacctcggcgagtagtactatcagaagggaacagttgagacgccttcggctcaaaaggaggaactggtgaacatgataaacaccgcacactc
 ctatggcataaaaggtgatagcggaatagtcataaaaccaccgcgcgggtggagaccttgagtgaacccctttgtaaacactatacttggaca
 gacttclccaaggctcgcctccggttaataacacggccaactaccttgacttccacccaaacgaggtcaagtgtcgtgatgagggtacatttggga
 ctttccggacatcgcccacgagaaagagctgggatcagtagtctggtggaagcaatgagagctacgccgcatactccggagcataggga
 tcgatgcagtggttctgactacgtcaaaaggttacggagcgtgggtgttaatgactggctcagctggtggggaggctggccgttgagagta
 ctgggacacgaacgttgatgcactccttaactgggcatacgacagcgggtgccaaggtctttgacttcccgtctactacaagatggaggaagcc
 ttgacaacaccaacatcccgtttgtttacgccctccagaacggaggaaacagtcgttccgcgatccctcaaggcagtaactttcgttgc
 aaccacgatacagataatctggaacaagtatccggcttatgcgttcatccttacctatgagggacagcctgttatatttaccgcgactacgagg
 agtggctcaacaaggataagcttaacaaccttcttgatacagagcaccttgcggaggaaagtaacagatcccttactacgataacgatga
 gctaataatcatgagggagggtacggagcaagccgggcctacataaacctcggaaacgactggccgagcgtggtggaac
 gtcggctcaaaagttgcggctacacaatccatgaatacacaggcaatctcgggtgctgggtgacaggtgggttcagtagcatggtggtta
 aactgacggcacctctcatgatccagccaacggatattacggctactcagcttgagctacgcaggcgtcgatga

SEQ ID NO: 126

Val Val His Met Lys Leu Lys Tyr Leu Ala Leu Val Leu Leu Ala Val Ala Ser Ile Gly Leu Leu Ser
 Thr Pro Val Gly Ala Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp
 Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile
 Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr
 Asp Phe Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu
 Leu Val Asn Met Ile Asn Thr Ala His Ser Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg
 Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asn Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala
 Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr
 Phe Gly Asp Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Asn Glu
 Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly
 Ala Trp Val Val Asn Asp Trp Leu Ser Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn
 Val Asp Ala Leu Leu Asn Trp Ala Tyr Asp Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys
 Met Asp Glu Ala Phe Asp Asn Thr Asn Ile Pro Ala Leu Val Tyr Ala Leu Gln Asn Gly Gly Thr Val
 Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys
 Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu
 Trp Leu Asn Lys Asp Lys Leu Asn Asn Leu Ile Trp Ile His Glu His Leu Ala Gly Gly Ser Thr Lys
 Ile Leu Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Met Arg Glu Gly Tyr Gly Ser Lys Pro Gly Leu Ile
 Thr Tyr Ile Asn Leu Gly Asn Asp Trp Ala Glu Arg Trp Val Asn Val Gly Ser Lys Phe Ala Gly Tyr
 Thr Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Arg Trp Val Gln Tyr Asp Gly Trp Val
 Lys Leu Thr Ala Pro Pro His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr Ala Gly
 Val Gly

SEQ ID NO: 127

gtgtgcatgaattattgaaaaaagtgtggtgtattacgtatcgtcgtacctaattactttcttaccgcccttttcaactgcacaagccaacac
 tgcaccagtcacaggaacgatgatgcaatatttgaatgggtattaccgaatgatggcacactttggacgaaagtaaaaaacgaagcaagcagt
 ctttctttaggtattactcgttatgttaccacctgcatacaaaaggaacgagccaaggggatgtcgggtatggcgtgtacatttgtatgactt
 aggaagaatttaatacaaaaaggacgattcgaacgaataacggaacaaaaacgcaatatttacaagccattcaagcggcaaaaagcgtggcat
 gcaagtatacgtgtatgtctatttaatacaaggcggggcagatagtagaagaatgggtgacgcagtcgaagtgaatccttctaatacgaacc
 aagaaacatctggcacatatacaattcaagcatggacaaaatttgatttccctggccgtgggaacacatactcaagctttaaattggcgtatgatac
 attttgacggtagcggattgggatgaaagccgaaactaaatcgtatttacaatttctgggcacagggaaagcatgggattgggaagttagacaca
 ggaacaggaaactatgactacttaattgttctgatttagatattgataccctgaagctgtgacagagctaaaaaactggggaaacaggtacgtc

FIGURE 16YY

aatacgacaaatgctgatgggttcgcttagatgcagtaagcataaataatagcttccagattggtaacacatgtgcgttcacaaacag
 aaaaaatcttttgcagtaggagaattttggagctacgatgtcaataaactgcataactacattacaaaaaagaaggaaacatgtcgttattgatg
 cgccactcacaacacttttactgctcaaaatctagcgggtatttgacatgcgctatttgtaataatacgttgatgaaagaccagccttctct
 tgcgggtcacactcgttgataatcatgacacgcaaccgggacaatctttacaatcatgggtagagccttgggttaagccgcttgcttatgcctttattt
 gacaagacaagaaggatattccttgcgtattttacggcgactattacggcatccctaaatacaacattccgggattgaaaagtaaaatcgateccgt
 tctcattgcccgtagagactacgcatacggaaacacaacgtgattattgacatcaagacattattggatggacacgggaaggaattgactcaa
 aaccgaactctggacttgcggctttaattactgacggccctgggtggaagtaaatggatgtatgtaggtaaaaagcatgctggaaaagtgttttacg
 atctcactggaatcgaagcgatacggtaacgattaatgcagacggctggggagagtttaagtaaacgggtgctccgttccatttgggttgc
 aaaacatcacaaatgcacgttaccgtcaacaatgcgacaacgataagcggacaaaatgtgtatgtcgttgtaacattccagagctcggaattg
 gaacacagcaaacgcaatcaaatgacccatcttctatccaacgttggaagcaaccattgcttccacaaggaaaagcattgaatttaatt
 tattaanaagaccaatcgggaattgttttgggaagcattccaaaccgaacatacaccgttccattttatcaacaggctcatatacagctagt
 ggaatgtacctaa

SEQ ID NO: 128

Val Cys Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val Ala Thr Leu Ile Ile Tyr Phe
 Leu Thr Pro Phe Ser Thr Ala Gln Ala Asn Thr Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu
 Trp Asp Leu Pro Asn Asp Gly Thr Leu Trp Thr Lys Val Lys Asn Glu Ala Ser Ser Leu Ser Ser Leu
 Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Thr Ser Gln Gly Asp Val Gly Tyr Gly Val
 Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly Thr Lys Thr
 Gln Tyr Leu Gln Ala Ile Gln Ala Ala Lys Ser Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asn
 His Lys Ala Gly Ala Asp Ser Thr Glu Trp Val Asp Ala Val Glu Val Asn Pro Ser Asn Arg Asn Gln
 Glu Thr Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr Tyr
 Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg
 Ile Tyr Lys Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp
 Tyr Leu Met Phe Ala Asp Leu Asp Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly
 Thr Trp Tyr Val Asn Thr Thr Asn Val Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr Ser
 Phe Phe Pro Asp Trp Leu Thr His Val Arg Ser Gln Thr Arg Lys Asn Leu Phe Ala Val Gly Glu Phe
 Trp Ser Tyr Asp Val Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Ser Gly Thr Met Ser Leu Phe Asp
 Ala Pro Leu His Asn Asn Phe Tyr Thr Ala Ser Lys Ser Ser Gly Tyr Phe Asp Met Arg Tyr Leu Leu
 Asn Asn Thr Leu Met Lys Asp Gln Pro Ser Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln
 Pro Gly Gln Ser Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr
 Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Gly
 Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln Arg Asp Tyr
 Ile Asp His Gln Asp Ile Ile Gly Trp Thr Arg Glu Gly Ile Asp Ser Lys Pro Asn Ser Gly Leu Ala Ala
 Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Lys His Ala Gly Lys Val Phe
 Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ala Asp Gly Trp Gly Glu Phe Lys Val
 Asn Gly Gly Ser Val Ser Ile Trp Val Ala Lys Thr Ser Gln Val Thr Phe Thr Val Asn Asn Ala Thr
 Thr Ile Ser Gly Gln Asn Val Tyr Val Val Gly Asn Ile Pro Glu Leu Gly Asn Trp Asn Thr Ala Asn
 Ala Ile Lys Met Thr Pro Ser Ser Tyr Pro Thr Trp Lys Ala Thr Ile Ala Leu Pro Gln Gly Lys Ala Ile
 Glu Phe Lys Phe Ile Lys Lys Asp Gln Ser Gly Asn Val Val Trp Glu Ser Ile Pro Asn Arg Thr Tyr
 Thr Val Pro Phe Leu Ser Thr Gly Ser Tyr Thr Ala Ser Trp Asn Val Pro

SEQ ID NO: 129

ttcggttgcgcgctggcagggaagggtgtgtgtgcggcggttaagcgtgcccgcacaccgcgtgaacaaaaaatatgaattattt
 aataggatgggggtgtcaagaatgacaaaatctcgagagttgcggtgttcaggaaagtatttgttgggtgcttggatggcttggga
 tcttccgcgtccgcccgtattgatgcaaggcttctactgggacgccagttaccgggaccagtgattcgtgttgagcgcatttggccaagcaag
 ccaacggtctaaacggcggggttcaccgcccgtattgattcctcgggtgcttaaaagggttcagggggtattccaacgggtacgatccct
 tgacgactatgatacgggaagcaaggaccagaagggtaccgtggcgacgcgatgggggacgcgagaagaactgcaacgtccgtggccgt
 gatgcgcgcgaacggtctggatgtgtatgtggtgctgaaccaccgcaacggggacgcgggaattggaattttcatcaaaagatgc
 gtacggcaaaagtgggttacggcggtttcaaaagggtttacgattttaccccactcaacattcaggatgccaatgttcccaacgaggattc

FIGURE 16ZZ

cagcttcgggcgcgatttagcccatgacaatccgtatgtggccgatggactgaagctgcaggcgattggctgaccaaagccctcgaatgta
 gggatactgcttgattacgtgaaaggcatcagctacaccttctgaaaagtatctgctctatggggccatgaacggaaaatttgcgcgtgga
 gtactgggatgccaacgggatacgtgaactgggtggcgaacacggcgatggaagggcgggccatgtgttatttgcgtgcgcgagg
 agctgaaaaacatgtgcaatgcggacgggtactacgacatgctgcgattggaccacgcgggtctggtcggaaatgaccctggaaggcgt
 gacgtttgtcgaataacatgatacggatcggcacgacccatctacaataaagcatttggcgtatgcctacatcttgacgtcgaagggtatc
 cgacgggtgttctggaaggattactaccaatcaggaatgaagccgatcgcacaacctcatttggatccacgaacacattgctgacggaacgac
 ccaagagcgttggaaagacgaagatgtcttgtgtatgagcgacggaggcaagcggctatttgggtggggttaacgacaatcgcgccacca
 gcaaacgggtcacggtacagaccggcttgggtgccaacgtggccttgcacgactacaccggcaacggccccgatctccgtaccgacgcctac
 ggtegggtaacctgaccattctgcaaacgggtacgtggcctattcgttccgggcatctccggatccttggcgggtcgagaaaaccgtgac
 gcaggagtttccggggcgctcgacttggatattcgtccggcgataacacgcaatttgcaggctggcggtatatacgaaggcaaacaa
 gccggttacagcgggaattgtattggatgccaagactggacgacccacgtcgatttcttagaagtgcttcgggtcgggaacgctcatc
 acgacaaagaccgtgacccaattgtctgccagggtaccgcgttcttcacgccttcggctaccggatggtacgtctttccattcgaagctat
 aacacgccttcgacgaacccaagccggcctactggttaaggtaacgtatacggcgccgaattgcttcagtaa

SEQ ID NO: 130

Met Arg Cys Arg Arg Gly Arg Asp Gly Cys Trp Cys Gly Arg Arg Asn Ala Leu Pro Arg His Pro
 Arg Glu Gln Asn Asn Met Asn Tyr Leu Asn Arg Met Gly Val Ser Arg Met Thr Lys Ser Arg Glu
 Leu Arg Cys Ser Trp Lys Val Phe Val Val Gly Cys Leu Leu Trp Met Ala Trp Gly Ser Ser Ala Ser
 Ala Gly Val Leu Met Gln Gly Phe Tyr Trp Asp Ala Ser Thr Gly Thr Ser Asp Ser Trp Trp Thr His
 Leu Ala Lys Gln Ala Asn Gly Leu Lys Arg Ala Gly Phe Thr Ala Val Trp Ile Pro Pro Val Leu Lys
 Gly Ala Ser Gly Gly Tyr Ser Asn Gly Tyr Asp Pro Phe Asp Asp Tyr Asp Ile Gly Ser Lys Asp Gln
 Lys Gly Thr Val Ala Thr Arg Trp Gly Thr Arg Glu Glu Leu Gln Arg Ala Val Ala Val Met Arg Ala
 Asn Gly Leu Asp Val Tyr Val Asp Leu Val Leu Asn His Arg Asn Gly Asp Asp Gly Asn Trp Asn
 Phe His Tyr Lys Asp Ala Tyr Gly Lys Val Gly Tyr Gly Arg Phe Gln Lys Gly Phe Tyr Asp Phe His
 Pro Asn Tyr Asn Ile Gln Asp Ala Asn Val Pro Asn Glu Asp Ser Ser Phe Gly Arg Asp Leu Ala His
 Asp Asn Pro Tyr Val Ala Asp Gly Leu Lys Ala Ala Gly Asp Trp Leu Thr Lys Ala Leu Asp Val
 Gln Gly Tyr Arg Leu Asp Tyr Val Lys Gly Ile Ser Tyr Thr Phe Leu Lys Ser Tyr Leu Ser Tyr Gly
 Ala Met Asn Gly Lys Phe Ala Val Gly Glu Tyr Trp Asp Ala Asn Arg Asp Thr Leu Asn Trp Trp
 Ala Asn Thr Ala Met Glu Gly Arg Ala His Val Phe Asp Phe Ala Leu Arg Glu Glu Leu Lys Asn
 Met Cys Asn Ala Asp Gly Tyr Tyr Asp Met Arg Arg Leu Asp His Ala Gly Leu Val Gly Ile Asp
 Pro Trp Lys Ala Val Thr Phe Val Glu Asn His Asp Thr Asp Arg His Asp Pro Ile Tyr Asn Asn Lys
 His Leu Ala Tyr Ala Tyr Ile Leu Thr Ser Glu Gly Tyr Pro Thr Val Phe Trp Lys Asp Tyr Tyr Gln
 Tyr Gly Met Lys Pro Ile Ile Asp Asn Leu Ile Trp Ile His Glu His Ile Ala Tyr Gly Thr Thr Gln Glu
 Arg Trp Lys Asp Glu Asp Val Phe Val Tyr Glu Arg Thr Gly Gly Lys Arg Leu Leu Val Gly Leu
 Asn Asp Asn Arg Ala Thr Ser Lys Thr Val Thr Val Gln Thr Gly Phe Gly Ala Asn Val Ala Leu His
 Asp Tyr Thr Gly Asn Gly Pro Asp Leu Arg Thr Asp Ala Tyr Gly Arg Val Thr Leu Thr Ile Pro Ala
 Asn Gly Tyr Val Ala Tyr Ser Val Pro Gly Ile Ser Gly Ser Phe Val Pro Val Gln Lys Thr Val Thr
 Gln Glu Phe Ala Gly Ala Ser Asp Leu Asp Ile Arg Pro Ala Asp Asn Thr Gln Phe Val Gln Val Gly
 Arg Ile Tyr Ala Lys Ala Asn Lys Pro Val Thr Ala Glu Leu Tyr Trp Asp Ala Lys Asp Trp Thr Thr
 Ser Thr Ser Ile Leu Leu Glu Val Arg Ser Ala Ser Gly Thr Leu Ile Thr Thr Lys Thr Val Thr Gln Leu
 Ser Ser Gln Gly Thr Arg Val Ser Phe Thr Pro Ser Ala Thr Gly Trp Tyr Val Phe Ser Ile Arg Ser Tyr
 Asn Thr Pro Ser Thr Asn Pro Lys Pro Ala Tyr Trp Leu Lys Val Thr Tyr Thr Ala Pro Gln Leu Leu
 Gln

SEQ ID NO: 131

atgccgcagctttaccattgccgcgcgtggcggcgcgccggcgaggcctggccgccttgacgtggccaccacggccctgggc
 atctcgacggccagggccagagtgcaccgcgcacggccttctgtcatctgttgaatggaagtggaccgacatcgcgcgagtgcgaga
 ccttctcgggcccgaagggttcgcggcggtgcaggtgtcgccccgaacgagcacaactgggtgaccagcggtgatggtgcaccttaccg
 tgggtggatgcctaccagcgggtgagctacagcctggaccgcagcgcgcggcgacgcgcggcgagttccaggacatggtcaaccgatgc
 aatgccgtggcggtgggcattcgtggacggcggtgatcaatcacatgtccggcgccggcgccggcgacctcgagcgctggggcgagctgg

FIGURE 16AAA

agctatcacaactacccctgggtctatggccccaacgacttccaccagccgggtgtgcagcatcaccaactacggggatgcgaacaatgtgcag
 cgttgcgagctctcgggttcgaggacctggacactgggagcgcttatgtgcgcggaagatcgccgactatctggtggatctgtgtaacatg
 ggggtcaagggttcgggtggatgcggccaagcacatcagcccagccgacctgggcccacatcagatcggtgtaacagccgaccggc
 gcgaaccgccccttctggttctggaggtgattggcgcggccggcgaggcagtgagccgaaccagctactctcgtcggcggcgccaggt
 caccgtgaccgagttcaactatgggaagcaaatcttcggcaagtgcgggtggcgccgtctggccgagctgcgcagcttcggtgaaacctg
 gggcctgatgccagcagcaagcgattgcttcatcgacaaccacgacaagcagcgcggtcatggcggcggtggcaactatctgacctacc
 accatggctcgacctagatctggccaacatcttcatgctggcttggccttatggctacccggcgctgatgtccagctatgcttcaaccgcagc
 acggcctacgacacgagcttggcccgccacacgacagtggtggcgccaccgtggccctgggatggtggcgccagccagccggcctgc
 ttaaccagagcatcggtggctgggtgtgagcaccgctggcgggcgatcgccaatatggtggccttcgcaacgccacgctgcccactg
 gaccgtgaccgactgggtgggacaacggcaacaaccagatcgcttccggcggggtgacaagggttcgtggtgtaaccgcgaagacgc
 cgcgctgacgcgcaactcaagaccagcctgccagccggccagctactgcgatgtcatctccgggacttcaacaatggtcagtgacggcc
 atgtggtgacggctgatcgccggggctacgtgacgctgacggcgggcccaatggtcgcgccgcatccacgtggcgcccgctggacg
 gcgctctcagccgcccagaccgctcggtagcttaacgcgtcgccgataccttttggggacagaacctgttcgtcgtgggcaaccaca
 gcgcactgggcaactggtcgccggcgccggccagccgatgacttgatttcgggttcgggcacgcgcgggaactggcgcgcggtgtca
 atttgcgggccaataccacctaacaatacaagttcatcaagaaggacggggctggaaacgtggttgggagggcggtggcaatcgctcgtga
 ccacgcccgtctggggcggtgagcacggcgccgaattggcagtag

SEQ ID NO: 132

Met Pro Gln Leu Tyr Pro Leu Pro Pro Arg Trp Arg Arg Ala Ala Arg Gln Gly Leu Ala Ala Leu Thr
 Leu Ala Thr Thr Ala Leu Gly Ile Ser Thr Ala Gln Ala Gln Ser Ala Pro Arg Thr Ala Phe Val His
 Leu Phe Glu Trp Lys Trp Thr Asp Ile Ala Arg Glu Cys Glu Thr Phe Leu Gly Pro Lys Gly Phe Ala
 Ala Val Gln Val Ser Pro Pro Asn Glu His Asn Trp Val Thr Ser Gly Asp Gly Ala Pro Tyr Pro Trp
 Trp Met Arg Tyr Gln Pro Val Ser Tyr Ser Leu Asp Arg Ser Arg Ser Gly Thr Arg Ala Glu Phe Gln
 Asp Met Val Asn Arg Cys Asn Ala Val Gly Val Gly Ile Tyr Val Asp Ala Val Ile Asn His Met Ser
 Gly Gly Thr Gly Gly Thr Ser Ser Ala Gly Arg Ser Trp Ser Tyr His Asn Tyr Pro Gly Leu Tyr Gly
 Pro Asn Asp Phe His Gln Pro Val Cys Ser Ile Thr Asn Tyr Gly Asp Ala Asn Asn Val Gln Arg Cys
 Glu Leu Ser Gly Leu Gln Asp Leu Asp Thr Gly Ser Ala Tyr Val Arg Gly Lys Ile Ala Asp Tyr Leu
 Val Asp Leu Val Asn Met Gly Val Lys Gly Phe Arg Val Asp Ala Ala Lys His Ile Ser Pro Thr Asp
 Leu Gly Ala Ile Ile Asp Ala Val Asn Ser Arg Thr Gly Ala Asn Arg Pro Phe Trp Phe Leu Glu Val
 Ile Gly Ala Ala Gly Glu Ala Val Gln Pro Asn Gln Tyr Phe Ser Leu Gly Gly Gly Gln Val Thr Val
 Thr Glu Phe Asn Tyr Gly Lys Gln Ile Phe Gly Lys Phe Ala Gly Gly Gly Arg Leu Ala Glu Leu Arg
 Ser Phe Gly Glu Thr Trp Gly Leu Met Pro Ser Ser Lys Ala Ile Ala Phe Ile Asp Asn His Asp Lys
 Gln Arg Gly His Gly Gly Gly Gly Asn Tyr Leu Thr Tyr His His Gly Ser Thr Tyr Asp Leu Ala Asn
 Ile Phe Met Leu Ala Trp Pro Tyr Gly Tyr Pro Ala Leu Met Ser Ser Tyr Ala Phe Asn Arg Ser Thr
 Ala Tyr Asp Thr Ser Phe Gly Pro Pro His Asp Ser Gly Gly Ala Thr Arg Gly Pro Trp Asp Gly Gly
 Gly Ser Gln Pro Ala Cys Phe Asn Gln Ser Ile Gly Gly Trp Val Cys Glu His Arg Trp Arg Gly Ile
 Ala Asn Met Val Ala Phe Arg Asn Ala Thr Leu Pro Asn Trp Thr Val Thr Asp Trp Trp Asp Asn
 Gly Asn Asn Gln Ile Ala Phe Gly Arg Gly Asp Lys Gly Phe Val Val Ile Asn Arg Glu Asp Ala Ala
 Leu Thr Arg Asn Phe Lys Thr Ser Leu Pro Ala Gly Gln Tyr Cys Asp Val Ile Ser Gly Asp Phe Asn
 Asn Gly Gln Cys Thr Gly His Val Val Thr Val Asp Ala Gly Gly Tyr Val Thr Leu Thr Ala Gly Pro
 Asn Gly Ala Ala Ala Ile His Val Gly Ala Arg Leu Asp Gly Ala Ser Gln Pro Pro Thr Thr Ala Ser
 Val Thr Phe Asn Ala Ser Ala Asp Thr Phe Trp Gly Gln Asn Leu Phe Val Val Gly Asn His Ser Ala
 Leu Gly Asn Trp Ser Pro Ala Ala Ala Arg Pro Met Thr Trp Ile Ser Gly Ser Gly Thr Arg Gly Asn
 Trp Arg Ala Val Leu Asn Leu Pro Ala Asn Thr Thr Tyr Gln Tyr Lys Phe Ile Lys Lys Asp Gly Ala
 Gly Asn Val Val Trp Glu Gly Gly Gly Asn Arg Val Val Thr Thr Pro Ser Gly Gly Gly Ser Val Ser
 Thr Gly Gly Asn Trp Gln

SEQ ID NO: 133

atgaataatgtgaaaaaagtatggtgtattattctataattgctacccttagttatttcttttcacacctttttcaacagcacaagctaaactgcacctg
 tcaacgggaacaatgatgcataatttcgaatgggattaccgaatgatgggacgcttggacgaaggaanaalgaagctaccaaatcttttctcgt

FIGURE 16BBB

aggtattacagcggttatggctccctccagcatataaagggaacgagccaaagcgatgtcggatatggcgtgtacgatttatgacctggggaatt
 taatcaaaaaggacgatccgaacgaatacggaaacaaagcacaatatattcaagccatccaagctgccaagccgcagggatgcaagt
 atgcagatgttgatttaataaaggcgggggctgacggcacagaattgtcgtatgaggttaaaccttctaatacgaatacaaaacat
 ctggcacatatcaaatcaagcatggacaaaattgatcttcggctgtgaaacacatactccagcttcaaatggcgtgtatcatttggcgt
 accgattgggatgaaagtcgtaaattaaatcgtattacaattccgcgtacaggaaaagcggtggactgggaagtcgatacagaaaacgga
 aactatgattttaaattgtcgtgatttagatggtacccctgaagttgtgacagagttaaaaactggggaaatggtatgtaatacgcacaa
 ttagacggatttcgttggatgccgtaaaacataataacacgttttccctgactggctaacatatgtacgtatcaaacaggaaaaattatt
 gctgtggggaatttggagctatgacgtcaataagctgcataactacattacaaaaacaaatggatcgtatgtatttgcaccccttcata
 caacttttatcgcctccaaatcgaagtggatatttgacatgcgttatttgaataatacattatgaagatcaaccttactcgtgtacactgt
 cgataaccatgatacacaaccaggctcaatcttacaatcatggtagaagcttggtttaaacgcgttctacgcctttatitaaacagacaag
 gggtatcccttgcgtatttaccgtgactattacggatcccgaatacaatattccgggattaaaaagtaaaattgacccgttttaattgctcgt
 gattatgcttatggaacacaacgtgattacattgatcatcaagacattatcggatggacacgagaaggcattgatcaaaaccgaactctggact
 gcggctttaaattaccgacggccctggcggagtaaatggatgtatgctggtaaaaaacatgctgggaaagtgtttatgatttaactggaatcga
 agtgacacagtaacgattaatgcggacgggtgggagaaattaaagtaaaggcggtccgttcgatttgggtggctaaaacatcaaacgtca
 cattacagtcataacgccacaacaagaaggacaaaacgtatatgtgttggcaacattccagagctaggaattcttg

SEQ ID NO: 134

Met Asn Asn Val Lys Lys Val Trp Leu Tyr Tyr Ser Ile Ile Ala Thr Leu Val Ile Ser Phe Phe Thr Pro
 Phe Ser Thr Ala Gln Ala Asn Thr Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp Leu
 Pro Asn Asp Gly Thr Leu Trp Thr Lys Val Lys Asn Glu Ala Thr Asn Leu Ser Ser Leu Gly Ile Thr
 Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Thr Ser Gln Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu
 Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile
 Gln Ala Ile Gln Ala Ala Lys Ala Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asn His Lys Ala
 Gly Ala Asp Gly Thr Glu Phe Val Asp Ala Val Glu Val Asn Pro Ser Asn Arg Asn Gln Glu Thr Ser
 Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr Tyr Ser Ser Phe
 Lys Trp Arg Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile Tyr Lys
 Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp Tyr Leu
 Met Phe Ala Asp Leu Asp Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly Lys Trp
 Tyr Val Asn Thr Thr Asn Val Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr Ser Phe Phe
 Pro Asp Trp Leu Thr Tyr Val Arg Asn Gln Thr Gly Lys Asn Leu Phe Ala Val Gly Glu Phe Trp Ser
 Tyr Asp Val Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu Phe Asp Ala Pro
 Leu His Asn Asn Phe Tyr Ile Ala Ser Lys Ser Ser Gly Tyr Phe Asp Met Arg Tyr Leu Leu Asn Asn
 Thr Leu Met Lys Asp Gln Pro Ser Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln Pro Gly Gln
 Ser Leu Gln Ser Trp Val Glu Ala Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Gln Glu
 Gly Tyr Pro Cys Val Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Gly Leu Lys Ser
 Lys Ile Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln Arg Asp Tyr Ile Asp His
 Gln Asp Ile Ile Gly Trp Thr Arg Gln Gly Ile Asp Ala Lys Pro Asn Ser Gly Leu Ala Ala Leu Ile Thr
 Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Lys His Ala Gly Lys Val Phe Tyr Asp Leu
 Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ala Asp Gly Trp Gly Glu Phe Lys Val Asn Gly Gly
 Ser Val Ser Ile Trp Val Ala Lys Thr Ser Asn Val Thr Phe Thr Val Asn Asn Ala Thr Thr Thr Ser
 Gly Gln Asn Val Tyr Val Val Gly Asn Ile Pro Glu Leu Gly Asn Ser Leu

SEQ ID NO: 135

gtgacaggcaccctgtttatacttccacataaaaataaccatacagcttcaaatgttgaaatgtataaaaataaaaatagtattgtaagc
 gttaacatccgcatataataactcaaacgcgtttatgttttaagcaaacgttgcactctctttttaaagaaaggatgtgtgcatgaattat
 tgaaaaagtggtgttattacgctatcgtcgtacctaatacttcttcttacgccctttcaactgcacaagccaacactgcaccagtcacg
 gaacgatgatgcaatatttgaatgggaattaccgaatgatggcacactttggacgaaagtaaaaaacgaagcaagcagccttcttcttaggtat
 tactgcgttatgttaccacctgcatacaaaaggaacgagccaaggggatgtcgggtatggcgtgtacgattttagtacttaggagaatttaac
 aaaaaggacgattcgaacgaaatcggacaaaaacgcaatatttacaagccattcaagcggcaaaaagcgtcggcatgcaagtatacgtg
 atgtcgtatttaatacacaaggcgggggcagatagtcagaatgggttgcgcagtcgaagtgaatccttcaatcgaaccaagaacatctgg

FIGURE 16CCC

cacatacaaatcaagcatggacaaaatttgattccctgaccgtgggaacacatactcaagctttaaatggcgctggtatcatttgacggtag
 gattgggatgaaagtcgaaaactaatcgattacaaatttcgtggcagcaggaaagcatgggattgggaagtagacacagagaacggaaac
 tatgactacttaattgttgcgatttagatatggatcacccgaaagtcgtgacagagclaaaaactggggaacatggtacgtcaatagacaaaatg
 tcgatgggttgccttagatgcagtaaaagcatataaataatagcttttccagattggttaacatatgtgcgtcacaaacacaaaaaatctgttg
 cagtaggagaattttggagctacgatgtcaataaactgcataactacattacaaaaacaagtggaacctatgctgtatttgatgcgccactcataa
 caacttttacactgcttcaaaactagcgggtattttgacatgcgctatttggtaataataacgttgatgaagaccagccttctcttgcggtcacactc
 gttgataatcatgacacgcaaccgggacaactcttacaatcatgggtagagccttgggttaagccgcttgcttatgctttattttgacaagacaaga
 aggataccttgcgtattttacggcgactattacggcatccctaaatacaataatccgggattgaaaagtaaaatcgatccgcttctcatlgtcccgta
 gagactacgcatacggaaacacaacgtgattatattgaccatcaagacattatggatggacacgggaaggaaatgactcaaaacgaactctgg
 acttgcggctttaattactgacggctcgtggggaagtaaatggatgtatgtaggtaaaaagcatgctggaaggtgtttacgaatcactggaat
 cgaagcgatacggtaacgattaatgcagacggctggggagagtttaagtaaacgggtggcctccgttccatttgggtggccaaacatcacaag
 tcacgtttaccgtcaacaatgcgacaacgacaagcggacaaaatgtgtatgtcgttggcaacattccagagctcggaaattggaaacagcaaa
 cgcaatcaaaatgaccccatcttctatccaacgttgaaaacaacattgctcttccacaaggaaaagcaattggcggcgtagccatggccctt
 ga

SEQ ID NO: 136

Val Thr Gly Thr Pro Ser Leu Tyr Ile Pro Pro His Lys Ile Thr Ile Gln Leu Ser Asn Leu Leu Lys Cys
 Ile Lys Ile Lys Asn Ser Ile Val Ser Val Asn Ile Arg His Tyr Asn Asn Phe Lys Arg Val Tyr Val Leu
 Met Gln Thr Phe Ala Ser Ser Phe Tyr Leu Lys Lys Gly Cys Val Cys Met Asn Tyr Leu Lys Lys Val
 Trp Leu Tyr Tyr Ala Ile Val Ala Thr Leu Ile Ile Ser Phe Leu Thr Pro Phe Ser Thr Ala Gln Ala Asn
 Thr Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp Leu Pro Asn Asp Gly Thr Leu
 Trp Thr Lys Val Lys Asn Glu Ala Ser Ser Leu Ser Ser Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro
 Ala Tyr Lys Gly Thr Ser Gln Gly Asp Val Gly Tyr Gly Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe
 Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly Thr Lys Thr Gln Tyr Leu Gln Ala Ile Gln Ala Ala
 Lys Ser Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala Asp Ser Thr Glu
 Trp Val Asp Ala Val Glu Val Asn Pro Ser Asn Arg Asn Gln Glu Thr Ser Gly Thr Tyr Gln Ile Gln
 Ala Trp Thr Lys Phe Asp Phe Pro Asp Arg Gly Asn Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His
 Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile Tyr Lys Phe Arg Gly Thr Gly Lys
 Ala Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Phe Ala Asp Leu Asp
 Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly Thr Trp Tyr Val Asn Thr Thr Asn Val
 Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr Ser Phe Phe Pro Asp Trp Leu Thr Tyr Val
 Arg Ser Gln Thr Gln Lys Asn Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Val Asn Lys Leu His
 Asn Tyr Ile Thr Lys Thr Ser Gly Thr Met Ser Leu Phe Asp Ala Pro Leu His Asn Asn Phe Tyr Thr
 Ala Ser Lys Ser Ser Gly Tyr Phe Asp Met Arg Tyr Leu Leu Asn Asn Thr Leu Met Lys Asp Gln
 Pro Ser Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln Pro Gly Gln Ser Leu Gln Ser Trp Val
 Glu Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe
 Tyr Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Gly Leu Lys Ser Lys Ile Asp Pro Leu Leu Ile
 Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln Arg Asp Tyr Ile Asp His Gln Asp Ile Ile Gly Trp Thr
 Arg Glu Gly Ile Asp Ser Lys Pro Asn Ser Gly Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser
 Lys Trp Met Tyr Val Gly Lys Lys His Ala Gly Lys Val Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp
 Thr Val Thr Ile Asn Ala Asp Gly Trp Gly Glu Phe Lys Val Asn Gly Gly Ser Val Ser Ile Trp Val
 Ala Lys Thr Ser Gln Val Thr Phe Thr Val Asn Asn Ala Thr Thr Thr Ser Gly Gln Asn Val Tyr Val
 Val Gly Asn Ile Pro Glu Leu Gly Asn Trp Asn Thr Ala Asn Ala Ile Lys Met Thr Pro Ser Ser Tyr
 Pro Thr Trp Lys Thr Thr Ile Ala Leu Pro Gln Gly Lys Ala Ile Gly Gly Val Arg His Gly Pro

SEQ ID NO: 137

gtgggacggggcaggcttggcgcatcactcgaacacttccgccaaggggacatacgggtcacctctcgaactcgcgtccggatcgcccgccgt
 ggcggggggcggtcagcttgaagatgtccagcggggagccgcccggaggatcaccccgcgcgctactcgccaggggcggggctcag
 cttagacggcggtggcggagccgctccagggagccagacgttggaggcccgccggatggcggtcgaaggagggtggcggtcggggctgt
 tctcgtactgcagacggcggtctcggaccagcggcggtcttccaggggcggggaaccggcgggccacccggcgccgggcttccggca

FIGURE 16DDD

ggggccgggggtgatcgtccgctgcgccggcgtgggagtcagtgggtcgcgcccggggtgtcgtccgccacctgaagccgcgggtgtcgtggtcc
gggggatgccgtagtatgataccgctcgcggagatcgaccagaccggacagccgccctcctggaagcgcgggtgcgcccgggcggtgtccga
agaagaacacctcttggcgggtgttgcggaggaaccgctaccgatcacgtcgggaacagccggccagccagggaccgcagggcgaag
acgtagaggtcggccgcgagagtggagcgtccgaaaggtgaagccgtccaaaggcccgggaccatggcggccttccgggtactcccc
gccccctgccctggaacagctccaccgggtccggcagggcgcgccggcggaacagggcgccggcttctctctgtaccagatcgtcgggac
ggcgtcgaatcgacctgggggaagcgggtccggccctccccctgagacagctcggcgaccggcagccccgcgtctccagaaaaaggaa
gggagtcgcggacgtagctgtcgtcctcgcgcacatccagaggacccgggtcctttgtacagccggtaaccggactggacttcggcgtccc
gccagagctcgaaggagcgggagcaccactccacgtacagacggtcgggtccgtaggcgcgggagatccgcgtctcggaccggag
ctggagcgggagtgtcccggaacccaggcgtccaggagggtcaccgggtccgcggcgaggagatcgaggcggtccagccggcg
aaggcgcgggcgccgacgacggcgatatgggagtgaggaggcatggcgggcgtaaaggtatcgagcccgatcttctgctggcatcccat
ctccgaccggagtatcttggaaaattcgaagaaggagatcgacatgcaatcgaaaggaaacctga

SEO ID NO: 138

Val Gly Arg Ala Gly Leu Ala His His Ser Asn Thr Ser Ala Lys Gly Thr Tyr Gly Ser Pro Leu Glu
Leu Arg Pro Asp Arg Pro Ala Val Ala Gly Ala Val Glu Leu Glu Asp Val Gln Arg Gly Ala Ala Ala
Glu Asp His Pro Gly Gly Val Leu Ala Gln Gly Gly Ala Gln Leu Glu Ala Val Ala Gly Ala Ala Ser
Gln Glu Pro Asp Val Gly Gly Pro Arg Met Ala Val Glu Glu Glu Val Ala Val Gly Ala Val Leu Val
Leu Ala Asp Ala Gly Leu Asp Gln Arg Arg Val Leu Gln Gly Arg Glu Pro Ala Gly His Leu Gly
Pro Gly Arg Phe Gln Gln Gly Arg Gly Asp Arg Pro Leu Ala Arg Arg Gly Ile Asp Gly Leu Ala Pro
Gly Val Val Arg His Leu Glu Ala Ala Val Leu Val Ala Gly Asp Ala Val Val Asp Pro Leu Ala Glu
Ile Asp Pro Asp Arg Thr Ala Ala Leu Leu Glu Ala Arg Val Ala Arg Arg Arg Ala Glu Glu Glu His
Leu Leu Ala Gly Val Ala Glu Glu Pro Leu Thr Asp His Val Arg Glu Gln Pro Gly Gln Pro Gly Thr
Ala Gly Glu Asp Val Glu Val Gly Arg Glu Ser Gly Ala Val Arg Lys Val Lys Pro Leu Gln Gly Pro
Arg Asp His Gly Gly Leu Pro Val Leu Pro Ala Leu Ala Leu Glu Gln Leu His His Gly Pro Ala Gly
Ala Pro Gly Glu Gln Gly Ala Gly Phe Leu Leu Val Pro Asp Arg Ala Asp Ala Val Glu Ile Asp Leu
Gly Glu Ala Ala Pro Gly Leu Pro Leu Arg Gln Leu Gly Asp Arg Gln Pro Arg Val Leu Gln Lys
Arg Lys Gly Val Ala Asp Val Ala Val Val Leu Ala Ala His Pro Glu Asp Pro Gly Pro Phe Val Gln
Pro Val Thr Gly Leu Asp Phe Gly Val Pro Pro Glu Leu Glu Gly Ala Gly Asp Pro Leu His Val Gln
Thr Val Gly Ser Val Gly Ala Ala Asp Asp Pro Arg Leu Ala Thr Gly Ala Gly Ala Gly Val Pro Arg
Thr Pro Gly Val Gln Glu Gly His Pro Gly Ser Ala Ala Glu Glu Met Gln Gly Gly Pro Ala Ala Glu
Gly Ala Gly Ala Asp Asp Gly Asp Met Gly Met Gly Gly His Gly Gly Arg Lys Val Ile Ala Ala Arg
Ser Phe Ala Gly Ile Pro Ser Pro Thr Gly Val Ser Trp Lys Ile Arg Arg Arg Arg Ser Thr Cys Asn
Arg Thr Glu Thr

SEQ ID NO: 139

atgaaaacattcaaccctaaaccacacctttacctttaactttgctgtgtgaggttgcgcgtatftggcgggcacaaaatggaaetaatgatgcagatttcc
cattgggtttggcgaatgagggggcattatggaaacaaagtggaaacaaatggcgcagcactatcggaaaacgggtttacagggctgggttgc
caccagcattaaagggtgcaggttggtagaacgacgttggtaacggtttatogatatgtaigacttggggagtttgatcaaaaaaggtcggta
cgaactaagtacggcaccaaaagaccaataatataatgccatcaaaagcagcacacaaaaacaatatccaaatttatggtagcgtatggttcaacca
tcgtggcgggtgcagatggcgaagtcgtgggtcgataccaagcgtgttgattggaataaccgcaatttgaacttggcgataaatggattgaagca
tgggttgaaatttagcttccaggacgtaacgataaatactcagacttcatttggacgtggatcactttgatggcgctgattgggatgacgcaggta
aagagaaagcgtatctttaaattcaaaagggtgaggttaaagcatgggattgggaagtcagttcgaaaaaggcaactatgactacctcatgtacgca
gacttagacatggatcaccagaagtgaagcaagagctgaagattgggtgaatggtaacttaaacatgacgggtgttgaaggcttccgaatgg
atgcagtgaaacacatcaaatatcagtaacctacaagaatggatcgattacttgcgtaagaaaacgggcaaaagagctcttaccgttggtagtac
tggaaactacgacgtgaacaatctgcacaactttatgactaagacttctggcagcatgtcattgttgatgcgccttacatatgaacttctataacgct
tcacgctctgttggcaactttgatatgcgccgaatcatggatggcaacttgatgaaagacaaccacgtgaaagcagtaacactgggtgagaacc
atgatacgcaaacactacaggccttagagctccgggtgattgtgtgttcaaacaccatitgcgtlacgcgttcatittgcttcgtgaggaaggttatcc
gtcagctctctacgcgattactacgggtgcgaatacagcgataaaagggcacgatatcaacatgggtgaaagtgccttacattgagcaattgggtga
aagcgcgtaaagattatgcttatgttaaacaacattcttacccttgaccactgggatgtgattgggttggcacacggaagggggtatgcggaacatccg

FIGURE 16EEE

aactctatggcggttatcatgagtgatggctctggcggacaagtggatgtacacaggttcaccgagcacacgttatgtcgataaactaggtatt
cgtaccgaagaagtaaggactaacgctagtggatgggccgaattcccgtagaacggcggatcggttctgtttgggttggcgtaaataa

SEQ ID NO: 140

Met Lys Thr Phe Asn Leu Lys Pro Thr Leu Leu Pro Leu Thr Leu Leu Ser Ser Pro Val Leu Ala
Ala Gln Asn Gly Thr Met Met Gln Tyr Phe Leu Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln
Val Glu Asn Asn Ala Pro Ala Leu Ser Asp Asn Gly Phe Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys
Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp
Gln Lys Gly Ser Val Arg Thr Lys Tyr Gly Thr Lys Asp Gln Tyr Leu Asn Ala Ile Lys Ala Ala His
Lys Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His Arg Gly Gly Ala Asp Gly Lys Ser Trp
Val Asp Thr Lys Arg Val Asp Trp Asn Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp
Val Glu Phe Ser Phe Pro Gly Arg Asn Asp Lys Tyr Ser Asp Phe His Trp Thr Trp Tyr His Phe Asp
Gly Val Asp Trp Asp Asp Ala Gly Lys Glu Lys Ala Ile Phe Lys Phe Lys Gly Asp Gly Lys Ala Trp
Asp Trp Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp
His Pro Glu Val Lys Gln Glu Leu Lys Asp Trp Gly Glu Trp Tyr Leu Asn Met Thr Gly Val Asp Gly
Phe Arg Met Asp Ala Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp Tyr Leu Arg Lys
Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn Tyr Asp Val Asn Asn Leu His Asn
Phe Met Thr Lys Thr Ser Gly Ser Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala
Ser Arg Ser Gly Gly Asn Phe Asp Met Arg Arg Ile Met Asp Gly Thr Leu Met Lys Asp Asn Pro
Val Lys Ala Val Thr Leu Val Glu Asn His Asp Thr Gln Pro Leu Gln Ala Leu Glu Ser Pro Val Asp
Trp Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser Val Phe Tyr
Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly His Asp Ile Asn Met Val Lys Val Pro Tyr Ile
Glu Gln Leu Val Lys Ala Arg Lys Asp Tyr Ala Tyr Gly Lys Gln His Ser Tyr Leu Asp His Trp Asp
Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser Met Ala Val Ile Met Ser Asp Gly
Pro Gly Gly Thr Lys Trp Met Tyr Thr Gly Ser Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg
Thr Glu Glu Val Trp Thr Asn Ala Ser Gly Trp Ala Glu Phe Pro Val Asn Gly Gly Ser Val Ser Val
Trp Val Gly Val Lys

SEQ ID NO: 141

[illegible]

SEQ ID NO: 142

Met Lys Pro Ile Asn Thr Leu Leu Ile Ser Ala Leu Ala Val Cys Ser Phe Ser Ser Ala Thr Tyr Ala
Asp Thr Ile Leu His Ala Phe Asn Trp Lys Tyr Ser Asp Val Thr Ala Asn Ala Asn Gln Ile Ala Gln
Ala Gly Tyr Lys Lys Val Leu Val Ala Pro Ala Met Lys Ser Ser Gly Ser Gln Trp Trp Ala Arg Tyr
Gln Pro Gln Asp Leu Arg Thr Ile Asp Ser Pro Leu Gly Asn Lys Gln Asp Leu Ala Ala Met Ile Ala

FIGURE 16FFF

Ala Leu Lys Gly Val Gly Val Asp Val Tyr Ala Asp Val Val Leu Asn His Met Ala Asn Glu Ser Trp
 Lys Arg Ser Asp Leu Asn Tyr Pro Gly Thr Glu Val Leu Asn Asp Tyr Ala Ser Arg Ser Ser Tyr Tyr
 Ala Asp Gln Thr Leu Phe Gly Asn Leu Ala Gln Gly Tyr Val Ser Ala Asn Asp Phe His Pro Ala Gly
 Cys Ile Ser Asp Trp Asn Asp Pro Gly His Val Gln Tyr Trp Arg Leu Cys Gly Ala Asp Gly Asp Val
 Gly Leu Pro Asp Leu Asp Pro Asn Asn Trp Val Val Ser Gln Gln Arg Leu Tyr Leu Lys Ala Leu
 Lys Asp Met Gly Ile Lys Gly Phe Arg Ile Asp Ala Val Lys His Met Ser Gln Tyr Gln Ile Asp Gln
 Val Phe Thr Ser Glu Ile Thr Ala Asn Met His Val Phe Gly Glu Val Ile Thr Ser Gly Gly Ala Gly
 Asn Ser Gly Tyr Glu Ser Phe Leu Ala Pro Tyr Leu Asn Asn Thr Asn His Ser Ala Tyr Asp Phe Pro
 Leu Phe Ala Ser Ile Arg Ser Ala Phe Ser Met Gly Gly Gly Leu Asn Gln Leu His Asp Pro Lys Ala
 Tyr Gly Gln Ala Leu Asp Asp Asn Arg Ser Ile Thr Phe Ala Ile Thr His Asp Ile Pro Thr Asn Asp
 Gly Phe Arg Tyr Gln Ile Met Asp Pro Gln Asp Glu Gln Leu Ala Tyr Ala Tyr Ile Leu Gly Lys Asp
 Gly Gly Thr Pro Leu Ile Tyr Ser Asp Asp Leu Pro Asp Ser Glu Asp Lys Asp Asn Gly Arg Trp Gly
 Asn Val Trp Asn Ser Ser Thr Met Lys Asn Met Leu Ser Phe His Asn Ala Met Gln Gly Lys Thr
 Met Thr Met Ile Ser Ser Asp His Cys Thr Leu Leu Phe Lys Arg Gly Lys Glu Gly Val Val Gly Ile
 Asn Lys Cys Gly Glu Thr Arg Gly Val Thr Val Asp Thr Tyr Gln His Glu Phe Asn Trp His Val Gln
 Tyr Lys Asp Val Leu Ser Ser Ala Thr Glu Thr Val Thr Ser Arg Tyr His Thr Phe Asn Leu Pro Pro
 Arg Ser Ala Arg Met Phe Lys Leu

SEQ ID NO: 143

atgccaaagagcacttttaccacaaatccataacaaatcacttcttgctactccgttggtgtaagcttattgcctgcctacgcacaggccgacactat
 ctgcatgccttaactggaaatacagcgacattaccgccaagcagagcaaatgcgcaagctggtataaaaaagtactgatttaccgccgc
 tgaagtcacagggccacaaatggtggcgacgttaccacacaggaacatcagtgattgactccctgtcggcaacaagcaagattacaag
 ccctcattgcagcctaaaggcacaaggcggtgaagtatacgcagacatcgactacacacatggccaacgaaagctggaaacgagacgatc
 tgaactaccgggaagtgtatttacttaccacaaatagcttacctgaaccagcaaaatggttgagattagagcaaaatcagtt
 ctctgcaatgatttaccggctggtgctgacttactgattgagtaacccgggcatgttcaactactggcgcttatgtggtggaatggtgacact
 gggttacctgatcttgcctaaactcgtgggtgatcgaacaaaaacgtatttactgcttgaagacatgggaataaagggttccgagttg
 atgcggtaaaacacatgagcgattaccacaaatcaaccaaggtttacgccagacatcatcgaggcttacctgatttggtagtgatcaccagtg
 gtggcaaggcgagcaatgactaccactctttctggaaccgtatttaataacacaaatcacgccgcgtatgacttccgctatttgccttatccg
 aaatgcatttagttatcatgagcgttgcctcaattacatgatccacaagcttacgggcaagcacttctaacgacagagccattacttaccatca
 ctacgacattccaaccaatgatgttccgttaccacaaatcatggatccaaccagtgaaaaactcgcgtacgcgtacatttaggcaaatgagg
 ggtagccacttactatagcgatgctttgacccaagtgaagataaagataaggccgctggcgatgtatggaaccaagaatacatggttaa
 catgatcagcttccacaacaagggtgaaggtaaaagcatggaggtcatgtacagcgatcaatgcttgcgttcttaaacgtgaaaaacaaggct
 tagtcggtattaataagtgctgaaagccgtacctaaccatagataccatcggtttgaatttaactgttaccacacgtacaacgacacattaag
 ccagcacagcgagacctttagcagccgttatcatgctctgaccattccggcgcaaacagcacgaatgttggcgctataa

SEQ ID NO: 144

Met Pro Lys Ser Thr Phe Thr Lys Ser Ile Thr Lys Ser Leu Leu Ala Thr Ser Val Val Val Ser Leu
 Leu Pro Ala Tyr Ala Gln Ala Asp Thr Ile Leu His Ala Phe Asn Trp Lys Tyr Ser Asp Ile Thr Arg
 Gln Ala Glu Gln Ile Ala Gln Ala Gly Tyr Lys Lys Val Leu Ile Ser Pro Pro Leu Lys Ser Thr Gly Pro
 Gln Trp Trp Ala Arg Tyr Gln Pro Gln Asp Ile Arg Val Ile Asp Ser Pro Val Gly Asn Lys Gln Asp
 Leu Gln Ala Leu Ile Ala Ala Leu Lys Ala Gln Gly Val Glu Val Tyr Ala Asp Ile Val Leu Asn His
 Met Ala Asn Glu Ser Trp Lys Arg Asp Asp Leu Asn Tyr Pro Gly Ser Asp Leu Leu Thr Gln Tyr Ser
 Gln Asn Met Ala Tyr Met Asn Gln Gln Lys Leu Phe Gly Asp Leu Glu Gln Asn Gln Phe Ser Ala
 Asn Asp Phe His Pro Ala Gly Cys Ile Thr Asp Trp Ser Asn Pro Gly His Val Gln Tyr Trp Arg Leu
 Cys Gly Gly Asn Gly Asp Thr Gly Leu Pro Asp Leu Asp Pro Asn Ser Trp Val Ile Asp Gln Gln Lys
 Arg Tyr Leu Arg Ala Leu Lys Asp Met Gly Ile Lys Gly Phe Arg Val Asp Ala Val Lys His Met Ser
 Asp Tyr Gln Ile Asn Gln Val Phe Thr Pro Asp Ile Ile Ala Gly Leu His Val Phe Gly Glu Val Ile Thr
 Ser Gly Gly Lys Gly Ser Asn Asp Tyr His Ser Phe Leu Glu Pro Tyr Leu Asn Asn Thr Asn His Ala
 Ala Tyr Asp Phe Pro Leu Phe Ala Ser Ile Arg Asn Ala Phe Ser Tyr His Gly Ser Leu Ser Gln Leu
 His Asp Pro Gln Ala Tyr Gly Gln Ala Leu Pro Asn Asp Arg Ala Ile Thr Phe Thr Ile Thr His Asp

FIGURE 16GGG

Ile Pro Thr Asn Asp Gly Phe Arg Tyr Gln Ile Met Asp Pro Thr Ser Glu Lys Leu Ala Tyr Ala Tyr
 Ile Leu Gly Lys Asp Gly Gly Ser Pro Leu Ile Tyr Ser Asp Ala Leu Asp Pro Ser Glu Asp Lys Asp
 Lys Gly Arg Trp Arg Asp Val Trp Asn Gln Glu Tyr Met Val Asn Met Ile Ser Phe His Asn Lys Val
 Gln Gly Lys Ser Met Glu Val Met Tyr Ser Asp Gln Cys Leu Leu Val Phe Lys Arg Glu Lys Gln
 Gly Leu Val Gly Ile Asn Lys Cys Ala Glu Ser Arg Thr Tyr Thr Ile Asp Thr His Arg Phe Glu Phe
 Asn Trp Tyr Gln Pro Tyr Asn Asp Thr Leu Ser Gln His Ser Glu Thr Phe Ser Ser Arg Tyr His Ala
 Leu Thr Ile Pro Ala Gln Thr Ala Arg Met Leu Ala Leu

SEQ ID NO: 145

atgttgaaaaggattacggtagctctgtttattatttttgccttttccataatataatggagggaataaggcgggaagcagcaacgataataatgga
 acattaatgcagttattttagtggtacgctccgaatgatgggaatcatgttggaatcggttgctatgatgctgaaagtttagctcataagggaatcac
 atctgtatggataccacctgcataataagggaacttcgcaaaatgatgtagggtatggggcctatgattatacagtttaggggaagttcaatcaaaaa
 ggaacgggtgcggacgaaatatgggacaaaggcacagttgaaatctgcaattgacgctttacataagcaaaacatcgacgtataggtgatgta
 gttatgaatcataaagggtgggctgattatactgaaaccgtaacagctgttgaggtagaccgtaacaatcgaaatattgaagtatcagggtgattatg
 aaattagtgctggacgggttttaacttccaggggcgagagatgctatttcaatttcaaatggaaatggatcattttgacggaacggattgggat
 gaaggaaggaataaaccgaattataaatttaggggtataggtaaagcgtgggactgggaagtgtcagcgaaatggaaattatgattattg
 atgtatgcagatcttatttgcacatccagatgttgcgaatgaaatgaaaagttggggaacgtggtatgcgaatgaataaatttagatggattcgt
 ttatagctgttaaacatattgatcatgaatatttacgcgattgggttaaactgtcagacagcaaacggggaagaaatgtttacgggtgctgaat
 attggcaaatgataccagacttaaacatatttggcgaaagtcgaattataatcaatctgtatttgatgcaccgcttcattacaatttcattatgctt
 caacaggaaatgggaattatgatatgagaaatattttaaattggaacagtaataaaaaatcctgcactgcagttactctcgttgagaatcatga
 ttctcaacctgggcaatcattggaatctgtagtaagtcggtgttttaagccgctggcatatgcatttttaactcgtgcagagggtatccttcagt
 ttttatgggtgattactatgggacaagcggaatagtagttatgaaatccagcgttaaaagataaaatgatccaattttgacggcacgaaaaaact
 ttgcataatggtagcgcggtgattatttagaccatccagatgtgattggctggacaagagaaggagatagtgatcatgtaagctggttagcgg
 cattaatciccgatggaccaggagatcaaatggatggatgttgaaagaataacgctggggaagtatggtagatattacgggtaatacaaac
 aaatactgtaacaataataaagatggatcggggcaattccatgtaagtggaggctctgtttctatatattgttaacagtaa

SEQ ID NO: 146

Met Leu Lys Arg Ile Thr Val Val Cys Leu Leu Phe Ile Leu Leu Phe Pro Asn Ile Tyr Gly Arg Asn
 Lys Ala Glu Ala Ala Thr Ile Asn Asn Gly Thr Leu Met Gln Tyr Phe Glu Trp Tyr Ala Pro Asn Asp
 Gly Asn His Trp Asn Arg Leu Arg Tyr Asp Ala Glu Ser Leu Ala His Lys Gly Ile Thr Ser Val Trp
 Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Asn Asp Val Gly Tyr Gly Ala Tyr Asp Leu Tyr Asp Leu
 Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Leu Lys Ser Ala Ile
 Asp Ala Leu His Lys Gln Asn Ile Asp Val Tyr Gly Asp Val Val Met Asn His Lys Gly Gly Ala Asp
 Tyr Thr Glu Thr Val Thr Ala Val Glu Val Asp Arg Asn Asn Arg Asn Ile Glu Val Ser Gly Asp Tyr
 Glu Ile Ser Ala Trp Thr Gly Phe Asn Phe Pro Gly Arg Arg Asp Ala Tyr Ser Asn Phe Lys Trp Lys
 Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Gly Arg Lys Leu Asn Arg Ile Tyr Lys Phe Arg Gly
 Ile Gly Lys Ala Trp Asp Trp Glu Val Ser Ser Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp
 Leu Asp Phe Asp His Pro Asp Val Ala Asn Glu Met Lys Ser Trp Gly Thr Trp Tyr Ala Asn Glu Leu
 Asn Leu Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Asp His Glu Tyr Leu Arg Asp Trp Val Asn
 His Val Arg Gln Gln Thr Gly Lys Glu Met Phe Thr Val Ala Glu Tyr Trp Gln Asn Asp Ile Gln Thr
 Leu Asn Asn Tyr Leu Ala Lys Val Asn Tyr Asn Gln Ser Val Phe Asp Ala Pro Leu His Tyr Asn
 Phe His Tyr Ala Ser Thr Gly Asn Gly Asn Tyr Asp Met Arg Asn Ile Leu Asn Gly Thr Val Met Lys
 Asn His Pro Ala Leu Ala Val Thr Leu Val Glu Asn His Asp Ser Gln Pro Gly Gln Ser Leu Glu Ser
 Val Val Ser Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Ala Glu Gly Tyr Pro Ser
 Val Phe Tyr Gly Asp Tyr Tyr Gly Thr Ser Gly Asn Ser Ser Tyr Glu Ile Pro Ala Leu Lys Asp Lys
 Ile Asp Pro Ile Leu Thr Ala Arg Lys Asn Phe Ala Tyr Gly Thr Gln Arg Asp Tyr Leu Asp His Pro
 Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ser Val His Ala Lys Ser Gly Leu Ala Ala Leu Ile Ser
 Asp Gly Pro Gly Gly Ser Lys Trp Met Asp Val Gly Lys Asn Asn Ala Gly Glu Val Trp Tyr Asp Ile
 Thr Gly Asn Gln Thr Asn Thr Val Thr Ile Asn Lys Asp Gly Ser Gly Gln Phe His Val Ser Gly Gly
 Ser Val Ser Ile Tyr Val Gln Gln

FIGURE 16HHH

SEQ ID NO: 147

atgagcttaaataactttaaggtaaactgcttagtttgcgtgtcttctgctgtattgtcactggctccaaatttagccaatgctgcaaattttgaag
 tgagatgggtgataatccatccgtttcagtgagacatatgacaatatagcaaaagagtgtacagagtaccttggccagccggatttgacgggtgaca
 gatttccagccagcgggaacataagcgggctgaaggagtagtggtggccgtatatcagccggtaattataagaattttacaacatgaccggta
 acgaggagcagcttaaggcaatgatcaagacctgtaatgatgcagggttaagggttcgctgacgctgttttcaacaaaaggctacagacgg
 ttaggctggggcgggttaacttgagttataagaactaccctgacggatctccggatcagatttccatggagactgttccatgacaaaagctat
 actgatgcaataatgtcagaacctgtgcactcaggtatgccggacgttgccacagataactccgctactcaggaaaagattgcagattacct
 cgcttcttaataatgaatatgggggtctatgtttccgtattgacgctgcgaagcacatgggatacaacgatcaactccattctttcaaaaactgcac
 agaagactggaagaagacctcctgcatactggaagtaactggagccggtaacgaagctgccgacattcagccggacaagatactttattga
 gaatgcgggtgtaactgacttcgggtatgtctgggatgcaaatgagagttcggaaagggttaattacggtaaggcactggaactcagttacgtgct
 cgggtgcaaatcagaacattcgtaaacaatcatgatgatgaatggggcagatgctcagccggtagctgctcaatgaaaactcagaattatgctg
 attataatctggctcagctcgtgctgtatggcctgtaggtagacagtaagacagatatattccgggtattcattccctgtaaaagataatgatcctta
 tcgctcagtgatgcaactcatgatcagggcgggctcttgggtgccgaccgctgtgaagggtggtggtgtgacgaccggtgtgtcttctgttct
 caattccccagatttgcgagagctaccagaggtactgctgtatcaaccaagggttgcgaatggtgcttgggttaacagagggaagcaag
 gttttatgcacagaatactaccaacagctctataaccagacattctgttgaagtacgtgacggaattactgtgatacttaggaacatcagat
 cctaaagagcaatccatgcggagcagagcgtgtcgtgaagcggcggtgaaggctacatttactattcctgcaaaagacagctgtggctatctgtacaga
 ctacagactggtgcggcaagggttgatccttgaagtgtacggcgggtgctgctgtgttgaagggggaaccaccgttaattggtgtgt
 gcgtcagctggtgtaatgcgcattcatcaaatgaggaaatgcacctgtgtattgaatccgaatgatccaactgtcaggctgataattgaacctacca
 agggtaaacctgtttacgcccgtacttcaaacgggtggaacaggatccttaacataataaccgtaaaacagggttcttgactattaatctgactctt
 gacgggtgcaggtgataccagcggagctcagcgttcaagggttacagcggatgttcatggaccggaacagttacgggttctcaggtactgccg
 gaaaggttgatgtaaaatcatcatcaaccggcgatgaacctgtgtcttctgtgtgattatgttcttccattaaacgataagaccatggaatatacat
 tcaccaaggcagatgaagtaactaatcagccaccggtgcatcattaccgcgacagttaacggctgaccgttcttttccaataattcatccga
 ccctgagaatgatgaattaacctacagctggaattcggtaatggtataaacatcatccgagaagctcctgataacctatgaagaatccggta
 agtatactgttactttaagggttactgattcagtaataacactgatacatttactaaagataataactgtaacagcaccttctagtgccaagtactttaa
 ggtgcagtcagaggttcgcatgataattacggaactgatctgttaaccaagaacgggtctgattggaccggcgtcttgaattcttggatccacta
 gtgtgcacctgcagggcgcgcgagctc

SEQ ID NO: 148

Met Ser Leu Asn Asn Phe Lys Val Lys Leu Leu Ser Phe Ala Val Ser Ser Ala Val Leu Ser Leu Ala
 Pro Asn Leu Ala Asn Ala Ala Asn Phe Glu Ser Glu Met Val Ile Ile His Pro Phe Gln Trp Thr Tyr
 Asp Asn Ile Ala Lys Glu Cys Thr Glu Tyr Leu Gly Pro Ala Gly Phe Asp Gly Val Gln Ile Ser Gln
 Pro Ala Glu His Lys Arg Ala Glu Gly Val Trp Trp Ala Val Tyr Gln Pro Val Asn Tyr Lys Asn Phe
 Thr Thr Met Thr Gly Asn Glu Glu Gln Leu Lys Ala Met Ile Lys Thr Cys Asn Asp Ala Gly Val Lys
 Val Phe Ala Asp Ala Val Phe Asn Gln Lys Ala Thr Asp Gly Val Gly Trp Gly Gly Ser Thr Trp Ser
 Tyr Lys Asn Tyr Pro Asp Gly Phe Ser Gly Ser Asp Phe His Gly Asp Cys Ser Ile Asp Lys Ser Tyr
 Thr Asp Ala Asn Asn Val Arg Thr Cys Ala Leu Ser Gly Met Pro Asp Val Ala Thr Asp Asn Ser Ala
 Thr Gln Glu Lys Ile Ala Asp Tyr Leu Ala Ser Leu Met Asn Met Gly Val Tyr Gly Phe Arg Ile Asp
 Ala Ala Lys His Met Gly Tyr Asn Asp Ile Asn Ser Ile Leu Ser Lys Thr Ala Gln Lys Thr Gly Arg
 Arg Pro Pro Ala Tyr Leu Glu Val Ile Gly Ala Gly Asn Glu Ala Ala Asp Ile Gln Pro Asp Lys Tyr
 Thr Phe Ile Glu Asn Ala Val Val Thr Asp Phe Gly Tyr Val Trp Asp Ala Asn Glu Ser Phe Gly Lys
 Gly Asn Tyr Gly Lys Ala Leu Glu Leu Ser Thr Trp Leu Gly Ala Asn Ser Glu Thr Phe Val Asn Asn
 His Asp Asp Glu Trp Gly Arg Cys Ser Ala Gly Ser Cys Ser Met Lys Thr Gln Asn Tyr Ala Asp Tyr
 Asn Leu Ala Gln Ser Trp Leu Ala Val Trp Pro Val Gly Thr Val Arg Gln Ile Tyr Ser Gly Tyr Ser
 Phe Pro Val Lys Asp Asn Asp Pro Tyr Arg Val Ser Asp Ala Thr His Asp Gln Gly Gly Pro Leu Gly
 Ala Asp Arg Cys Glu Gly Gly Trp Leu Cys Gln His Arg Val Ser Phe Val Leu Asn Ser Pro Arg Phe
 Ala Arg Ala Thr Arg Gly Thr Ala Val Ser Thr Lys Gly Phe Asp Asn Gly Ala Leu Trp Phe Asn Arg
 Gly Ser Lys Gly Phe Tyr Ala Gln Asn Thr Thr Asn Ser Pro Ile Thr Gln Thr Phe Ser Val Glu Val
 Pro Asp Gly Asn Tyr Cys Asp Ile Leu Gly Thr Ser Asp Pro Lys Ser Asn Pro Cys Gly Ala Asp Val
 Val Val Ser Gly Gly Lys Ala Thr Phe Thr Ile Pro Ala Lys Thr Ala Val Ala Ile Cys Thr Asp Ser

FIGURE 16III

Asp Trp Cys Gly Lys Gly Val Asp Pro Cys Glu Ser Asp Pro Thr Gly Ala Ala Cys Val Cys Lys Gly
 Glu Thr Thr Val Asn Gly Val Cys Val Ser Trp Cys Asn Ala His Ser Ser Asn Glu Glu Cys Thr Cys
 Val Leu Asn Pro Asn Asp Ala Asn Cys Gln Ala Asp Ile Glu Pro Thr Lys Gly Lys Leu Cys Tyr Ala
 Gly Thr Ser Asn Gly Trp Lys Gln Asp Pro Leu Thr Tyr Asn Arg Lys Thr Gly Phe Trp Thr Ile Asn
 Leu Thr Leu Asp Gly Ala Gly Asp Thr Ser Gly Ala Gln Arg Phe Lys Val Thr Asp Gly Cys Ser Trp
 Thr Gly Thr Val Tyr Gly Ser Ser Gly Thr Ala Gly Lys Leu Asp Val Asn Thr Ser Ser Thr Gly Asp
 Glu Pro Val Ser Leu Val Gly Asp Tyr Val Leu Ser Ile Asn Asp Lys Thr Met Glu Tyr Thr Phe Thr
 Lys Ala Asp Glu Val Thr Asn Gln Pro Pro Val Ala Ser Phe Thr Ala Thr Val Asn Gly Leu Thr Val
 Ser Phe Ala Asn Asn Ser Ser Asp Pro Glu Asn Asp Glu Leu Thr Tyr Ser Trp Asn Phe Gly Asn Gly
 Lys Thr Ser Ser Glu Lys Ala Pro Ser Ile Thr Tyr Glu Glu Ser Gly Lys Tyr Thr Val Thr Leu Lys
 Val Thr Asp Ser Ala Asn Asn Thr Asp Thr Phe Thr Lys Asp Ile Thr Val Thr Ala Pro Ser Ser Gly
 Lys Tyr Leu Lys Val Ala Val Arg Gly Ser His Asp Asn Tyr Gly Thr Asp Leu Leu Thr Lys Asn Gly
 Ser Asp Trp Thr Gly Val Phe Glu Phe Phe Gly Ser Thr Ser Val Asp Leu Gln Ala Arg Glu Leu

SEQ ID NO: 149

atgacctaagtaatttaaggtaaaactcttagtttctgtgtcttctgtgtactgacactggctgcaaatgtcgccaatgccaagaattatgaaa
 gtgaaatgggtattatcattcattcagtgacatgacaatagcaaaagaatgtactgagtatctgggacctgcccggattgacggggtgca
 gatttcccaggcggctgagcataaagatgccggtggtgcatggtgggtaccctaccagcctgtaaactcaagagtttactaccatggttgta
 atgaagaacagcttagagcaatgattaaacctgtaacgaggcaggtgttaagggtttgccgatgccgtgtaaatcagaagccggcgacgg
 ttaggtataggtgttcaacttccgaaattataattatcctgacggatttaccagtgatgatttcatcataaactgcagtataggttaataatt
 cagatgcatgggtagtaagattctgtgacctcagtgccatgccgatatagcaactgataacgacagtaccagaaataagattgctgattacttcg
 ccagccctatgaatatgggggtatagcgttccgtatgctgccaagcactttagctatgatgatatagacgctattgtagagaaacagcaa
 ccaaaagcaggcaggagacctcctgtctatatggagggtatcgtaalcgggtcaagaggcggatgataccagccgaacaagtatacatgga
 ttgataatgccgttgtaacagattttacttatgctaatagcatgcataatattttaacggaaagggttatgccaaggccttgaaacatggggctagggc
 atgttgatgctgaaaatgccgaagtcttataagtaatcatgataatgaatggggaagaaagctgcccgttctgtcctaataagaacccagaata
 atccggattaccatctggctcagtcctgctcgcagttggcccttaggcaagggttagacagattatctgcatacagttcccggtcttgaagata
 gttgtgagcgggtcagtcageaagcccatgatcaggcgggtcctatcggggcagcccgtgtgaaagggtggtgtgtcagcaccgtgtac
 cgtttgtgctcaattctcctagatttgcaagagcaaccagaggacagtcgttactactaaaggtttgatgacggaacttgtgttaacagagg
 aagcaagggtcttatgccagaatactaccggcagttctataactcatacattctcagttgaattacctgatgaaattactgtgataccttgga
 caaccgatccgaagaataactcttgcggagcggatgtcactgtaagcggagggttaagcaacctttaccattccggcaagaccggcgtagcta
 tctgtactgatgaaaatgtgtgtggaagggttgaccctgtgaaagcgtactaccgggtccgctgtgtatgtaagggtgaaaccacagtt
 aacggcgtatgtgtaagctggtgtaatgtcactcactcactaatgaagaatgtgcctgtgtgctaaatcctaagctgagtgagtgacggcgacatt
 gagccgaccaagggttaactctgctatgtaggtaccctcaacaaggagtcaggaaccttaacctataatgcagaccggtttctgagactct
 caacgttgaacttgacggttaaggggataccagcggggcgcagcgtttaaagttaccgcagcggctgttcagtcagggtactgtttacgggtca
 tcaggagtgaaggcagacttgacgttaatacttcagccaccggagatgaaccgggttactgacaggttaaatgttcttccataaatgataag
 accatggaatacacattccttgcaggcagtggaacaagccctccggttgcgtcattactccgactgtttaaagatctgactgtatcttgc
 taattcaccgaacctgagaatgagaaatcctcagctggaatttcggttaacggtaaaccttctggaagaatcaggtgttaataatgat
 aaagcgggttaataatctgttctcactcaagtaactgatactgcaaacacactgtaacaaaacacttggaatcgaattacatctctgttaacg
 gaaaatattccaaggttgcatgcagaggttcacatgataactacggaacaaactgttaaccagggaatggttcagaatggaccggtatcttgaatt
 cagtaagacaacaaattcaagcttgaagctctgctcctcagctgaccagtgatctctcggcggttaacgaggtgagccattgactgcct
 ccggtggatttatctctctcctgcccgaaggatataaagttaatgaggaaagcaagggttcttactgcagcggatgttactgcaccggg

SEQ ID NO: 150

Met Ile Leu Ser Asn Phe Lys Val Lys Leu Leu Ser Phe Ala Val Ser Ser Ala Val Leu Thr Leu Ala
 Ala Asn Val Ala Asn Ala Lys Asn Tyr Glu Ser Glu Met Val Ile Ile His Pro Phe Gln Trp Thr Tyr
 Asp Asn Ile Ala Lys Glu Cys Thr Glu Tyr Leu Gly Pro Ala Gly Phe Asp Gly Val Gln Ile Ser Gln
 Ala Ala Glu His Lys Asp Ala Gly Gly Ala Trp Trp Gly Thr Tyr Gln Pro Val Asn Phe Lys Ser Phe
 Thr Thr Met Val Gly Asn Glu Glu Gln Leu Arg Ala Met Ile Lys Thr Cys Asn Glu Ala Gly Val Lys
 Val Phe Ala Asp Ala Val Ile Asn Gln Lys Ala Gly Asp Gly Val Gly Ile Gly Gly Ser Thr Phe Gly
 Asn Tyr Asn Tyr Pro Asp Gly Phe Thr Ser Asp Asp Phe His His Asn Cys Ser Ile Gly Asn Asn

FIGURE 16JJJ

Tyr Ser Asp Ala Trp Val Val Arg Phe Cys Asp Leu Ser Gly Met Pro Asp Ile Ala Thr Asp Asn Asp
 Ser Thr Arg Asn Lys Ile Ala Asp Tyr Phe Ala Ser Leu Met Asn Met Gly Val Tyr Gly Phe Arg Ile
 Asp Ala Ala Lys His Phe Ser Tyr Asp Asp Ile Asp Ala Ile Val Glu Lys Thr Ala Thr Lys Ala Gly
 Arg Arg Pro Pro Val Tyr Met Glu Val Ile Gly Asn Pro Gly Gln Glu Ala Asp Asp Ile Gln Pro Asn
 Lys Tyr Thr Trp Ile Asp Asn Ala Val Val Thr Asp Phe Thr Tyr Ala Asn Ser Met His Asn Ile Phe
 Asn Gly Ser Gly Tyr Ala Lys Ala Leu Asn Met Gly Leu Gly His Val Asp Ala Glu Asn Ala Glu Val
 Phe Ile Ser Asn His Asp Asn Glu Trp Gly Arg Lys Ser Ala Gly Ser Cys Ser Ile Arg Thr Gln Asn
 Asn Pro Asp Tyr His Leu Ala Gln Ser Trp Leu Ala Val Trp Pro Leu Gly Lys Val Arg Gln Ile Tyr
 Ser Ala Tyr Gln Phe Pro Val Phe Glu Asp Ser Cys Glu Arg Val Ser Gln Gln Ala His Asp Gln Gly
 Gly Pro Ile Gly Ala Ala Arg Cys Glu Gly Gly Trp Leu Cys Gln His Arg Val Pro Phe Val Leu Asn
 Ser Pro Arg Phe Ala Arg Ala Thr Arg Gly Thr Val Val Thr Thr Lys Gly Phe Asp Asp Gly Ala Leu
 Trp Phe Asn Arg Gly Ser Lys Gly Phe Tyr Ala Gln Asn Thr Thr Gly Ser Ser Ile Thr His Thr Phe
 Ser Val Glu Leu Pro Asp Gly Asn Tyr Cys Asp Ile Leu Gly Ala Thr Asp Pro Lys Asn Asn Pro Cys
 Gly Ala Asp Val Thr Val Ser Gly Gly Lys Ala Thr Phe Thr Ile Pro Ala Lys Thr Ala Val Ala Ile
 Cys Thr Asp Glu Lys Trp Cys Gly Lys Gly Val Asp Pro Cys Glu Ser Asp Pro Thr Gly Ser Ala Cys
 Val Cys Lys Gly Glu Thr Thr Val Asn Gly Val Cys Val Ser Trp Cys Asn Ala His Ser Ser Asn Glu
 Glu Cys Ala Cys Val Leu Asn Pro Asn Asp Ala Glu Cys Gln Ala Asp Ile Glu Pro Thr Lys Gly Lys
 Leu Cys Tyr Val Gly Thr Ser Asn Lys Trp Thr Gln Glu Pro Leu Thr Tyr Asn Arg Lys Thr Gly Phe
 Trp Thr Leu Asn Val Glu Leu Asp Gly Lys Gly Asp Thr Ser Gly Ala Gln Arg Phe Lys Val Thr
 Asp Gly Cys Ser Trp Gln Gly Thr Val Tyr Gly Ser Ser Gly Val Glu Gly Arg Leu Asp Val Asn Thr
 Ser Ala Thr Gly Asp Glu Pro Val Ser Leu Thr Gly Lys Tyr Val Leu Ser Ile Asn Asp Lys Thr Met
 Glu Tyr Thr Phe Ile Pro Ala Gly Ser Gly Asn Lys Pro Pro Val Ala Ser Phe Thr Pro Thr Val Lys
 Asp Leu Thr Val Ser Phe Val Asn Asn Ser Ser Asp Pro Glu Asn Asp Glu Leu Thr Tyr Ser Trp Asn
 Phe Gly Asn Gly Lys Thr Ser Ser Glu Lys Asn Pro Ser Val Thr Tyr Asp Lys Ala Gly Lys Tyr Thr
 Val Ser Leu Lys Val Thr Asp Thr Ala Asn Asn Thr Asp Thr Lys Thr Leu Glu Ile Asp Leu Thr Ser
 Pro Val Asn Gly Lys Tyr Ser Lys Val Ala Val Arg Gly Ser His Asp Asn Tyr Gly Thr Asn Leu Leu
 Thr Arg Asn Gly Ser Glu Trp Thr Gly Ile Phe Glu Phe Ser Lys Thr Thr Lys Phe Lys Leu Glu Ala
 Leu Pro Pro Ala Ala Asp Gln Cys Ile Phe Leu Gly Gly Asn Arg Gly Glu Ala Leu Thr Ala Ser Gly
 Gly Phe Ile Ser Leu Pro Ala Gly Arg Tyr Thr Ile Lys Phe Asn Glu Glu Ser Lys Val Leu Thr Ala
 Gly Asp Val Asp Cys Thr Gly

SEQ ID NO: 151

atgaaaactattcttcaacaatcatggtgatggcggtgcggctgccaccaccgtagaggetcaaggctggccggaaaactacggcggcgct
 atgttcagggtattctactgggttcttaccagccaccaagtggaactaaactggaagcacaggctgacgagatctgcaactattctcgtctgta
 tgggtaccacagtcggcctataccggcagcagctacccatgggtacgaccgcgtgtattacttcgaccagcattcatcgttcggcaccgaag
 agcagctacggctcgttcacagctacacagcagaagaaggactggcctacatagcggatgtatgttcacacgaaagaatgtctcaaacg
 gatgatttcctggcgagacacacaggtgtaacctatcagatgtaagcaccgacatggttgaacgagtgaggcgaaaggaagcga
 ctgggcaaatcaaacggctacagctctctcctcaatgccgaaggaaggctgggacggcagcagcctggatcacaagtgca
 gaacgtgcagaaatcggtcttgctacaccaaatatctggttgacgacttaggctataccggattcggctacgatatggtaaagggttgacgg
 atcgcatgtagccgactacaacaccaatgccggcgtgcagttctctgcggcgaatatgggacggcactgcacgaaagttagcgttgatca
 acagcaccacaaagagcgatgtgcgcagtcggcagccttcgacttcgcttccgatacacctgccgcgatgcgctcaacaacaagaactgg
 gcgaacctgaagaacacttcgggtatcagcgatgccgattacaggcgtcttccggttacggttgtaaaatcacgatacggaaatccgttcagct
 accggttcccaggatcccatcaagggtgatacgggtgcctcaatgcctggatgctggtatgccgggcacacctgtgtttctgaaacattgg
 accgactgcaaggagatcaagaatctatcgaggcagctgcctggtcgggtattcacaaccagacacctatgccgaatggatgagcgg
 tgcagcctacatcggtacgtacgtaacaggtacgaacggcacccttactgttctgtgcggctctatcagataatgtagccgccactacattca
 gattctctcaggcaaaaactataatactactgactcaacacgctcgaggtccctggatcgggaaagggttcggctcgtacaccgaagggtgaa
 accgtaacggttcgctcagccatcggccgatgccaatgccagctggtatataccaccgaaggcagacacccaccgaacctcaaca
 gccgtaacagcggaacggaaactgaccatcacttcggacggcgtctgaagggttggtctgttccggcggcagcaggaacatacagagc
 cgtacattcaccttcaggctgcaaacacctccgagtattacagccacctgacgctatgcaaccagtcggagctctcaatccgctgtttgc
 ctatgtttggcaggacggacacgagcagatlaacggcaactggcggcaccacagctcaccgtatccattaccgaaacacaccccttccct

FIGURE 16KKK

ggtaacacgcagtcgttccagattccgaagaacgtggactatgtcgtgaactttgtttaccacaaccggcggttacgcagacagtgatgtt
 accggcatgaaggccgatgtctgttacattattaacagtaccaagagcggaacaagtacacggtaaccgacgttacctcacagtattctcgtt
 agaggccatctttgatgaagaaaactccggctcctcctgtctatgacctgcagggacgcgcgtcagcgaattagaaacaggacaattatat
 ctccagaacggaaagaagatactatcagataaacagagggtccgaaccattctcctattatgaaaatcagacacttagtaatctcagcactgctg
 ggtttggggggctgtacaccatcagctgctcctcgtcggg

SEQ ID NO: 152

Met Lys Thr Ile Leu Ser Thr Ile Met Val Met Ala Ala Ala Ala Thr Thr Val Glu Ala Gln Gly
 Trp Pro Glu Asn Tyr Gly Gly Val Met Leu Gln Gly Phe Tyr Trp Asp Ser Tyr Ser Ala Thr Lys Trp
 Thr Lys Leu Glu Ala Gln Ala Asp Glu Ile Cys Asn Tyr Phe Ser Leu Val Trp Val Pro Gln Ser Ala
 Tyr Thr Gly Ser Ser Thr Ser Met Gly Tyr Asp Pro Leu Tyr Tyr Phe Asp Gln His Ser Ser Phe Gly
 Thr Glu Glu Gln Leu Arg Ser Phe Ile Ser Thr Tyr Lys Gln Lys Gly Thr Gly Ile Ile Ala Asp Val Val
 Val Asn His Arg Lys Asn Val Ser Asn Trp Val Asp Phe Pro Ala Glu Thr Tyr Asn Gly Val Thr Tyr
 Gln Met Val Ser Thr Asp Ile Val Ser Asn Asp Asp Gly Gly Lys Thr Ala Thr Trp Ala Asn Gln Asn
 Gly Tyr Ser Leu Ser Ser Asn Ala Asp Glu Gly Glu Gly Trp Asp Gly Met Arg Asp Leu Asp His
 Lys Ser Gln Asn Val Gln Lys Ser Val Leu Ala Tyr Thr Lys Tyr Leu Val Asp Asp Leu Gly Tyr Thr
 Gly Phe Arg Tyr Asp Met Val Lys Gly Phe Asp Gly Ser His Val Ala Asp Tyr Asn Thr Asn Ala
 Gly Val Gln Phe Ser Val Gly Glu Tyr Trp Asp Gly Thr Ala Ser Lys Val Tyr Ser Trp Ile Asn Ser
 Thr Lys Lys Ser Asp Val Pro Gln Ser Ala Ala Phe Asp Phe Ala Phe Arg Tyr Thr Cys Arg Asp Ala
 Val Asn Asn Lys Asn Trp Ala Asn Leu Lys Asn Thr Ser Gly Ile Ser Asp Ala Asp Tyr Arg Arg Tyr
 Ser Val Thr Phe Val Glu Asn His Asp Thr Glu Tyr Arg Ser Ala Thr Ala Ser Gln Asp Pro Ile Lys
 Gly Asp Thr Val Ala Leu Asn Ala Trp Met Leu Ala Met Pro Gly Thr Pro Cys Val Phe Leu Lys His
 Trp Thr Asp Cys Lys Glu Glu Ile Lys Asn Leu Ile Glu Ala Arg Arg Leu Val Gly Ile His Asn Gln
 Ser Thr Tyr Ala Glu Trp Met Ser Gly Ala Ala Tyr Ile Gly Arg Thr Val Thr Gly Thr Asn Gly Thr
 Leu Arg Val Leu Cys Gly Ser Tyr Gln Tyr Asn Val Ala Ala Asn Tyr Ile Gln Ile Leu Ser Gly Lys
 Asn Tyr Lys Tyr Tyr Val Leu Asn Thr Leu Glu Ala Pro Trp Ile Gly Lys Gly Ser Gly Ser Tyr Thr
 Glu Gly Glu Thr Val Thr Val Pro Leu Ile Ala Ile Ser Ala Asp Ala Asn Ala Lys Leu Val Tyr Thr
 Thr Asp Gly Thr Asp Pro Thr Ala Thr Ser Thr Ala Val Thr Ser Gly Thr Glu Leu Thr Ile Thr Ser
 Asp Ala Val Leu Lys Val Gly Leu Leu Ser Gly Gly Ile Val Arg Asn Ile Gln Ser Arg Thr Phe Thr
 Phe Gln Ala Ala Asn Thr Ser Glu Tyr Tyr Thr Ala Thr Met His Val Cys Asn Gln Ser Gly Ala Leu
 Asn Pro Leu Phe Ala Tyr Val Trp Ala Gly Pro Asp Asn Glu Gln Ile Asn Gly Asn Trp Pro Gly Thr
 Lys Leu Thr Ala Thr Ile Thr Glu Asn Asn Leu Thr Trp Tyr Thr Gln Ser Phe Gln Ile Pro Lys Asn
 Val Asp Tyr Val Val Asn Phe Val Phe Thr Thr Thr Gly Gly Gly Thr Gln Thr Val Asp Val Thr Gly
 Met Lys Ala Asp Val Trp Tyr Ile Ile Asn Ser Thr Lys Ser Gly Asn Lys Tyr Thr Val Thr Asp Val
 Thr Ser Gln Tyr Ser Ser Leu Glu Ala Ile Phe Asp Glu Glu Asn Ser Gly Ser Phe Pro Val Tyr Asp
 Leu Gln Gly Arg Arg Val Ser Glu Ile Arg Asn Arg Thr Ile Ile Ser Ser Glu Arg Lys Glu Asp Thr
 His Gln Ile Asn Arg Gly Ser Glu Pro Phe Ser Tyr Tyr Glu Asn Gln Thr Leu Ser Asn Leu Ser Thr
 Ala Gly Phe Gly Gly Leu Val His His Gln Leu Leu Leu Val Gly

SEQ ID NO: 69

atgttgaaggattacggtagctgtttattgtttattgtttcttctaatatataatgaggaaataaggcagaagcagcaacagtgaaacaatgga
 acattatgcagtatgttggtacgtccgaatgatgggaatcattggaatcgtttgcgttcgatgctgaaagtttagctcataaaggaaatcac
 atctgtatggataccacctgcatataaagggaactcgcataaatgatgtagggtatgggcctatgattatattgattgggagttcaatcaaaaa
 ggaaacggtgcggcagcaaatatgggacaaaagcacagttgaaatcgcattgacgctttacataagcaaacatcgacgtatacgggtatgtatg
 ttatgaatcataaagggtgggctgattatactgaaccgtaacagctgttgaggtagaccgtaacaatcgaataatgaatcaggtgattatca
 aattagtgatggacggggttttaatttccagggcgcggagatgcttattcctaattcaaatggaatgtatcatttgacggaacggattgggatg
 aaggaaaggaaattaaatcgaattataaatttaggggtgtagataaagcgtgggattgggaagtgtctagcgaataatggaattatgattattgat
 gtatgcagatcttgatttgatcatcctgatgttgcgaatgagatgaaaaattggggaacatggtatgcgaatgaattaaatttagtggtttcgttt
 ggacgtctgtaaacatattgatcatgaattttacgcgattgggtaaatcatgccagacagcaaacggggaaagaatgttacagtagctgataa
 ttggcaaaatgatgttcagggttttaaacattatttagcgaaggcatttaataatcaatcgtgtgttgatgcacggcttcattacaatttcattatgcttc

FIGURE 16LLL

aacaggaaatgggaattatgatagagaaatatttaaatggaaacagtaaatgaaaaatcacctgcactcgcagttactctcgttgagaatcatgat
tctcagcctgggcagtcattggaatctgtagtaagtcctggtttaagccgctggcatatgcaittatttaactcgtgcagagggctatccttcagtt
ttctatggtgattactatgggacaagcggaaatagtagttatgaaattccagcgttaaaagataaaattgatccaatttgacggcacgaaaaaactt
tgcatatggtacgcagcgtgattatttagaccatccagatgtgattggctggacaagagaaggcgatgggtacatgctaaitcigtgttagcgac
attactctcgacggaccaggaggatcaaatggtgatggttggaagaataacgctggggaagtatggtacgatattacgggtaatacaaac
aaatactgtaacaattaataaggacggatgggggcagttctatgtaagtggcggtcagttccatatatgttcagcggtaa

SEQ ID NO: 70

Met Leu Lys Arg Ile Thr Val Val Cys Leu Leu Phe Ile Leu Leu Phe Pro Asn Ile Tyr Glu Gly Asn
Lys Ala Glu Ala Ala Thr Val Asn Asn Gly Thr Leu Met Gln Tyr Phe Glu Trp Tyr Ala Pro Asn Asp
Gly Asn His Trp Asn Arg Leu Arg Ser Asp Ala Glu Ser Leu Ala His Lys Gly Ile Thr Ser Val Trp
Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Asn Asp Val Gly Tyr Gly Ala Tyr Asp Leu Tyr Asp Leu
Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Leu Lys Ser Ala Ile
Asp Ala Leu His Lys Gln Asn Ile Asp Val Tyr Gly Asp Val Val Met Asn His Lys Gly Gly Ala Asp
Tyr Thr Glu Thr Val Thr Ala Val Glu Val Asp Arg Asn Asn Arg Asn Ile Glu Val Ser Gly Asp Tyr
Gln Ile Ser Ala Trp Thr Gly Phe Asn Phe Pro Gly Arg Gly Asp Ala Tyr Ser Asn Phe Lys Trp Lys
Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Gly Arg Lys Leu Asn Arg Ile Tyr Lys Phe Arg Gly
Val Asp Lys Ala Trp Asp Trp Glu Val Ser Ser Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp
Leu Asp Phe Asp His Pro Asp Val Ala Asn Glu Met Lys Asn Trp Gly Thr Trp Tyr Ala Asn Glu
Leu Asn Leu Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Asp His Glu Tyr Leu Arg Asp Trp Val
Asn His Ala Arg Gln Gln Thr Gly Lys Glu Met Phe Thr Val Ala Glu Tyr Trp Gln Asn Asp Val Gln
Ala Leu Asn Asn Tyr Leu Ala Lys Val Asn Tyr Asn Gln Ser Val Phe Asp Ala Pro Leu His Tyr Asn
Phe His Tyr Ala Ser Thr Gly Asn Gly Asn Tyr Asp Met Arg Asn Ile Leu Asn Gly Thr Val Met Lys
Asn His Pro Ala Leu Ala Val Thr Leu Val Glu Asn His Asp Ser Gln Pro Gly Gln Ser Leu Glu Ser
Val Val Ser Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg Ala Glu Gly Tyr Pro Ser
Val Phe Tyr Gly Asp Tyr Tyr Gly Thr Ser Gly Asn Ser Ser Tyr Glu Ile Pro Ala Leu Lys Asp Lys
Ile Asp Pro Ile Leu Thr Ala Arg Lys Asn Phe Ala Tyr Gly Thr Gln Arg Asp Tyr Leu Asp His Pro
Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Gly Val His Ala Asn Ser Gly Leu Ala Thr Leu Leu Ser
Asp Gly Pro Gly Gly Ser Lys Trp Met Asp Val Gly Lys Asn Asn Ala Gly Glu Val Trp Tyr Asp Ile
Thr Gly Asn Gln Thr Asn Thr Val Thr Ile Asn Lys Asp Gly Trp Gly Gln Phe Tyr Val Ser Gly Gly
Ser Val Ser Ile Tyr Val Gln Arg

SEQ ID NO: 153

ttgccttaattaatgcaagcgattgcaaaaaaaggagataggagtatgaagaggaaaaatggactgcgttagcactatcttaccactagtt
atgagcttatcaacaacatacaagcagaaacattacataataaagggtcaaaaggcgcaaacaggaaataaagacggaattttatgaact
gtatgtaattcttttatgatactgataagcaatggacatggtgatttaaaaggcgacacaaagaaacttgatttttaaatgatggaaatccaagaac
aaataatgattcttaataaaggatctggatgatgcctattacacctctcctagttatcaaaatagatgtaacgattatataatcgaact
cagttatggaatttcaagatttccgtgaacttacaacagaagcgataaacgcgaacgtaaaagggtgtaatagatctgttataatcaaacagc
agtgaagcattccttggttgcgaigcattaaaaataaaaacagtaagtatcgagattactatatttgggtgataaaaatacagacttaaatgaaaa
aggcccatggggtcaacaagtatggcacaagcgtcgaacggagagtattttacgcaacgttctgggaagggtatgccggacttaactatga
caaccctaaagtaagagaagaaatgattaaaatcgggaattttggctcaaacaggagctgatggcttctctagatgcagccatgcacatctt
taaagggcaaacacctgaaggagcaaaagaaaaatattgaatgggtgaatgaattccgcgacgcgatgagagaacgaatccaaatcgtatct
agttggtagaaatgggataaccagaagtgtgtccgttatcaatcgttagttctacatttaacttcgacttagcatataaaatcgttaattcc
gttaaaaaatggtactgatcaagggttagccgcggcagctgttgcaacggatgagttatataaaacataataccaaataaaattgatggaacgttt
ttaacgaatcatgaccaaaatcgtgtaatgagtgagttaaatgggtgatgaacaagcaaaatcagcagcctctattctgttgacactccctggta
atccgttatttatttggcgaagaaatcggcatgacaggccaaaaccagatgagttgattcgtgagccttccgttggtatgaagtataaag
aaggtaaacgagctgggagactccagatatataacattgatcataatggtgtttcagttgagacacaagataaaacaaaaagcttctcttaagcc
attatcgtaaaaatgattcgttctgcagcaacacgatgaactgtcaaaagtaattagaacctatttctgtcaataattcagaggttggtgcctataat
cgtacgtataaaaaataatcaattcaagtgtaccataatatttcagacaagccggttacattaactgtttcaaacaaaggaaaactgatttttctagt
gaattaggagcaaaaaaggaaaaatcaacattagtaattccagcgaatcagacagtgctagtaaaagtaa

FIGURE
16MMM

SEQ ID NO: 154

Met Pro Ser Ile Asn Ala Ser Asp Cys Lys Lys Lys Gly Asp Arg Ser Met Lys Arg Lys Lys Trp Thr
 Ala Leu Ala Leu Ser Leu Pro Leu Val Met Ser Leu Ser Thr Asn Ile Gln Ala Glu Thr Leu His Asn
 Asn Lys Gly Gln Lys Ala Gln Thr Gly Asn Lys Asp Gly Ile Phe Tyr Glu Leu Tyr Val Asn Ser Phe
 Tyr Asp Thr Asp Ser Asn Gly His Gly Asp Leu Lys Gly Val Thr Lys Lys Leu Asp Tyr Leu Asn
 Asp Gly Asn Pro Arg Thr Asn Asn Asp Leu Gln Ile Asn Gly Ile Trp Met Met Pro Ile Asn Thr Ser
 Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Ser Leu Gln Asp
 Phe Arg Glu Leu Thr Thr Glu Ala His Lys Arg Asn Val Lys Val Val Ile Asp Leu Val Ile Asn His
 Thr Ser Ser Glu His Pro Trp Phe Val Asp Ala Leu Lys Asn Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr
 Ile Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Pro Trp Gly Gln Gln Val Trp His Lys Ala
 Ser Asn Gly Glu Tyr Phe Tyr Ala Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Lys
 Val Arg Glu Glu Met Ile Lys Ile Gly Lys Phe Trp Leu Lys Gln Gly Ala Asp Gly Phe Arg Leu Asp
 Ala Ala Met His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Glu Trp Trp Asn Glu
 Phe Arg Asp Ala Met Arg Glu Thr Asn Pro Asn Thr Tyr Leu Val Gly Glu Ile Trp Asp Gln Pro Glu
 Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Thr Phe Asn Phe Asp Leu Ala Tyr Lys Ile Val Asn
 Ser Val Lys Asn Gly Thr Asp Gln Gly Val Ala Ala Ala Val Ala Thr Asp Glu Leu Tyr Lys Thr
 Tyr Asn Pro Asn Lys Ile Asp Gly Thr Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu
 Asn Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Phe Ile
 Tyr Tyr Gly Glu Glu Ile Gly Met Thr Gly Gln Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp
 Tyr Glu Asp Asp Lys Glu Gly Gln Thr Ser Trp Glu Thr Pro Val Tyr Asn Ile Asp His Asn Gly Val
 Ser Val Glu Ala Gln Asp Lys Gln Lys Ala Ser Leu Leu Ser His Tyr Arg Lys Met Ile Arg Val Arg
 Gln Gln His Asp Glu Leu Val Lys Gly Asn Leu Glu Pro Ile Ser Val Asn Asn Ser Gln Val Val Ala
 Tyr Asn Arg Thr Tyr Lys Asn Lys Ser Ile Gln Val Tyr His Asn Ile Ser Asp Lys Pro Val Thr Leu
 Thr Val Ser Asn Lys Gly Lys Leu Ile Phe Ser Ser Glu Leu Gly Ala Lys Lys Glu Lys Ser Thr Leu
 Val Ile Pro Ala Asn Thr Thr Val Leu Val Lys

SEQ ID NO: 155

gtgtcaagaatgtttgcaaaacgattcaaaaccttactgcccgttattcgtgtgattttattgctgtttcatttggcttggcaggaccaacggcgtg
 cgaatgctgaacaggctaaacaaatcaaatgagcttacgaccgctgatcaaaagggaaccattcttcatgcttggaaatggctgttcaatcacg
 taaaacacaaatgatgaaggatattcatgatgcaggatatacagcgattcagacgctccgattaaaccaagtaagggaacgaaggaaataa
 aaacatgtcgaactggctatggctctatcagccgacatcgtaccaaattggcaaccgttacttaggtactgaacagaattaaagaaatgtgtgc
 agccgctgaagaatatggcataaaggttattgtgaacggtcatcaatcaccaccagtactatgccgcgattccaatgagattaagagtatt
 ccaaactggacacatggaacacacaaaataaaaactggctgatcgtgggatgtcacgcagaatgcattgctcggcgtgtatgactggaata
 cacaaaaacacaaagtacagtcctatttgaacggcttctagaagagcattgaatgacggggcagacgggtttcatttgatgccgcaacata
 tagagcttccggatgatggcagttacggcagtcatttggccgaatatcacaataacatcgcagagttccaatcggagaatcctgcaggat
 agtgcctgaagaatgcttcatgcaatattgattgtacagcgtctaacatagggaattccataaggctcgccttgaagaatcctgaatcggg
 cgtgtgaattatctccactatgcatgattgtgtggaagcgtatgacatgggtagaategcattgataatgctccaatgatgatgag
 agtcgacatggatgagcgtatgatatacgttttaggtggcgggtgatagcttcgttcaggcagtagcctcttttctttcagacctgaggg
 aggcggaaatggtgtgagattcccggggaaagccaaataggcgatcggggagtgcttattgaagatcaggctatcactgcggtaaatg
 attcacaatgtgatggctggacagcctgaggaactctgaacccaatggaacaaccagataattatgaatcagcggcgtcacatggcgttg
 tgctggcaaatgcagggttcatcctctgtttctatcaatcgcacaacaaatggcctgatggcaggatgataataaagctggggcaggttcattca
 agtaaatgacgggtaactgacaggcagcatcaatgccagggtctgtgctgtgttctctgatgataattgcaaaagcgcctcatgtttccttgag
 aattacaaaacagggtgaacacattcttcaatgatcaactgacgattacactgcgtgcagatgcgaatacaacaaaagcgtttatcaaatcaata
 atggaccagagacggcgtttaaggatggagatcaattcacaatcgaaaaggagatccatttggcaaacataccatcatgttaaaagggaac
 gaacagtgatggtgaacgaggaccgaggaatacgtttgttaaaagagatccagcttcggccaaaaccatcggctatcaaaatccgaatcatt
 ggagccaggtaaatgcttatactataacatgatggggccgggca

SEQ ID NO: 156

FIGURE 16NNNN

Val Ser Arg Met Phe Ala Lys Arg Phe Lys Thr Ser Leu Leu Phe Leu Phe Ala Gly Phe Leu Leu Leu
Phe His Leu Val Leu Ala Gly Pro Thr Ala Ala Asn Ala Glu Thr Ala Asn Lys Ser Asn Glu Leu Thr
Ala Pro Ser Ile Lys Ser Gly Thr Ile Leu His Ala Trp Asn Trp Ser Phe Asn Thr Leu Lys His Asn
Met Lys Asp Ile His Asp Ala Gly Tyr Thr Ala Ile Gln Thr Ser Pro Ile Asn Gln Val Lys Glu Gly
Asn Gln Gly Asn Lys Asn Met Ser Asn Trp Tyr Trp Leu Tyr Gln Pro Thr Ser Tyr Gln Ile Gly Asn
Arg Tyr Leu Gly Thr Glu Gln Glu Phe Lys Glu Met Cys Ala Ala Ala Glu Glu Tyr Gly Ile Lys Val
Ile Val Asp Ala Val Ile Asn His Thr Thr Ser Asp Tyr Ala Ala Ile Ser Asn Glu Ile Lys Ser Ile Pro
Asn Trp Thr His Gly Asn Thr Gln Ile Lys Asn Trp Ser Asp Arg Trp Asp Val Thr Gln Asn Ala Leu
Leu Gly Leu Tyr Asp Trp Asn Thr Gln Asn Thr Gln Val Gln Ser Tyr Leu Lys Arg Phe Leu Glu
Arg Ala Leu Asn Asp Gly Ala Asp Gly Phe Arg Phe Asp Ala Ala Lys His Ile Glu Leu Pro Asp Asp
Gly Ser Tyr Gly Ser Gln Phe Trp Pro Asn Ile Thr Asn Thr Ser Ala Glu Phe Gln Tyr Gly Glu Ile
Leu Gln Asp Ser Ala Ser Arg Asp Ala Ser Tyr Ala Asn Tyr Met Asn Val Thr Ala Ser Asn Tyr Gly
His Ser Ile Arg Ser Ala Leu Lys Asn Arg Asn Leu Gly Val Ser Asn Ile Ser His Tyr Ala Ser Asp
Val Ser Ala Asp Lys Leu Val Thr Trp Val Glu Ser His Asp Thr Tyr Ala Asn Asp Asp Glu Glu Ser
Thr Trp Met Ser Asp Asp Asp Ile Arg Leu Gly Trp Ala Val Ile Ala Ser Arg Ser Gly Ser Thr Pro
Leu Phe Phe Ser Arg Pro Glu Gly Gly Gly Asn Gly Val Arg Phe Pro Gly Lys Ser Gln Ile Gly Asp
Arg Gly Ser Ala Leu Phe Glu Asp Gln Ala Ile Thr Ala Val Asn Arg Phe His Asn Val Met Ala Gly
Gln Pro Glu Glu Leu Ser Asn Pro Asn Gly Asn Asn Gln Ile Phe Met Asn Gln Arg Gly Ser His Gly
Val Val Leu Ala Asn Ala Gly Ser Ser Ser Val Ser Ile Asn Thr Pro Thr Lys Leu Pro Asp Gly Arg
Tyr Asp Asn Lys Ala Gly Ala Gly Ser Phe Gln Val Asn Asp Gly Lys Leu Thr Gly Thr Ile Asn Ala
Arg Ser Val Ala Val Leu Tyr Pro Asp Asp Ile Ala Lys Ala Pro His Val Phe Leu Glu Asn Tyr Lys
Thr Gly Val Thr His Ser Phe Asn Asp Gln Leu Thr Ile Thr Leu Arg Ala Asp Ala Asn Thr Thr Lys
Ala Val Tyr Gln Ile Asn Asn Gly Pro Glu Thr Ala Phe Lys Asp Gly Asp Gln Phe Thr Ile Gly Lys
Gly Asp Pro Phe Gly Lys Thr Tyr Thr Ile Met Leu Lys Gly Thr Asn Ser Asp Gly Val Thr Arg Thr
Glu Glu Tyr Ser Phe Val Lys Arg Asp Pro Ala Ser Ala Lys Thr Ile Gly Tyr Gln Asn Pro Asn His
Trp Ser Gln Val Asn Ala Tyr Ile Tyr Lys His Asp Gly Gly Arg Ala

SEQ ID NO: 157

SEQ ID NO: 157

atgcaaacgattgcacacggaaggggatgaaacgatgaaaggggaaaaatggacagcattagctctaacactgccgctggctgctgacttatca
acaggcggttcacgccgaaccgtacataaaggtaaagctccaacagcagataaaaacgggtgcttttatgagggtgtagtaaacctcttttacgat
gcaaaataaagatggacatgggtgatttaaaaggctctacacaaaagctggattattgaaatgacggcgcaattctcataccaaaaatgatcttcaagtaa
acgggaatttggatgatgccggtaaaccccttccttagctatcataaataatgatgtaacggactattataacattgatccgcgtagcggaaatctgca
agattttcgcaagctgatgaaagaagcagataaacgagacgtaaagggtattattggacctgtgtgtaatcatacaagcagtgacaatccttggt
tcaagctgcattaaaaagataaaaacagcaagtacagagattactatatttgggccgataaaaatactgatttaaatgaaaaaggatcttgggggca
gcaagtagtaggcataaaagctccaaacgggagtagtattttatggtacgttttgggaaggaaatgcctgacttaattcagataatcccgaaagtaagaaa
agaaatgattaacgtcpggaaatttggctaaagcaaggcgttgacgggttccgcttagatgctgcgcgttcataattttaaaaggctaaacacctgaa
ggcgctaaagaaaaatagggtggtggaatgatttggatgcgaatgaaaaagataaacctaacgtatatctaacgggggaagtagtgggac
aacgggtgtaggtgctctctacataaagctggggttcttatttaacattgatttagagggaaggaatgtaaacctctgtaaaatcagggaatgatca
aggaatcgcgactgcagcagccgcaactgatgagctgttcaaatcatacaatccaaataaaattgacggcatttcttaaccacacatgaccaa
atcgcgtcatgagtgcgtaagcggcgatgtgaaataaagcaaaagtcagctgcctctatcttacttacgcttccggcaaccgtagatttattacgg
tgaagaaattggaatgaccgggtgaaaagcctgatgagttaatccgtgaaccgttccgctggtacgaaggcaatggacttggacaaaccagctg
ggaaacatccgtatacaacaaaggcgggcaatgggtgtgtagtagagacacaaacaaaaaaggattcttgttaaatacttaccgtgaaatga
ttcgcgtgctgcagcagcatgaagagtgtagtaaaaaggaaaccttcaactattttagtagacagtaaagaagtcgttgcctatagccgcacgtata
aaggcaaatcgattagcgtgtagtataatatttcaaatcaaccggtaaaagtagtctgtaacagcgaaaggtaaattgatttttctagtgaaaaagggt
gcaaaaaaagtcaaaaatcagcttgtgttccagctatacaacgggtttaataaaaaaa

SEQ ID NO: 158

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ala Pro Thr
Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His

FIGURE 16000

Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
 Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
 Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
 Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
 Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
 Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
 Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly
 Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Val Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
 Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser
 Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Asn Ser Val Lys Ser Gly Asn Asp Gln
 Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly
 Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys
 Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
 Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr
 Ser Trp Glu Thr Ser Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Thr Gln Thr Lys Gln Lys
 Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly
 Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser Ile
 Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Thr Ala Lys Gly Lys Leu Ile Phe Ala
 Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Val Pro Ala Asn Thr Thr Val Leu Ile Lys

SEQ ID NO: 159

ttgcaaaaaaaggggatgaacgatgaagggaataatggacagcttagcttaacactgccgctggctgctagcttatcaacaggcgttc
 acgccgaaaccgtacataaaggtaaatctccaacagcagataaaaacgggtatatttatgaggtgatgtaaactcttttacgatgcaataaaga
 tggacatgggtgatttaaaaggcttacacaaaagggtgattatttaaatgatggcaattctcatacaagaatgatcttcaagtaaaccgggattggat
 gatgccgggtcaacccttctccagctatcataaatgatgtaacggactattataattgatccgcagatggaatctgcaagattttcgcaaac
 tgaatgaagaagcagataaacgagatgtaaaagtcattatggacctggtgtgaatcatagcagcagtgaaacaccttgggttcaagctgcattaa
 aagataaaaacgacgaatgacagagattactatctgggctgataaaaataccgactgaatgaaaaaggatctggggacagcaagatggca
 taaagctccaaacggagagatttttacggaacgttttgggaagggaatgccggacttaattacgataatcctgaagtaagaaaagaatgattaa
 cgtaggaaagtttggctaaagcaaggagtgatgggttccgtctagatgctgcgttcataattttaaggccaacacacctgaaggcgctaagaa
 aatctcctgtgtgggaatgaatttagagatgcaatgaaaaaggaaaaccctaactatacttaacgggtgaagtgatggatcaaccggaagta
 gtgctccttactatcaatcgcttgatctttatttaacittgatttagcaggaaagattgtaaacctgttaaatcaggaatgatcaaggaaatcgca
 ctgcagcagcggcgaacggatgaactgttcaaatcataaatcaataaattgacgggtattttctaaccaacctgacccaaatcgctcatga
 gtgagctaaacggcgatgtaataagcaaaagtcagctgcctctacttactacgcttctgccaacccgtatattattacgggtgaagaatcgg
 catgaccgggtgaaaagcctgatgagtaaatcggtgaaccgttccctgttacgaaggaaacggacttggacaaccagctgggaacacacctgt
 atatacaaaaggcgcaacggcggtgtctgtgagcacaacacaaaaggactcttggtaaatcattaccgtgaatgattcgcgtgcgtc
 agcagcaggaagagttgtaaaaggaaacgtttaaactatttaagtagacagtaagaagtcgttgcctatagcgtgaagtaaaaggcaaatg
 attagcgtgataataatttaaatcaatcggtaaaagtagtagcgaaggtaaaatgattttgctagtgaanaagggtcgaagaaagt
 caaaaatcagcttgattcggcgaataaacgggttaataaaataa

SEQ ID NO: 160

Met Gln Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr Leu Pro
 Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr Ala Asp
 Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp
 Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu
 Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp
 Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala Asp
 Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Gln
 Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu
 Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr

FIGURE 16PPP

Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly
 Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln
 Thr Pro Glu Gly Ala Lys Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
 Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser
 Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Asn Ser Val Lys Ser Gly Asn Asp Gln
 Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly
 Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Asn Gly Asp Val Asn Lys Ala Lys
 Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
 Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Pro Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr
 Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys
 Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly
 Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser Ile
 Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala
 Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys

SEQ ID NO: 161

gtggatccaaagaattgtatgcaatttatgcaaacgattgcaaaaaaggggatgaaacgatgaaagggaataatggacagctttagctctaa
 cactgccgctggctgctagcttatcaacaggtgttcacgccgaacccgtacataaaggtaaagctccaacagcagataaaaaagggtgtctttat
 gaggtatattgtaaacctttttacgatgcaataaagatggacatgggtatttaaaggccttacacaaaagttggactatttaaatgacggaaattc
 tcatacaagaatgatcttcaagtaaacgggatttggatgatgccggtcaaccccttcctagctatcataaatagatgtaacggactattataat
 tgatccgcagtgatggaactctgcaagatttgcgaacttatgaagaagcagataaacgagacgtaaaagtcattatggaccttgggtgaatcat
 acgagcagtgaaacaccttggttcaagctgcgttgaaagataaaaacagcaagtacagagattactataatttgggctgataaaaaactgacttg
 aatgaaaaaggatcttggggacaacaagatggcataaagctccaacggagagatattttacggaacgttctgggaaggatgcctgacttaa
 attacgataacctgaagtaagaaaagaatgattaacgtcggaaagtttggctaaaacaaggcgttgacggcttccgcttagatgctgcccttc
 atattttaaaggtaaacgcctgaaggcgctgaagaaaacattctatgggtgaatgagtttagatgcgatgaaaaagaaaacccgaacgta
 tatctaacgggtgaagtggtggaccagcagaagtagtagcccttactatcaatcacttgattctctatttaatttggatttagcaggaaaaattgc
 agctctgtaaaagcaggaaatgatcaaggaatcgccactgcagcagcgccaaactgatgagctgttcaaatcataaatccaaataaattgacg
 gcatttttaaccaacctgacaaaaatcgctcatgagtgagtgtaagcggcgatgtgaataaagcaaaatcagccgctctacttactacgct
 tcttgaaatccgtatatttatcgggtgaagaaattggcatgacaggtgaaaagcctgatgaattaatccgtgaaccgttccgctggtacgaagg
 caacggaattggcacaactagctgggaaacacctgtatatacaaaaggcggtaacggcgtgtctgtagaagcacaacaaaaaaaaggatt
 ccttgttaaatcattaccgtgaatgattcgtgtgcgcagcagcacgaaggttagtaaaaggacgcttcaatccatttcagtagacagtaaaag
 aagtcgttgctatagccgcagctacaaggcaaatcgattagcgtgtatcataatatttcaaatcaacctgtaaaagtatctgtagcagcgaaag
 gtaacttgattttgctagtgaaaaagggtgtaagaaagtcataaatcagcttgtgattccggcgaaatgcgacgggtttataaaaataa

SEQ ID NO: 162

Val Asp Pro Lys Asn Cys Ser Gln Phe Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly
 Lys Lys Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu
 Thr Val His Lys Gly Lys Ala Pro Thr Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser
 Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu
 Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro
 Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln
 Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val
 Asn His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp
 Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His
 Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp
 Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe
 Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Leu Trp
 Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp
 Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly
 Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp Gln Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu

FIGURE 16QQQ

Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val
 Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly
 Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu
 Pro Phe Arg Trp Tyr Glu Gly Asn Gly Ile Gly Gln Thr Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly
 Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met
 Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser Ile Ser Val Asp Ser Lys
 Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro
 Val Lys Val Ser Val Ala Ala Lys Gly Asn Leu Ile Phe Ala Ser Glu Lys Gly Ala Lys Lys Val Lys
 Asn Gln Leu Val Ile Pro Ala Asn Ala Thr Val Leu Ile Lys

SEQ ID NO: 163

atggtacgtcccgaacgacgggctgcattggaaccgactatcgaacgactcgcagcactgaaagacattgggtgacgacgggtggattccg
 ccggcgtaacaaaggcacgtcacagaacgatgtcgggtatggggcgtagatttatc gatctcggcgaattcaacaaaaaggacgacccg
 gacgaagtacgggacgaaagcgacgtccagaccgccatctcgaacttcgcggtaaaaggatcgggtgtacggcgacgtcgtatgaat
 cacaaggcgggggccgattataccgaatccgttcaggcgatcaggtcaatccgtcgaaccggaaccaagaacgtccggtagtatggcat
 ctggcctggactgggttcaacttcggggcgcaacaatacatctcgcgttcaaatggcgctgtaccattttgacgggtaccgattgggac
 agtcacgcagcttgagccgcatctataagttcaagagcacaggcgaaggcgtgggacacggacgtgtcgaacgagaacggcaactatgattat
 ctatgtatgccgacgtcgatttcgagcatcccgaggctcgccaagagatgaagaactggggcaaatggtagccgactcgtcgggctcgac
 gggttcgggttgatcggtcaaatacagccactcgtactgaaggagtggtgacgagcgtgcgccagacgacgggaaagagatgttc
 acggtcgcccagatttgaagaacgatctcgggtccatcaacgactatctgtataagacgggctacacgcactcgtctcgtatgtcgcgtcc
 attataactccaagcggccggtaacggcgccgggtattacgatatgcgaacatcttgaaggcaccgtcaccgaacagcatccgtcgtcgtc
 cgtgacgattgtcgaaccacgactcacagccggcgccagtcgctgagtcgacggtcgccaactgggtcacaaccgctcgcctacgcgacga
 tcatgacgcgcgggtcagggtatccggccctctctatggagactattatggcacgaaaggacgacgaaccgcgaatccggaacatgtcgg
 gcacgctccaaccgattttgaaggcacgaaaagacttcgctacgggacgcagcatgactacctcgtatcatcaggacgtcatcggctggacac
 gtgaagggtgtgacggaccgtgccaatcgggtctcgcgacgattctatcgacgggtcggcggtcgaagtggatgtacgtcggcaaacag
 aacgcggcgaggtatggaagacatgacgaacaacaacgccctcgtcgtacgatcaatgctgacgggtgggtcagttcttcgtaacgg
 aggtcgtgctcgtattatagcaacaataa

SEQ ID NO: 164

Met Val Arg Pro Glu Arg Arg Ala Ala Leu Glu Pro Thr Ile Glu Arg Leu Ala Ala Leu Glu Arg His
 Trp Val Thr Thr Val Trp Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Asn Asp Val Gly Tyr Gly Ala
 Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Thr Arg Thr Lys Tyr Gly Thr Lys
 Ala Gln Leu Gln Thr Ala Ile Ser Asn Leu Arg Gly Lys Gly Ile Gly Val Tyr Gly Asp Val Val Met
 Asn His Lys Gly Gly Ala Asp Tyr Thr Glu Ser Val Gln Ala Ile Glu Val Asn Pro Ser Asn Arg Asn
 Gln Glu Thr Ser Gly Glu Tyr Gly Ile Ser Ala Trp Thr Gly Phe Asn Phe Ala Gly Arg Asn Asn Thr
 Tyr Ser Pro Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Thr Asp Trp Asp Gln Ser Arg Ser Leu Ser
 Arg Ile Tyr Lys Phe Lys Ser Thr Gly Lys Ala Trp Asp Thr Asp Val Ser Asn Glu Asn Gly Asn Tyr
 Asp Tyr Leu Met Tyr Ala Asp Val Asp Phe Glu His Pro Glu Val Arg Gln Glu Met Lys Asn Trp
 Gly Lys Trp Tyr Ala Asp Ser Leu Gly Leu Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Ser His
 Ser Tyr Leu Lys Glu Trp Val Thr Ser Val Arg Gln Thr Thr Gly Lys Glu Met Phe Thr Val Ala Glu
 Tyr Trp Lys Asn Asp Leu Gly Ala Ile Asn Asp Tyr Leu Tyr Lys Thr Gly Tyr Thr His Ser Val Phe
 Asp Val Pro Leu His Tyr Asn Phe Gln Ala Ala Gly Asn Gly Gly Gly Tyr Tyr Asp Met Arg Asn Ile
 Leu Lys Gly Thr Val Thr Glu Gln His Pro Ser Leu Ser Val Thr Ile Val Asp Asn His Asp Ser Gln
 Pro Gly Gln Ser Leu Glu Ser Thr Val Ala Asn Trp Phe Lys Pro Leu Ala Tyr Ala Thr Ile Met Thr
 Arg Gly Gln Gly Tyr Pro Ala Leu Phe Tyr Gly Asp Tyr Tyr Gly Thr Lys Gly Thr Thr Asn Arg Glu
 Ile Pro Asn Met Ser Gly Thr Leu Gln Pro Ile Leu Lys Ala Arg Lys Asp Phe Ala Tyr Gly Thr Gln
 His Asp Tyr Leu Asp His Gln Asp Val Ile Gly Trp Thr Arg Glu Gly Val Thr Asp Arg Ala Lys Ser
 Gly Leu Ala Thr Ile Leu Ser Asp Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Gln Asn Ala
 Gly Glu Val Trp Lys Asp Met Thr Asn Asn Ala Arg Leu Val Thr Ile Asn Ala Asp Gly Trp Gly
 Gln Phe Phe Val Asn Gly Gly Ser Val Ser Ile Tyr Thr Gln Gln

FIGURE 16RRR

SEQ ID NO: 165

atgcagtatttcgagtggtacgtgccaaatgatggggaacattggaatcgtttgcgtaatgatgctgaaaatttagctcataaaggaaattacatctgt
 atggataccacccgtatataaaggaaacttcacaaaatgatgtagggtatggagtgtatgatgtatatgattgggagaattcaatcaaaaaggaaac
 gatacggacaaaatattgggacaaaagcacaattaaaatctgcaattgaggctttacataatcaaaaatcgaatgatacgggtgatgttgatgaac
 cataaagggtgggagcagattatactgaggtgtaacagccgttgaggtagaccgtaacaatcgaaatattgaacatcgaagtattcaaatagat
 gcgtggacgggattgattttccaggacgcagggacicttaictaattttaaatggagatgggttcattttgatggaacagattgggatgaggga
 ggaaattaaatagaatttataaatttaaaggcgtaggtaaagctgggactgggaagtgtctagtgaatggtaactgattatttaattgtatgca
 gatcttgatttcgatcatcttgaagttgcaaaatgaaatgaaaaactggggaacctgggtatcgggacgaattaaatttagatggcttctgttagacg
 cagttaaacattattgacatgagatcttctgattgggtaaatcatgttagaagcaaacggggaaggaaatgtttacagtagctgaatttgcca
 aaatgatattctgtactttaacaattatttagggaaagtaaatataatcaatctgtgttcgatgcaccttctcattataatttcattatgcttcaacagg
 gaatggaaattatgataggaataatttaaagggtacggtagtagaaagtcctcactctgctgttactctgttgagaatcatgattctcagcc
 tggacagtcattagaatctgtgtgagtcctgtttaaagccgttgccatgcaattttaacgcgtgcagaagggtatctctgtttttatggag
 attactatggcacaatggaaatagtagttatgaaattccaacgttaaaggataaaattgatccaattctgacggcacgaaaaactttgcatatgg
 tacgcaacatgattatttagaccatccagatgtgattggctggacaagagaaggggatagatacatgctaattctgttttagcaacattatctctg
 atggaccaggaggatcaaaatggatgaatgttggaaagaacaacgcaggggaatatgtgtacgatattacgggcaatcaacaataactgtaa
 cgattataaagatggatggggcgatgccatgtaaatggggctctgtttcaatatatgttcagaagtaa

SEQ ID NO: 166

Met Gln Tyr Phe Glu Trp Tyr Val Pro Asn Asp Gly Glu His Trp Asn Arg Leu Arg Asn Asp Ala
 Glu Asn Leu Ala His Lys Gly Ile Thr Ser Val Trp Ile Pro Pro Val Tyr Lys Gly Thr Ser Gln Asn
 Asp Val Gly Tyr Gly Val Tyr Asp Val Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Ile Arg Thr
 Lys Tyr Gly Thr Lys Ala Gln Leu Lys Ser Ala Ile Glu Ala Leu His Asn Gln Asn Ile Asp Val Tyr
 Gly Asp Val Val Met Asn His Lys Gly Gly Ala Asp Tyr Thr Glu Val Val Thr Ala Val Glu Val Asp
 Arg Asn Asn Arg Asn Ile Glu Thr Ser Ser Asp Tyr Gln Ile Asp Ala Trp Thr Gly Phe Asp Phe Pro
 Gly Arg Arg Asp Ser Tyr Ser Asn Phe Lys Trp Arg Trp Phe His Phe Asp Gly Thr Asp Trp Asp Glu
 Gly Arg Lys Leu Asn Arg Ile Tyr Lys Phe Lys Gly Val Gly Lys Ala Trp Asp Trp Glu Val Ser Ser
 Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Phe Asp His Pro Glu Val Ala Asn
 Glu Met Lys Asn Trp Gly Thr Trp Tyr Ala Asp Glu Leu Asn Leu Asp Gly Phe Arg Leu Asp Ala
 Val Lys His Ile Asp His Glu Tyr Leu Arg Asp Trp Val Asn His Val Arg Lys Gln Thr Gly Lys Glu
 Met Phe Thr Val Ala Glu Tyr Trp Gln Asn Asp Ile Arg Thr Leu Asn Asn Tyr Leu Gly Lys Val Asn
 Tyr Asn Gln Ser Val Phe Asp Ala Pro Leu His Tyr Asn Phe His Tyr Ala Ser Thr Gly Asn Gly Asn
 Tyr Asp Met Arg Asn Ile Leu Lys Gly Thr Val Val Glu Ser His Pro Thr Leu Ala Val Thr Leu Val
 Glu Asn His Asp Ser Gln Pro Gly Gln Ser Leu Glu Ser Val Val Ser Pro Trp Phe Lys Pro Leu Ala
 Tyr Ala Phe Ile Leu Thr Arg Ala Glu Gly Tyr Pro Ser Val Phe Tyr Gly Asp Tyr Tyr Gly Thr Asn
 Gly Asn Ser Ser Tyr Glu Ile Pro Thr Leu Lys Asp Lys Ile Asp Pro Ile Leu Thr Ala Arg Lys Asn
 Phe Ala Tyr Gly Thr Gln His Asp Tyr Leu Asp His Pro Asp Val Ile Gly Trp Thr Arg Glu Gly Asp
 Ser Ile His Ala Asn Ser Gly Leu Ala Thr Leu Ile Ser Asp Gly Pro Gly Gly Ser Lys Trp Met Asn
 Val Gly Lys Asn Asn Ala Gly Glu Ile Trp Tyr Asp Ile Thr Gly Asn Gln Thr Asn Thr Val Thr Ile
 Asn Lys Asp Gly Trp Gly Gln Phe His Val Asn Gly Gly Ser Val Ser Ile Tyr Val Gln Lys

SEQ ID NO: 167

atgcacacgattgcaaaaaaggggatgaaacgatgaaagggaataatggacagcttttagcttaacactgccgctggctgctagcttatca
 acaggcgttcacgcccgaaccgtacataaaggtaaatctccaacagcagataaaaacgggtattttatgagggtatgtaaacctctttttagatg
 caataaagatggacatgggtatttaaaggcttctacacaaaagtggattatttaaatgatggcaattctcatacaaaagatgacttcaagtaaac
 gggatttggatgatgccgtcaaccctctccagctatcataaattatgatgtaacggactattataattatgatccgagatggaatctgcaag
 attttcgcaactgatgaaagaagcagataaacgagatgtaaaagtcattatggacctgtgtgtaacatacagcagtgaaacacctgtgttc
 aagctgcattaaagataaaaacagcaagtacagagattactatctgggctgataaaaataccgacttgaatgaaaaaggatcttggggaca
 gcaagtatggcataaagccccaacggagagatattttacggaacgttttgggaaggaaatgccggacttaattacgataatcctgaagtaagaa
 aagaaatgattaacgtaggaaagtgttggctaagcaaggaggtgacgggttcgcttagatgctgcgcttcatattttaaaggccaacacctg

FIGURE 16SSS

aaggcgctaagaaaaatctctgtgtggaatgaattagagatgcaatgaaaaggaaaaccctaactatatactaacgggtgaagtatggga
 tcaaccggaagtagtagctccttactatcaatcgcttgattcttatttaactttgatttagcaggaaagattgtaaaactctgtaaaatcaggaaatgat
 caaggaaatcgcgactgcagcagcggcaacggatgaacigtcaaatcatacaatccaaataaaattgacgggtatttcttaaccaaccatgacca
 aaatcgcgctcatgagtgagctaagcggcgatgtgaataaagcaagtcagctgcctctatcttactacgcttcctggcaaccctatattttac
 ggtgaagaaatcgcatgaccgggtgaaaagcctgaigagttatccgtgaaccgttccgctgtacgaaggaaacggacttgacaaaccag
 ctgggaaacacctgtatacaacaaaggcggcaacggcggtgtctgtagaagcacaacaaaacaaaaggactcttgttaaatcattaccgtgaa
 atgattcgctgctgcagcagcacgaagagttagtaaaaggaacgcttcaatctatttcagtagacagtaagaagtcgttgcctatagccgcac
 gtataaaggcaaatcgattagcgtgtatcataatatttcaaatcaaccggtaaaagtatctgtacgagcaaaaggtaaatgattttggtagtga
 aagggtgctaagaagtcaaaaatcagcttgtgattccggcgaatacaacggttttaataaaataa

SEQ ID NO: 168

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
 Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr
 Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His
 Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
 Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
 Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
 Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
 Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
 Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
 Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly
 Gln Thr Pro Glu Gly Ala Lys Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys
 Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln
 Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Asn Ser Val Lys Ser Gly Asn Asp
 Gln Gly Ile Ala Thr Ala Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp
 Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala
 Lys Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met
 Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln
 Thr Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln
 Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys
 Gly Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys
 Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile
 Phe Gly Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu
 Ile Lys

SEQ ID NO: 169

atgaaaacattcaatataaaagcaactttttacogctaaccttctgtctcagtgctcttgcctttgttgggcaaaaaggcaaccatgatgcagtattt
 cattggtatgtacctaatgatggcgattatggacgcagggttgaagcaatgctccagcactcgtgaaaaagggtttacagcgctctggctacc
 gccagcttacaaggcgcggcgagtaatacgcgcggttatggcgtctatgatgtacgatttaggtgagttgatcaaaaaggctcagtac
 gaaccaaataggcaccgaaggtcagtagctcttgaatcaatgccgcgcaacaacaatatccaaatctacggcgaigtgtgttaaacac
 cgaggtgtgtgatgggaagtcgtgggtcgaaccaagcgcgttgattgggacaaccgtaacattgaactgggcgacaaatggattgaagct
 tgggttgagtttaatttctggccgaacgacaaaatacicaaacttccattggacttggtatcactttgacgggtgtgactgggatgacggcga
 agaaaaagcgatcttaaatcaaaaggcgaaggaaaagcatggattgggaagtcagctctgaaaaaggcaattacgactacctaatgtacgc
 cgatttagacatggatcaccaagaagttaaacaagagctgaagattgggtgagtggtacatcaacatgaccggcggttagtgcttagaatg
 gatccgtgaagcacattaaatatcagtagctacaagagtggtgatcatttacgttgaaaacaggcaagagcgtttaccggttggtgagatt
 ggaattacgacgtaaatcaactgcataactttattactaagacctctggcagtagtgcgtgttcgatgcgcccttcacatgaacttcacaacgcg
 tcaaaatctggcggcaattacgatatgcgcaaatcatgaatggcacgttgatgaaggacaacccagtcgaagctgtgactctcgtgaaaacc
 acgatacacagccattgcagcggttagatgcacagtggtggttgaagcctcttgccttacgcattcattttatgctgtaagaagggtatcc
 atcagtggtctacgcagattactacggcgcagtagcgcgacaaaggctacaaccatcaatattggcgaaggcttaccattgaagacnlgtaa

FIGURE 16TTT

cactgcgtaaagagtagtcgtatggcaaacagaattcttatctcgaccactgggatgtgattggctggaccgcagaggggcgaigctgaacatcc
aaactcaatggcggtagcatgagtgatggaccagggtggcaaaaatggatgtataccggtaagccaagcacgcgctatgtcgacaagctgg
gtatccgaactgaagaagtttggaccgataccaatggctgggcagaaattcctgtcaatgggtgttcagctcggtttgggtgggcgttaagtaa

SEO ID NO: 170

Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu Leu Ser Ala Pro Ala Phe Ala Gly Gln Asn Gly Thr Met Met Gln Tyr Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu Ser Asn Ala Pro Ala Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ser Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile Ser Ala Ile Asn Ala Ala His Asn Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His Arg Gly Gly Ala Asp Gly Lys Ser Trp Val Asp Thr Lys Arg Val Asp Trp Asp Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp Val Glu Phe Asn Phe Pro Gly Arg Asn Asp Lys Tyr Ser Asn Phe His Trp Thr Trp Tyr His Phe Asp Gly Val Asp Trp Asp Asp Ala Gly Lys Glu Lys Ala Ile Phe Lys Phe Lys Gly Glu Gly Lys Ala Trp Asp Trp Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp His Gln Glu Val Lys Gln Glu Leu Lys Asp Trp Gly Glu Trp Tyr Ile Asn Met Thr Gly Val Asp Gly Phe Arg Met Asp Ala Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp His Leu Arg Trp Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn Tyr Asp Val Asn Gln Leu His Asn Phe Ile Thr Lys Thr Ser Gly Ser Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala Ser Lys Ser Gly Gly Asn Tyr Asp Met Arg Gln Ile Met Asn Gly Thr Leu Met Lys Asp Asn Pro Val Lys Ala Val Thr Leu Val Glu Asn His Asp Thr Gln Pro Leu Gln Ala Leu Glu Ser Thr Val Asp Trp Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser Val Phe Tyr Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly Tyr Asn Ile Asn Met Ala Lys Val Pro Tyr Ile Glu Glu Leu Val Thr Leu Arg Lys Glu Tyr Ala Tyr Gly Lys Gln Asn Ser Tyr Leu Asp His Trp Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser Met Ala Val Ile Met Ser Asp Gly Pro Gly Gly Lys Lys Trp Met Tyr Thr Gly Lys Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr Glu Glu Val Trp Thr Asp Thr Asn Gly Trp Ala Glu Phe Pro Val Asn Gly Gly Ser Val Ser Val Trp Val Gly Val Lys

SEO ID NO: 171

gtgtatgtaaacctcttttacgatgcaaaataaagatggacatgggtgattlaaaaggcttacacaaaagtggattttaaatgatggcaattcctacataaaagaatgatcttcaagtaaacgggatttggatgatgccggtaacccctctccagctatcataaatatgatgtaacggactattataatattgatccgcagataggaaatctgcaagattttcgc aaactgaigaagaagcagataaacgagatgtaaaagtcattatggaccctgtgttgatcatagcagcagtgaaacccctggittcaagctgcattaaaagataaaaacagcagatcacagagattactatctgggctgtataaaaataccgactgcatgaaaaaggatcttggggacagcaagataggcataaaagcccaaacggagagatttttacggaaacgttttgggaaggatgccggacttaaaatcagataalectgaagtaagaaaagaatgattaacgtaggaaagtttggctaaaggcaaggagtggacgggtccgctagatgctgcgttcatatttttaaaaggcaaaacccctgaaggcgctagaaaaatatctgttggggaaatgaatttagagatgcaatgaaaaggaaaacccctaaagctatataaagggtgaagatgggatgaacggagtagtagctcttaataaattgctgattcttaataactttgatttagcaggaaagattgtaaactctgtaaaatcaggaaatgatcaagggaatcgcgactgcagcagcggcaacggatgaactgttcaaatcatacaatccaaataaaaattgacggtattttcttaaccaaccatgacccaaaatcgcgctatgagtgaactaagcggcgatgtgaataaaggcaaaagtcagctgcctctactttacttacgcttccgtggcaaccggtatattttattacggtgaa gaaatcggcatgaccggtgaaaaagcctgatgagtaatccgtgaaccgttccgctgtgacgaaggaaacggacttggacaaaacagcttgggaaacacctgtatacaaaaaggcggaacggcggtgtctgtagaagcacaacaaaaacaaaaggactctttgtaaatcattaccgtgaaatgattcgcgtgcgtcagcagcacgaaaggttagtaaaaggaaacgcttc aatctatttcagtagacagtaaaagagtcgttgccctatagccgcacgtataaaaggcaaatcgattagcgtgtatcataalatttcaaatcaaccggtaaaagtactctgtagcagcaaaaaggtaaattgatttttgtagtgaaaaagggtgctaagaaagtcaaaaatcagcttgtattccggcgaatacaacggtttaataaaaaataa

SEO ID NO: 172

Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln
Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp
Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro

FIGURE 16UUU

Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val
 Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys
 Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp
 Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly Met Pro
 Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys
 Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala
 Lys Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu Asn Pro Asn Val Tyr
 Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe
 Asn Phe Asp Leu Ala Gly Lys Ile Val Asn Ser Val Lys Ser Gly Asn Asp Gln Gly Ile Ala Thr Ala
 Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn
 His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile
 Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro
 Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr Ser Trp Glu Thr
 Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys Asp Ser Leu Leu
 Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser
 - Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser Ile Ser Val Tyr His
 Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Gly Ser Glu Lys
 Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys

SEQ ID NO: 173

atgcaaacgattgcaaaaaaggggatgaacgatgaagggaataatggacagcttagctctaacactgccgctggtgctagcttatca
 acaggcggttcacgcagaaactgtacataaaggtaaaagctccaacagcagataaaaaacgggtgtttttatgagggtgatgtaaactcttttacgatg
 caaataaagatggacatggtgatttaaaaggctgcacaaaagtggattatataatgacggcaattctcatcaaaagaatgatcttcaagtaaa
 cgggatttggatgatgccggtaaaccttctctagctatcataaatatgatgaacggactattataacattgatcctcagtagcggaaagtctgcaa
 gatttcgcaaaactgatgaaagaagcagataaacgagacgtaaaagtattatggacctgtgtgaatcatacagcagtgaaacaccttggtt
 caagctgcactaaaagataaaaacagcaagtagagatactataatttggctgataaaaaatccgattgaatgaaaaggatcttggggaca
 gcaagtattggcataaagctccaaacggagagatattttacggaacgtctggaaggaatgcctgactaaattacgataaacctgaagtaagaa
 aagaaatgattaacgtcggaaagtttggctaaagcaaggcgttgatggcttccgcttagatgctgcccttcatactttaaggctcaaacctctga
 aggcgctaagaaaaatctctgtgtggaatgagtttagagatgcaatgaaaaagaaaaccctaactatatacgggtgaagtatgggat
 cagccggaagttagctccttattatcaatcgttgattccctatttaactttagcaggaaaaattgtcagctctgtaaaagcaggaaatgat
 caaggaaatgccactgcagcagcggcaacggatgagctgttcaatcataatccaaataaaattgacggcatttcttaaccaacctgacca
 aaaccgcgcatgagtgaagcggagatgtgaataaagcaaaatcagctgcttctacttactacgcttctggaatccgtatattttacg
 gtgaagaaattggcatgaccgggtgaaaagccgatgaattaatccgtgaaccgttccgctgtgacgaaggcaacggaattggacaaactagct
 gggaaacacctgtatatacaaaaggcggcaatggtgtgtctagaaagcacaacaaacaaaggattcttgttaaatcattaccgtgaaatg
 attcgcgtgcgcagcagcagaagagtagtaaaaggaacgcttcagctctattcagtagacagtaaaagaggtgtcgttatagccgtacgtat
 aaaggcaactecattagtgtgtatcataatattcaaatcaacctgtaaaagtatctgtagggcgcaaggtaaatgattttgttagtaaaaagg
 tgetraaaaaggcaaaaatcagcttggattcgggaatgagacgggttaataaataa

SEQ ID NO: 174

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
 Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ala Pro Thr
 Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His
 Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
 Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
 Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Ser Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
 Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
 Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
 Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
 Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly

FIGURE 16VVV

Gln Thr Pro Glu Gly Ala Lys Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys
 Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln
 Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp
 Gln Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp
 Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala
 Lys Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met
 Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Ile Gly Gln
 Thr Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln
 Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys
 Gly Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Asn
 Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile
 Phe Ala Ser Glu Lys Gly Ala Lys Lys Gly Lys Asn Gln Leu Val Ile Pro Ala Asn Ala Thr Val Leu
 Ile Lys

SEQ ID NO: 175

atgaaaaataatacgaactttgtgtgccagcgctatcctcacggtgtccacgccagttacgccagcaattttacacgcgtttaactggcaat
 ataccgatgaaccgcaatgcaaatcaaatgccgcaatggcttaaaaaagtcctcattcaccgcaatgaaatccagcggcagtcgaatgg
 tgggcccgtatcaaccgcaagacttgcgtgtcattgattctccgctgggcaacaacagatttagtcgcgatgatcaatgcgctcaacagcgt
 tggggtgcagcgtgtatgctgacgtgtgtcctaaccatattgctaacgagtcagtggaagcgcagtgacctgaactaccggggagtgagggtc
 caacgactatcaatcccgcagtgcttactatcaaaaggcaaacacttttcggcaattacaggagaacctttttccgagaatgatttccatccggca
 ggctgtattaccaattggaatgatcctggccacgtccagttatggcgctgtgcggcgacagggcgatactgggctaccggatctcgatcctaa
 tcaatgggttgtagtcagcagaagagttactgaacgcactcaaatcaatgggaatcaaaagggttcgatacgcggtcaaacatatgagtc
 aatatcaaatagaccaagtgttaccacagacattaccgctggtatgcataattcggagaagtcattaccagtggtggcgaaggatagcggct
 atgaggttttcttgccttaccctaataatccgatcacgccgttatgacttccgctatttgcacgacgcgcgttttcaatctctgtgg
 gttaaatcagctacacaatccacaagcctatggccaagcgttacaggactcacgtgcgacacattacgattaccacgacattccaaccaatg
 acgggttccgctaccagatcatggatccaaccgatgaacagctcgcctatgcctacatttgggcaagatggaggaacgccactgtctatagt
 gatgacctacctgacagcgaagacaaagacagtggtcgttggccgatgtgtggaagatccgaacatgattaacatgcttgccttccacaacg
 cgatcaaggacaaagcatgactgtatggttagcgcacatgtaccttgcattaaagcgcggcaagcaaggcgtgtaggaatcaataaatg
 tggcgagagtaagtcggtgactgtcgatacttaccagcatgagtttaactgtacacccgtaccaagacgtattgagcggcgacatcaccaca
 gtgagttctcgttatccaatgttttgcagcgcgcagtgcaaggatgtggaactataa

SEQ ID NO: 176

Met Lys Asn Ile Ile Arg Leu Cys Ala Ala Ser Ala Ile Leu Thr Val Ser His Ala Ser Tyr Ala Asp Ala
 Ile Leu His Ala Phe Asn Trp Gln Tyr Thr Asp Val Thr Ala Asn Ala Asn Gln Ile Ala Ala Asn Gly
 Phe Lys Lys Val Leu Ile Ser Pro Ala Met Lys Ser Ser Gly Ser Gln Trp Trp Ala Arg Tyr Gln Pro
 Gln Asp Leu Arg Val Ile Asp Ser Pro Leu Gly Asn Lys Gln Asp Leu Val Ala Met Ile Asn Ala Leu
 Asn Ser Val Gly Val Asp Val Tyr Ala Asp Val Val Leu Asn His Met Ala Asn Gln Ser Trp Lys Arg
 Ser Asp Leu Asn Tyr Pro Gly Ser Glu Val Leu Asn Asp Tyr Gln Ser Arg Ser Ala Tyr Tyr Gln Arg
 Gln Thr Leu Phe Gly Asn Leu Gln Glu Asn Leu Phe Ser Glu Asn Asp Phe His Pro Ala Gly Cys Ile
 Thr Asn Trp Asn Asp Pro Gly His Val Gln Tyr Trp Arg Leu Cys Gly Gly Gln Gly Asp Thr Gly
 Leu Pro Asp Leu Asp Pro Asn Gln Trp Val Val Ser Gln Gln Lys Ser Tyr Leu Asn Ala Leu Lys Ser
 Met Gly Ile Lys Gly Phe Arg Ile Asp Ala Val Lys His Met Ser Gln Tyr Gln Ile Asp Gln Val Phe
 Thr Pro Asp Ile Thr Ala Gly Met His Ile Phe Gly Glu Val Ile Thr Ser Gly Gly Gln Gly Asp Ser Gly
 Tyr Glu Ala Phe Leu Ala Pro Tyr Leu Asn Asn Thr Asp His Ala Ala Tyr Asp Phe Pro Leu Phe Ala
 Ser Ile Arg Ala Ala Phe Ser Phe Ser Gly Gly Leu Asn Gln Leu His Asn Pro Gln Ala Tyr Gly Gln
 Ala Leu Gln Asp Ser Arg Ala Ile Thr Phe Thr Ile Thr His Asp Ile Pro Thr Asn Asp Gly Phe Arg
 Tyr Gln Ile Met Asp Pro Thr Asp Glu Gln Leu Ala Tyr Ala Tyr Ile Leu Gly Lys Asp Gly Gly Thr
 Pro Leu Val Tyr Ser Asp Asp Leu Pro Asp Ser Glu Asp Lys Asp Ser Gly Arg Trp Ala Asp Val Trp
 Gln Asp Pro Asn Met Ile Asn Met Leu Ala Phe His Asn Ala Met Gln Gly Gln Ser Met Thr Val Val
 Ala Ser Asp Gln Cys Thr Leu Leu Phe Lys Arg Gly Lys Gln Gly Val Val Gly Ile Asn Lys Cys Gly

2013201807

FIGURE
16WWW

Glu Ser Lys Ser Val Thr Val Asp Thr Tyr Gln His Glu Phe Asn Trp Tyr Thr Pro Tyr Gln Asp Val
Leu Ser Gly Asp Ile Thr Thr Val Ser Ser Arg Tyr His Gln Phe Val Leu Pro Ala Arg Ser Ala Arg
Met Trp Lys Leu

SEQ ID NO: 177

atgaaacattcaaattaaacgcactttttaccgctgacctgtgcgtcagtgctctgaccttgctgggcgaaaatggcaccatgatgcagtaattt
cattggtagctgacctaatgatggcgacattatggacgcaggttgaaagcaatgctccagtactcgtgaaaacggttttacagcgctctggctacc
gcccgcatacaaaaggcgccggcgccagtaaatgacgtcggttatggcgctctatgatatgacgatttaggtgagttgacaaaaaggctcagta
cgaaccaataatcgccaccaaggctcagtagcatctctgcaatcaatgccggcgacacaacaacaatatccaaattacggcgacgttggtttaacca
ccgaggtggcgctgatgggaagtcgtgggtcgataccaagcgcggttgattgggacaaccgcaatatgtaactggcgacaaatggatggaag
ctgggttgagtttaatttcttggccgcaacgacaaaatacgaacttccattggacttggtatcactttgacggtgttgactgggatgatgccggc
aaagaaaaagcgatcttaaatcaaaaggcggaaggaagaaagcatgggatgggaagtcagctctgaaaaaggcaattacgactacctaattgac
gccgatttagacatggatcacccagaaggttaaacaagagctgaagattggggtagtggtacatcaaatgaccggcggttgatggctttaaga
tggatgccgtgaagcacattaaatatcagtatctacaagagtggttgatgactttacggttggaacaaggcgaagagctttaccggttggtgagta
ttggaattacgacgtaaatcaactgcacaactttattactaagaccttggcagtagtgcgtgtgtcgaatgcgccgttcaatgaatttcacaacgc
gtcaaaatctggcggcaccttacgatatgcgccaaatcatgaatggcacgttgatgaaggacaaccacgacaaagcagtgactctctgtagaaac
cacgatacgacgccattgcaggcggttagatgcacagtagatgggtgttaagcccttgccttacgcaattcatttattgcgtgaagaagggtatc
catcggtgttctacgcagattactacggcgccagtagacgcgacaaaggttaacaacattaatggccaagtgccttacattgaagaactgtta
cacitgcgtaaaagagtagtgcgtatggcaacagaattctatctcgaccattgggaatgtgattggctggaccggagaggcgatgcigaacatcc
aaactcaatggcggtagatcatgagtgaaggaccggcgccgacaaaatggatgtataccggtaagccaagtacgcgctatgtcgacaagctgg
gtatccgaaactgaagatgtttggaccgatgccaatggctgggcagaatttctgtcaatgggtgttcagctcgtgttgggtggcggttaagtaa

SEO ID NO: 178

SEQ ID NO: 178

Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu Leu Ser Ala Pro Ala Phe Ala Gly Gln Asn Gly Thr Met Met Gln Tyr Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu Ser Asn Ala Pro Val Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ser Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile Ser Ala Ile Asn Ala Ala His Asn Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His Arg Gly Gly Ala Asp Gly Lys Ser Trp Val Asp Thr Lys Arg Val Asp Trp Asp Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp Val Glu Phe Asn Phe Pro Gly Arg Asn Asp Lys Tyr Ser Asn Phe His Trp Thr Trp Tyr His Phe Asp Gly Val Asp Trp Asp Asp Ala Gly Lys Glu Lys Ala Ile Phe Lys Phe Lys Gly Glu Gly Lys Ala Trp Asp Trp Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp His Pro Glu Val Lys Gln Glu Leu Lys Asp Trp Gly Glu Trp Tyr Ile Asn Met Thr Gly Val Asp Gly Phe Arg Met Asp Ala Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp His Leu Arg Trp Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn Tyr Asp Val Asn Gln Leu His Asn Phe Ile Thr Lys Thr Ser Gly Ser Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala Ser Lys Ser Gly Gly Thr Tyr Asp Met Arg Gln Ile Met Asn Gly Thr Leu Met Lys Asp Asn Pro Val Lys Ala Val Thr Leu Val Glu Asn His Asp Thr Gln Pro Leu Gln Ala Leu Glu Ser Thr Val Asp Trp Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser Val Phe Tyr Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly Tyr Asn Ile Asn Met Ala Lys Val Pro Tyr Ile Glu Glu Leu Val Thr Leu Arg Lys Glu Tyr Ala Tyr Gly Lys Gln Asn Ser Tyr Leu Asp His Trp Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser Met Ala Val Ile Met Ser Asp Gly Pro Gly Gly Thr Lys Trp Met Tyr Thr Gly Lys Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr Glu Asp Val Trp Thr Asp Ala Asn Gly Trp Ala Glu Phe Pro Val Asn Gly Gly Ser Val Ser Val Trp Val Gly Val Lys

SEQ ID NO: 179

SEQ ID NO: 179
atgaaaacattcaaaftaaacgcactttttaccgctaacttgcctcagtgctcctgcccttgccgggcaaaatggcaccatgatgcagtactt
tcattggtacgtaccctaagatggcgcatatggacgcagggttgaaagcaaatgcttcagcactcgcctgaaaacggcttttacagcgctctggcacc

FIGURE 16XXX

gccagcttacaaggcgcgggcgccagtaatgatgtcgggtatggcgtctacgatattgattaggtgagttgatcaaaaggctcagtac
 gaaccaataacggtaccaaggctcagtacatcttgcataatgtcgcgcacaacaacaatatccaaatttacggcgacgttggttaaccatc
 gtggtggcgctgatgggaagtcgtgggtcgataccaagcgcgtgattgggacaaccgtaacattgaactggcgacaaatggatgaagcgt
 ggggtgagtttaatttccclagccgcaacgacaaatactcgaacttccattggacttggtatcacittgacgggtgtgactgggatgatccggcaa
 agaaaaagcgatctttaaattcaaggcggaaggaaaagcatgggattgggaagtcagctctgaaaaggcaattacgactacctaattgtacgc
 cgatttagacatggatcaccagaagttaaacaagagctgaagattgggtgagtggtacatcaacatgaccggcggtgatggcttagaatg
 gatgccgtaagcacattaatatcagtatctacaagagtgattgatcattacgttggaacaggcaagagccttaccggttggtgagattg
 gaattacgacgtaaatcaactgcataacttattactaagacctctggcagatgtcgttgcgatgcgccgttcacatgaacttctacaacgcgt
 caaaactggcggaattacgatatgcgcaaatcatgaatggcaggtgatgaaggacaaccagtcgaagctgtgactctctagaaaacca
 cgatacgagccattgcagcggttagagtcgacagtggttggttgcaagccttctgctacgcattcatctgttgcgtgaagaaggttatcca
 tgggtgtctacgcagattactcggcgcgagtagcagcgacaaaggttacaacattaatatggcaaaagtgcttaccattgaagaactgttaaca
 ctgcgtaaagagtagtgcgtatggcaaacagaattcttctcgaacctgggagtgattggctggactcagagggcgatgtgaacatccaaa
 ctcaatggcggtgatcatgagtgatggaccggcggaacaaatggatgtataccggtaatccaagcacgcgclatgtcgacaagctgggtat
 ccgaactgaagatgtttggaccgatgccaatggctggcgagaatttctgcaatgggtgtcagtcggttgggtggcggttaagtaa

SEQ ID NO: 180

Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu Leu Ser Ala Pro Ala Phe Ala
 Gly Gln Asn Gly Thr Met Met Gln Tyr Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln
 Val Glu Ser Asn Ala Pro Ala Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys
 Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp
 Gln Lys Gly Ser Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile Ser Ala Ile Asn Ala Ala His
 Asn Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His Arg Gly Gly Ala Asp Gly Lys Ser Trp
 Val Asp Thr Lys Arg Val Asp Trp Asp Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp
 Val Glu Phe Asn Phe Pro Ser Arg Asn Asp Lys Tyr Ser Asn Phe His Trp Thr Trp Tyr His Phe Asp
 Gly Val Asp Trp Asp Ala Gly Lys Glu Lys Ala Ile Phe Lys Phe Lys Gly Glu Gly Lys Ala Trp
 Asp Trp Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met Asp
 His Pro Glu Val Lys Gln Glu Leu Lys Asp Trp Gly Glu Trp Tyr Ile Asn Met Thr Gly Val Asp Gly
 Phe Arg Met Asp Ala Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp His Leu Arg Trp
 Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn Tyr Asp Val Asn Gln Leu His Asn
 Phe Ile Thr Lys Thr Ser Gly Ser Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala
 Ser Lys Ser Gly Gly Asn Tyr Asp Met Arg Gln Ile Met Asn Gly Thr Leu Met Lys Asp Asn Pro Val
 Lys Ala Val Thr Leu Val Glu Asn His Asp Thr Gln Pro Leu Gln Ala Leu Glu Ser Thr Val Asp Trp
 Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser Val Phe Tyr Ala
 Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly Tyr Asn Ile Asn Met Ala Lys Val Pro Tyr Ile Glu
 Glu Leu Val Thr Leu Arg Lys Glu Tyr Ala Tyr Gly Lys Gln Asn Ser Tyr Leu Asp His Trp Asp Val
 Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser Met Ala Val Ile Met Ser Asp Gly Pro
 Gly Gly Thr Lys Trp Met Tyr Thr Gly Asn Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr
 Glu Asp Val Trp Thr Asp Ala Asn Gly Trp Ala Glu Phe Pro Val Asn Gly Gly Ser Val Ser Val Trp
 Val Gly Val Lys

SEQ ID NO: 181

tggccagaggccttcggcctggccattacgccgtcacatagccggcgggggaggttggtggcggtgcgcggcgggcgagcctgccgatgc
 cggtctccactggcgcggttcctcgtccggcggttcgtcgccggtcatccgaacagcacaagaccggagttgcgatgagccaca
 ccttgcgtgcggcgtattggcgggcgatcctgctgcggttccccctcgtgaccaggccggcaagagcccgccggcggtgcgtacca
 cgcgcgcgacgaatcatctccagggttccactggaacgtctccgcgaagcgcccaacgactggtacaacatcttcgccagcaggcct
 cgacgatcgccggcgagggcttcggcaatctggatgccggtgccctggcgtgacttctcagctggaccgacggcggaagtgcaggcg
 cgcggaaggctactcttggcacgacttcaacaagaacggcgctacggcagcgacgccagctgcgccaggccggcgccgactcggtg
 cgccgggtgaaggtgctctacgatgtggtgcccatacatgaaccggcgctatccggacaaggagatcaacctgcggcgccggaaggc
 ttctggcgcaacgactgcaccgacccgggcaactacccaacgactgcgatgacggtgaccgcttcatcgcgcggaagtcggacctgaaca
 ccggccatccgcagatctcggcatgttcgcgacgaggttgccaactcgcgagcggttacggcgccggcggttccgcttcgacttcgttc

FIGURE 16YYY

gcggctatgcgccgaacgggtcgacagctggatgagcgacagcgccgacagcagtttctgcgttgccgagctgtgaaaagcccgtccga
gtaccgagctgggactggcgcaacacggcgagctggcagcagatcatcaaggactggccgaccgggccaagtccccgtgttcgacttc
gcgctcaaggagcgcatgcagaacggctcggtcgccgactggaagcatggcctcaatggcaaccggaccgcgctggcgaggtggc
ggtgacctttgtcgacaaccacgacaccggctattcgccgggcagaacggcgccagcaccactggcgctgcaggacgggtgatccg
ccaggcctacgcctacatcctcaccagcccgggcacgcccgtgtgtactggcgacatgtacgactggggtacggcgacttcattcgcca
gctgatccagggtgcggcgaccgctggcggtgcgcccgttcggcgatcagcttcacagcggtacagcgccgtgctgctaccgtcagc
ggcagccatcagaccctgggtggcgctcaactccgatctggccaaccggccaggtcgccagcgcgagcttcagcgaggcggtcaac
gccagcaacggccaggtgcgctgttcggcgagcggttagcgcgatggcgccgcaatgacggcgccgaggcggtctgtgtaatgtgaa
cttcgctgcgacaacggcgtagcgagatggcgacagcgctacggtgggcaacgtcagccagctcggaactggagcccgccctc
cgcggtacgggtgaccgacaccagcagctatccgacctggaaggcgagcatcgccctgcctgacggtcagaacgtggaatggaagtgcctg
atccgtaacgaggcggaagcgacgctgtgtgcgcaatggcgccgaacaaccagggtccaggccgctgcccggcgagcacca
gcggctcgttctga

SEQ ID NO: 182

Met Pro Glu Ala Phe Gly Leu Ala Ile Thr Pro Ser His Ser Arg Arg Gly Arg Leu Val Gly Val Ser
Arg Gly Gly Ser Leu Pro Met Pro Val Leu His Trp Pro Ala Phe Ile Leu Val Arg Arg Phe Val Ala
Gly His Pro Asn Lys His Lys Asn Arg Ser Ile Ala Met Ser His Thr Leu Arg Ala Ala Val Leu Ala
Ala Ile Leu Leu Pro Phe Pro Ala Leu Ala Asp Gln Ala Gly Lys Ser Pro Ala Gly Val Arg Tyr His
Gly Gly Asp Glu Ile Ile Leu Gln Gly Phe His Trp Asn Val Val Arg Glu Ala Pro Asn Asp Trp Tyr
Asn Ile Leu Arg Gln Gln Ala Ser Thr Ile Ala Ala Asp Gly Phe Ser Ala Ile Trp Met Pro Val Pro Trp
Arg Asp Phe Ser Ser Trp Thr Asp Gly Gly Lys Ser Gly Gly Gly Glu Gly Tyr Phe Trp His Asp Phe
Asn Lys Asn Gly Arg Tyr Gly Ser Asp Ala Gln Leu Arg Gln Ala Ala Gly Ala Leu Gly Gly Ala
Gly Val Lys Val Leu Tyr Asp Val Val Pro Asn His Met Asn Arg Gly Tyr Pro Asp Lys Glu Ile Asn
Leu Pro Ala Gly Gln Gly Phe Trp Arg Asn Asp Cys Thr Asp Pro Gly Asn Tyr Pro Asn Asp Cys
Asp Asp Gly Asp Arg Phe Ile Gly Gly Lys Ser Asp Leu Asn Thr Gly His Pro Gln Ile Tyr Gly Met
Phe Arg Asp Glu Leu Ala Asn Leu Arg Ser Gly Tyr Gly Ala Gly Gly Phe Arg Phe Asp Phe Val
Arg Gly Tyr Ala Pro Glu Arg Val Asp Ser Trp Met Ser Asp Ser Ala Asp Ser Ser Phe Cys Val Gly
Glu Leu Trp Lys Ser Pro Ser Glu Tyr Pro Ser Trp Asp Trp Arg Asn Thr Ala Ser Trp Gln Gln Ile Ile
Lys Asp Trp Ser Asp Arg Ala Lys Cys Pro Val Phe Asp Phe Ala Leu Lys Glu Arg Met Gln Asn
Gly Ser Val Ala Asp Trp Lys His Gly Leu Asn Gly Asn Pro Asp Pro Arg Trp Arg Glu Val Ala Val
Thr Phe Val Asp Asn His Asp Thr Gly Tyr Ser Pro Gly Gln Asn Gly Gly Gln His His Trp Ala Leu
Gln Asp Gly Leu Ile Arg Gln Ala Tyr Ala Tyr Ile Leu Thr Ser Pro Gly Thr Pro Val Val Tyr Trp Ser
His Met Tyr Asp Trp Gly Tyr Gly Asp Phe Ile Arg Gln Leu Ile Gln Val Arg Arg Thr Ala Gly Val
Arg Ala Asp Ser Ala Ile Ser Phe His Ser Gly Tyr Ser Gly Leu Val Ala Thr Val Ser Gly Ser His Gln
Thr Leu Val Val Ala Leu Asn Ser Asp Leu Ala Asn Pro Gly Gln Val Ala Ser Gly Ser Phe Ser Glu
Ala Val Asn Ala Ser Asn Gly Gln Val Arg Val Trp Arg Ser Gly Ser Gly Asp Gly Gly Gly Asn Asp
Gly Gly Gln Gly Gly Leu Val Asn Val Asn Phe Arg Cys Asp Asn Gly Val Thr Gln Met Gly Asp
Ser Val Tyr Ala Val Gly Asn Val Ser Gln Leu Gly Asn Trp Ser Pro Ala Ser Ala Val Arg Leu Thr
Asp Thr Ser Ser Tyr Pro Thr Trp Lys Gly Ser Ile Ala Leu Pro Asp Gly Gln Asn Val Glu Trp Lys
Cys Leu Ile Arg Asn Glu Ala Asp Ala Thr Leu Val Arg Gln Trp Gln Ser Gly Gly Asn Asn Gln Val
Gln Ala Ala Ala Gly Ala Ser Thr Ser Gly Ser Phe

SEQ ID NO: 183

atgcaaacgattgcaaaaaagggatgaaacgatgaaagggaataatggacagctttagctctaactgccgctgctgctagcttatca
acaggcggttcacgccgaaaccgtacataaaggtaagctgaagcaacagataaaacgggtgcttttatgaggtgtatgtaaccttttacgata
caataaagatggacatggtgatttaaaaggctgcacacaaaagtggattattaaatgacggcaatttcatacaagaatgatcttcaagtaaa
cgggatttgatgatgccagtaaccctctcctagctatcataaatatgatgtaacggactattataacattgatcctcagtaggaaatctgcaag
attttcgcaagctgatgaaagaagcagacaaacgagacgtaaaagtcattatggaccttgtgtgaatcatagcagcagcaaacaccttgggtt
caagctgcattaaaaagataaaaacagcaagtagagattactatatttgggctgataaaaataccgatttgaatgaaaaaggatcttgggggca
gcaagtatggcataaagctccaaacgggagagtatatttccggaacgttttgggaaggaaatgcctgacttaaatatcagataacccctgaagtaagaa

FIGURE 16ZZZ

aagaaatgattaacgtcggaaagtgttggttaagcaaggcgtaagtgttcgcttagatgctgcgttcataattttaaggtaaacacctga
 aggcgctaagaaaaatactgttggtggaatgagtttagatgcgatgaaaaagaaaaccctaactatatacagggtgaagtatgggat
 cagcctgaagtggtagctccttactatcaatcgcttgatctttatattttgatttagcaggaaaaattgtcagctctgtaaaagcaggaaatgatc
 aaggaatgccactgcagcagcggcaacagatgaactgttcaaatcatacaatccaaataaaatgacggcattttcttaaccaacctgaccaa
 aatcgctcatgagtgagctgagcggcgatgtgaacaaagcaaaatcagctgcttcttacttactacgttcttgcaacccgtatattattacg
 gtgaagaaattggcatgaccgggtgaaaagccigatgagttatccgtgaaccattccgctggtagcgaaggaaacggactggacaaactagct
 gggaaacacctgtatatacaaaagggcggaacggcggtgtctgtagaagtacaaacaaacaaaaggattctttgtaaatcattatcgtaaatg
 attcgcgtgcgtcagcagcatgaagagttagtaaaaggaaacgcttaacatatttcagtagacagtaaaagagtggtgacctatagtcgcacgtat
 aaaggcaactcgattagcgtgtatcataatatttcaaatcaacctgtaaaagtagtctgtagcagcgaaaggtaaatgattttgctagtgaaaaagg
 tgcataaaaagtaaaaatcagcttgtaattccggctaatacaacgggttttaataaaataa

SEQ ID NO: 184

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
 Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Glu Ala
 Thr Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Thr Asn Lys Asp Gly His
 Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
 Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
 Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
 Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
 Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
 Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
 Gly Lys Phe Trp Leu Lys Gln Gly Val Asn Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly
 Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
 Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser
 Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp Gln
 Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly
 Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys
 Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
 Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr
 Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Val Gln Thr Lys Gln Lys
 Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly
 Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Asn Ser Ile
 Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala
 Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys

SEQ ID NO: 185

atgaanetgatgaagggaataatggacagcttagcttatacaactgccgctggtgctagcttataaacaggogttacgcgcgaacigtac
 ataaaggtaaaagctccaacagcagataaaaacgggtgtctttatgagggtgatgtaaaactcttttacgatgcaataaagatggcatggtgattta
 aaaggctttacacaaaagctggactattaaatgacggaaatttcatacaaaagaatgatcttcaagtaaacgggatttgatgatgcagcgaac
 ccttctcctagctatcataaatatgatgaacggattattataacattgatccgcagctacggaatctgcaagattttgcaagctgatgaagaagc
 agacaaacgagacgtaaaagtattatggacctgtgtgaatcatcagcagcgaacaccttggtttcaagctgcgttaaaagataaaaaca
 gcaagtacagagattactatatttggcgctgataaaaataccgacttgaatgaaaaggatcttggggacagcaagatggcataaagctccaac
 ggagagatttttacggacggtttgggaaggatgcctgacttaattacgataacctgaagtaagaaagaaatgattaacgtcggaaagtgtt
 ggctaaagcaaggcggttgatggcttcgcttagatgctgcgttcataattttaaaaggtaaacgcctgaaggcgctgaagaaaaatttctgtgt
 ggaaatgagtttagatgcgatgaaaaagaaaaccctaactatatacagggtgaagtatgggatcagcctgaagtggtagctccttactat
 caatcgcttgattccctatttaactttgatttagcagggaattgtcagttctgtaaaagcaggaaatgatcaaggaaatcgccactgcagcagcgg
 caacggatgagctgttcaaatcatacaatccaaataaaatgacggcattttcttaaccaacctgacccaaacggcgtcatgagtgaactgatcg
 gcgatgtgaacaaagcaaaatcagctgcttcttacttactacgttcttggaacccgtatattttatcgggtgaagaaattggcatgaccgggtga
 aaagccctgatgagttatccgtgaacggttcgctggtacgaaggaaacggccttgacaaacccagctgggaaacccctgtatatacaaaagg

FIGURE
16AAAA

cggaacggcgtgtctgtagaagcacaacccaacaaaaggattctttgttaaatcattaccgtgaaatgattcgcgtgcgtcagcagcatgaag
agttagtaaaaggaacgcttcaatctatttttagtagacagtaaaagaagttgtgcctatagccgtacgtataagacaactcgattagcgtgtatcat
aatatttcaaatcaaccggtaaaagtatctgtagcagcaaaaggtaaatattttgtctagtgaaggggtctaaaaagtcagaatcagcttg
tgattccggctaatacaacgggttttaataaaataa

SEQ ID NO: 186

Met Lys Leu Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser
Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ala Pro Thr Ala Asp Lys Asn Gly Val Phe Tyr
Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr
Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp
Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro
Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val
Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys
Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp
Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly Met Pro
Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys
Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala
Lys Lys Asn Ile Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu Asn Pro Asn Val Tyr Leu
Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn
Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp Gln Gly Ile Ala Thr Ala Ala
Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His
Asp Gln Asn Arg Val Met Ser Glu Leu Ile Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile Leu
Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro Asp
Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr Ser Trp Glu Thr Pro
Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys Asp Ser Leu Leu Asn
His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser Ile
Leu Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Asp Asn Ser Ile Ser Val Tyr His
Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala Ser Glu Lys
Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys

SEQ ID NO: 187

ttgtatctatccaggaggggcacatgcgtttccgccattattaccgccgttaccggcctggccgttcgggttgagctctgcgtaccgcacag
agctggcgcataggggagtttgcgacttgcgggttcttgcgaattctgcaaaaaagccggatttgatctgtacagcttcttcgggtcaatgac
accggcacagaaggttccatcacagcgcgtttctgccttgccttgcacccgctgtatcagcgtttccgacctgctgaagcagcgggttcc
gaaaagcagattacagatctgaaaagccgggttgaggacttgcctcgtttcagctatcggagctgcgccgtgccaaactggaatctctgcgtgc
agtgttgataaaaacaaggcaaccatcatcggcagtgccgaactggaagccctggatttcagataacccctggatcatcgaatatcggttttat
gaacagaaagacacggcacttgaagccgggttgaacatttggaaaaggttgcgaacccctactcacaagaatacaaaaaacctggag
ggtaaaaccttggcaggtgacatctatcttgcgttgcgtgagatgcggctggacagcagtttactgagcgtgacaggtgcaagccc
ctgggtgtctatcttaaggcgatatacctataatgatgaacgaggattccgcagatgccctggcgaaatccggaattcttccgtgacgatcttgg
gccggaggtccccctgacgggtgaaaacccccagggaacaaactggggcttccccatttataactgggaaaaccttgcaaatgacgggtacag
ctggtgaaaaaacgtctgaagcacagcgcacggtattaccatgcctaccgcatgaccatattcttgggttttccggatatgggtataccctat
ggcgaatactccggctacctgggatggcccttgcgcgatgaaccggtaagtgcagcagaactggcagaacggggcttttccaaggaccgctt
ggcgttggttaccgaacccccacttgcctacacgggcagccgaggaagcgaataactgggactatctgggaacacacggctatctgaatcaga
tcatgaaccgtatcgggtgaagaagaactatggctgttcaagcccagatcacctgcgaggcagatatacgaacacaaacctgccgatgcc
tgaaagagggttctgtacggcagtggaacacccggctgtgcaggttaccggccgcgacgaaaaggacggacaatctactatccgctgtgg
cgtttccgtgacagcactgcatggcagacgcttaccgatggcgagaacactccctggagagctgttcgccccaaaaagcggcgacacatga
aaccttgtggcgagaacaggcgggtggaacttctgggtgagctgacgcgatctacggatatgcttgcctgtgtgagatctgggaagtattccc
cacagtgtaccggaagtgccttcaaacctttcaattacagcttgcgggttaccgcgtggcccgccaatgggatgccccggcagcccttca
cagactggaggagtatccgctcatgtcggtagcgaacccatcgggtcatgattcttacccttgcgggatggtgggaaacccgaaggcggcgga
ccgggcccattatggacgcatggcctccgggaacaggatgcatacgcaggagcaggccgcctatgagttcgaaggcgttggggacccccgcc

Figure 16 (cont.)

SEQ ID NO: 188

SEQ ID NO: 189

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FIGURE
16CCCC

gaaacacctgtatacaataaaggcggcaacggcgtgtctgtagaagcacaaccaacaaaaggactctttgttaaatcattaccgtgaaatgat
tcggtggtcagcagcagcgaagagtttagtaaaaggaacgcttcaatctatttcagtagacagtaaaagaagttgttctatagccgtacgtataa
aggcaactccattagtgtgtatcataatatttcaaatcaacctgtataaaagtatctgtagcagcgaaaggtaaatgattttgctagtgtaaaaagggtg
ctaaaaagggtcaaaaatcagcttgtgttccggcgaatacaacgggttttagtaaaataa

SEQ ID NO: 190

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Ala
Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His
Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Ile Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
Thr Asp Tyr Tyr Asn Ile Asp Ser Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly
Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Val Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser
Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp Gln
Gly Ile Ala Thr Ala Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly
Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys
Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr
Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys
Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly
Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Asn Ser Ile
Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala
Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Val Lys

SEQ ID NO: 191

atgcaaacgattgcacaaaaagggatgaaacgatgaaagggaataatggacagctttagcttaacactgccgctggtgctagcttatca
acaggcgttcacgccgaaacgtacataaaggtaaatcccaacagcagataaaaacgggtgtctttatgaagtgtatgtaaacctctttttagatg
caataaagatggacatggtgacttaaaaggctttacacaaaagttggactatttaaatgacggcaattctacatacaaaaatgatcttcaagtaaa
cgggatttggatgatgccagtaacccctctcctagctatcataaatatgatgtaacggactattataacatigatccgcagtagcgaaatctgcaa
gatttcgcaagctgatgaagaagcagacaaaacgagacgtaaaagtcattatggacettgttggatcatacagcagtgaaaccccttggtt
caagctgcgttaaaagataaaaaaggaagtaaacggaactatatttggcgtatataaacacgacttgatgaanaaggaattttaggtaac
acaagtatggcataaagctccaaacggagagatttttagggacgttttggaaaggaaatgacttaaatgaacacccctgaagttaagaa
aagaaatgattaacgtcggaaagtgttgctaaagcaaggcgttagcgggttcgcctagatgctgctgctcatattttaaagggtcaaacagctga
aggcgttaagaaaaatacctgtgtggaatgagtttagatgcgatgaaaaagaaatccgaatgtatactaacgggtgaagtatgggat
cagcctgaagtgtgctccttatcaatcgtgtattcttatttaattttagattagcaggaaaaattgtcagctctgtaaaagcaggaaatgac
aaggaatcgccactgcagcagcagcaacagatgaactgttcaaatcatacaatccaaacaaaattgatggcatattcttaaccaaccatgaccaa
aatcgctcatgagtgcgtgagcggcgatgtgagcaaaagcaaatcagctgttctatcttactttagcgttcttggcaaccgtatatttattacg
gtgaagaaatcgccatgaccggtgaaaagcctgatgaataatccgtgaaccgttccgctgtgacgaaggaaacggacttggaacaaacagtt
gggaaacacctgtatacaataaaggcggaaacgggtgtgtctgtagaagcacaacaaacaaaaggattctttgttaaatcattaccgtgaaatg
attcgctgcgtcagcagcatgaagagtttagtaaaaggaacgcttcaatctatttcagtagacagtaaaagaagttgttctatagccgtacgtata
aaggcaactccattagtgtatcataatatttcaaatcaaccggtataaaagtatctgtagcagcgaaaggtaaatgattttgctagtgtaaaaagggt
gctaagaaagtcacaaatcagcttgtgttccggcgaatacaacgggttttagtaataa

SEQ ID NO: 192

FIGURE
16DDDD

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr
Ala Asp Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His
Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn
Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val
Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu Ala
Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly
Thr Phe Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val
Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly
Gln Thr Ala Glu Gly Ala Lys Lys Asn Ile Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser
Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp Gln
Gly Ile Ala Thr Ala Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp Gly
Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Ser Lys Ala Lys
Ser Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr
Ser Trp Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys
Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly
Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Asn Ser Ile
Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala
Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln Leu Val Val Pro Ala Asn Thr Thr Val Leu Met Lys

SEQ ID NO: 193

[illegible]

SEQ ID NO: 194

Met Lys Phe Lys Lys Ser Leu Ser Ala Gly Leu Leu Leu Phe Gly Gly Leu Ser Gly Val Thr Pro Ser
Val Ala Ala Glu Val Pro Arg Thr Ala Phe Val His Leu Phe Glu Trp Ser Trp Pro Asp Ile Ala Thr

FIGURE 16EEEE

Glu Cys Glu Thr Phe Leu Gly Pro Lys Gly Phe Ser Ala Val Gln Val Ser Pro Pro Gln Lys Ser Val
 Ser Asn Ala Ala Trp Trp Ala Arg Tyr Gln Pro Val Ser Tyr Ser Phe Glu Gly Arg Ser Gly Thr Arg
 Ala Gln Phe Ala Asp Met Val Gln Arg Cys Lys Ala Val Gly Val Asp Ile Tyr Leu Asp Ala Val Ile
 Asn His Met Ala Ala Gln Asp Arg Tyr Phe Pro Glu Val Pro Tyr Ser Ser Asn Asp Phe His Ser Cys
 Thr Gly Asp Ile Asp Tyr Ser Asn Arg Trp Ser Ile Gln Asn Cys Asp Leu Val Gly Leu Asn Asp Leu
 Lys Thr Glu Ser Glu Tyr Val Arg Gln Lys Ile Ala Asp Tyr Met Asn Asp Ala Leu Ser Leu Gly Val
 Ala Gly Phe Arg Ile Asp Ala Ala Lys His Ile Pro Ala Gly Asp Ile Ala Ala Ile Lys Ser Lys Leu Asn
 Gly Ser Pro Tyr Ile Tyr Gln Glu Val Ile Gly Ala Ala Gly Glu Pro Val Gln Thr Ser Glu Tyr Thr Tyr
 Ile Gly Asp Val Thr Glu Phe Asn Phe Ala Arg Thr Ile Gly Pro Lys Phe Lys Gln Gly Asn Ile Lys
 Asp Leu Gln Gly Ile Gly Ser Trp Ser Gly Trp Leu Ser Ser Asp Asp Ala Val Thr Phe Val Thr Asn
 His Asp Glu Glu Arg His Asn Pro Gly Gln Val Leu Ser His Gln Asp Phe Gly Asn Leu Tyr Phe Leu
 Gly Asn Val Phe Thr Leu Ala Tyr Pro Tyr Gly Tyr Pro Lys Val Met Ser Gly Tyr Tyr Phe Ser Asn
 Phe Asp Ala Gly Pro Pro Ser Thr Gly Val His Ser Gly Asn Ala Cys Gly Phe Asp Gly Gly Asp Trp
 Val Cys Glu His Lys Trp Arg Gly Val Ala Asn Met Val Ala Phe Arg Asn His Thr Ala Ala Gln Trp
 Gln Val Thr Asp Trp Trp Asp Asp Gly Tyr Asn Gln Val Ala Phe Gly Arg Gly Gly Leu Gly Phe
 Val Val Ile Asn Arg Asp Asp Asn Lys Gly Ile Asn Gln Ser Phe Gln Thr Gly Met Pro Ala Gly Glu
 Tyr Cys Asp Ile Ile Ala Gly Asp Phe Asp Thr Gln Ser Gly His Cys Ser Ala Thr Thr Ile Thr Val
 Asp Ser Gln Gly Tyr Ala His Phe Thr Val Gly Ser His Gln Ala Ala Ala Ile His Ile Gly Ala Lys Leu
 Gly Ser Val Cys Gln Asp Cys Gly Gly Thr Ala Ala Glu Thr Lys Val Cys Phe Asp Asn Ala Gln
 Asn Phe Ser Gln Pro Tyr Leu His Tyr Trp Asn Val Asn Ala Asp Gln Ala Val Ala Asn Ala Thr Trp
 Pro Gly Val Ala Met Thr Ala Glu Asn Gly Gly Tyr Cys Tyr Asp Phe Gly Val Gly Leu Asn Ser Leu
 Gln Val Ile Phe Ser Asp Asn Gly Ala Ser Gln Thr Ala Asp Leu Thr Ala Ser Ser Pro Thr Leu Cys
 Tyr Gln Asn Gly Thr Trp Arg Asp Ser Asp Phe Cys Gln Ser Ser Asn Val Gly Asn Glu Ser Trp Tyr
 Phe Arg Gly Thr Ser Asn Gly Trp Gly Val Ser Ala Leu Thr Tyr Glu Ala Ala Thr Gly Leu Tyr Thr
 Thr Val Gln Ser Phe Asn Gly Glu Glu Ser Pro Ala Arg Phe Lys Ile Asp Asp Gly Asn Trp Ser Glu
 Ser Tyr Pro Ser Ala Asp Tyr Gln Val Gly Asp Tyr Ala Thr Tyr Thr Ile Thr Phe Asp Ser Gln Thr
 Lys Ala Ile Thr Val Thr Ser Gln

SEQ ID NO: 195

atgctgacagaccgtttcttgatggcgatacatcaacaacgacccttacaaccagaactacgatgctaaaaacgaccggggaacttatcagg
 gcggcgattttaagaatcacgcaaaaattggattatctcgataagctaggcgtgaacacaatctggatcagccccgacgttggaataatcaag
 catgatgctcggttatgacaactctgaagggcattcactatgcttaccacggctactgggcaagcaacttcgggtgcgttaaacccacacttcggt
 acaatggaagattccatacactgattgacgctgcccagaaaaggcatcaagatcattggtgacgtatgtaaacacacttggttatggctta
 aaagatatcaacggagaaatttccaatctccagccggttacccaactgacgcagaacgcagcacatatagcagcctgcttcgccagggttca
 aatgtcggctctgatgaggttggttggaattagctggcctacctgacttaaaaaacagaagaccocgcagtcgccagacaatcatcgactggc
 aaacagactgacacgaagcactacagctaaaggaaacacaattgacttccgttcgacactgtgaagcaggttgaaagacgcaacat
 ggatggcattcaaaaatgacccactgaaataatgacacacaaaatgattggggaagtgggggaagagggccnaaaagaaatggat
 acctgaaacagggtatgatggactcactgottgacttcgactcaaaaggatgacgacgatttcgtaacggcaagcttaaggcagcaaacgat
 gccctgactgcccgcaacggtaaaattgacaacacagctactttaaggttcattccttggagccatgacgaagatggttctatitaaagaagga
 aatgacaaaaggcaagcttaagggtgctgcttcctgcaagcaacatcaaaaggccagccggtcatctattatggtgaagagcttggtcaaaagt
 gagcaaaacactatccgcaatacgaataccgttatgacctggcatgggacaaggtgaaaacaacgacgtccttgagcactacactaaggtcct
 gaacttcagaagcgctcattcagaagtgttcgtaaaaggtaacgcgcaacaattggcggcttgacgctgataaattctacttttgcgtgtaa
 aatggaacgaagctgcttacgtcggcttgacgttgacacagcaaaagacgtaacactgactgtttcgcaggtgcagtcgtaactgacc
 actatgcagataaaaacttactgcttcagaagctggagaatcacattgacgatcccgcaaaagctgatggcggtactgtttactaacgggtg
 aaggcgagaaatcacagctgctaaagcggaagcgaagggcagggcacagttgagccagtcctcgcaaccacatccgattcactacaa
 ccgtacagacaacaactatgaaaactacgggtcatggctgtggaacgatgtagcctccctctgccaactggccgactggcgctacaatgttg
 aaaaacagacagctacgggtgcatatcgacgtaccacttaagaggcgctaaagacatcggcttcctggttatggtatgaacaaaagggtga
 tcagggttaagacggcggaacaaagggtttacgatctcatcctgaaatgaacgaatttgatcaagcaaggttgcgaaggtgtacatt
 acgagccagttgatcttcggcggaacactgtccgctccactatgtacgtgacaacgcagactacgaaaacttcggtatctggaactggggcga
 tgaacagcaccttcggaaaactggcctacagggcgagcgaaattcgttggtacagaccgttacgggtgctgtagcattacgctaaagaa

FIGURE 16FFFF

ggcgcaagaacattggaatgattgcttcaactgcaaatggagagaaagacggcgagataaatcttcaaccttctggataaatataatcg
catttgattaacaaggatgacaaatgtctacgtttctccatactgggagcaggcaacaggaatcaccaatgcagaggtaatctctgaagata
cgattctattaggcttcacaatgactgacggcttaacacctgaatctttaaaggaggtcttgaattaaagattcaactgggtgctgaagttgccatc
gaaagtgtgaaatcacaaagcgcaacctctgtaaaagtaaaagcaacattcgatttagaaaagcttcattatccatcacatacgcaggcagaac
agtttcagcttcaactggctggagaatgcttgatgaaatgtacgcttatgatgaaacgaccttgggtgcactacaaggacggagcagcgacg
cttaaatattgggtccgaaagcgagcaaggtaaccgctaactcttggataaaaaaatgccgctgaaaaaatcggcagcgtcaggttaacgaa
gggtgaaaaaggagctggtcagctatggtgctccggcgacctgaacgtaaccgacttgaagggtattttaccagtatgatgaacaaatga
cggataactcgcagggttagatccttatgcaaaatcaatggcagccttactgtgaatacagaaggcaatgctggtcctgacggggacactg
ttggcaaggcggaattcaaaaagcttctcgagagtacttctag

SEQ ID NO: 196

Met Leu Thr Asp Arg Phe Phe Asp Gly Asp Thr Ser Asn Asn Asp Pro Tyr Asn Gln Asn Tyr Asp
Ala Lys Asn Asp Arg Gly Thr Tyr Gln Gly Gly Asp Phe Lys Gly Ile Thr Gln Lys Leu Asp Tyr Leu
Asp Lys Leu Gly Val Asn Thr Ile Trp Ile Ser Pro Ile Val Glu Asn Ile Lys His Asp Val Arg Tyr Asp
Asn Ser Glu Gly His Ser Tyr Tyr Ala Tyr His Gly Tyr Trp Ala Ser Asn Phe Gly Ala Leu Asn Pro
His Phe Gly Thr Met Glu Asp Phe His Thr Leu Ile Asp Ala Ala His Glu Lys Gly Ile Lys Ile Met
Val Asp Val Val Leu Asn His Thr Gly Tyr Gly Leu Lys Asp Ile Asn Gly Glu Val Ser Asn Pro Pro
Ala Gly Tyr Pro Thr Asp Ala Glu Arg Ser Thr Tyr Ser Ser Leu Leu Arg Gln Gly Ser Asn Val Gly
Ser Asp Glu Val Val Gly Glu Leu Ala Gly Leu Pro Asp Leu Lys Thr Glu Asp Pro Ala Val Arg Gln
Thr Ile Ile Asp Trp Gln Thr Asp Trp Ile Thr Lys Ala Thr Thr Ala Lys Gly Asn Thr Ile Asp Tyr Phe
Arg Val Asp Thr Val Lys His Val Glu Asp Ala Thr Trp Met Ala Phe Lys Asn Asp Leu Thr Glu
Lys Met Pro Thr His Lys Met Ile Gly Glu Ala Trp Gly Ala Ser Ala Asn Asn Gln Leu Gly Tyr Leu
Glu Thr Gly Met Met Asp Ser Leu Leu Asp Phe Asp Phe Lys Gly Ile Ala His Asp Phe Val Asn Gly
Lys Leu Lys Ala Ala Asn Asp Ala Leu Thr Ala Arg Asn Gly Lys Ile Asp Asn Thr Ala Thr Leu Gly
Ser Phe Leu Gly Ser His Asp Glu Asp Gly Phe Leu Phe Lys Glu Gly Asn Asp Lys Gly Lys Leu
Lys Val Ala Ala Ser Leu Gln Ala Thr Ser Lys Gly Gln Pro Val Ile Tyr Tyr Gly Glu Glu Leu Gly
Gln Ser Gly Ala Asn Asn Tyr Pro Gln Tyr Asp Asn Arg Tyr Asp Leu Ala Trp Asp Lys Val Glu
Asn Asn Asp Val Leu Glu His Tyr Thr Lys Val Leu Asn Phe Arg Ser Ala His Ser Glu Val Phe Ala
Lys Gly Glu Arg Ala Thr Ile Gly Gly Ser Asp Ala Asp Lys Phe Leu Leu Phe Ala Arg Lys Asn Gly
Asn Glu Ala Ala Tyr Val Gly Leu Asn Val Ala Asp Thr Ala Lys Asp Val Thr Leu Thr Val Ser Ala
Gly Ala Val Val Thr Asp His Tyr Ala Asp Lys Thr Tyr Thr Ala Ser Glu Ala Gly Glu Ile Thr Leu
Thr Ile Pro Ala Lys Ala Asp Gly Gly Thr Val Leu Leu Thr Val Glu Gly Gly Glu Ile Thr Ala Ala
Lys Ala Ala Ser Glu Gly Asp Gly Thr Val Glu Pro Val Pro Ala Asn His Ile Arg Ile His Tyr Asn
Arg Thr Asp Asn Asn Tyr Glu Asn Tyr Gly Ala Trp Leu Trp Asn Asp Val Ala Ser Pro Ser Ala Asn
Trp Pro Thr Gly Ala Thr Met Phe Glu Lys Thr Asp Ser Tyr Gly Ala Tyr Ile Asp Val Pro Leu Lys
Glu Gly Ala Lys Asn Ile Gly Phe Leu Val Met Asp Val Thr Lys Gly Asp Gln Gly Lys Asp Gly Gly
Asp Lys Gly Phe Thr Ile Ser Ser Pro Glu Met Asn Glu Ile Trp Ile Lys Gln Gly Ser Asp Lys Val
Tyr Thr Tyr Glu Pro Val Asp Leu Pro Ala Asn Thr Val Arg Val His Tyr Val Arg Asp Asn Ala Asp
Tyr Glu Asn Phe Gly Ile Trp Asn Trp Gly Asp Val Thr Ala Pro Ser Glu Asn Trp Pro Thr Gly Ala
Ala Lys Phe Asp Gly Thr Asp Arg Tyr Gly Ala Tyr Val Asp Ile Thr Leu Lys Glu Gly Ala Lys Asn
Ile Gly Met Ile Ala Leu Asn Thr Ala Asn Gly Glu Lys Asp Gly Gly Asp Lys Ser Phe Asn Leu Leu
Asp Lys Tyr Asn Arg Ile Trp Ile Lys Gln Gly Asp Asp Asn Val Tyr Val Ser Pro Tyr Trp Glu Gln
Ala Thr Gly Ile Thr Asn Ala Glu Val Ile Ser Glu Asp Thr Ile Leu Leu Gly Phe Thr Met Thr Asp
Gly Leu Thr Pro Glu Ser Leu Lys Gly Gly Leu Val Ile Lys Asp Ser Thr Gly Ala Glu Val Ala Ile
Glu Ser Ala Glu Ile Thr Ser Ala Thr Ser Val Lys Val Lys Ala Thr Phe Asp Leu Glu Lys Leu Pro
Leu Ser Ile Thr Tyr Ala Gly Arg Thr Val Ser Ala Ser Thr Gly Trp Arg Met Leu Asp Glu Met Tyr
Ala Tyr Asp Gly Asn Asp Leu Gly Ala Thr Tyr Lys Asp Gly Ala Ala Thr Leu Lys Leu Trp Ala Pro
Lys Ala Ser Lys Val Thr Ala Asn Phe Phe Asp Lys Asn Asn Ala Ala Glu Lys Ile Gly Ser Val Glu
Leu Thr Lys Gly Glu Lys Gly Val Trp Ser Ala Met Val Ala Pro Gly Asp Leu Asn Val Thr Asp Leu
Glu Gly Tyr Phe Tyr Gln Tyr Asp Val Thr Asn Asp Gly Ile Thr Arg Gln Val Leu Asp Pro Tyr Ala

FIGURE
16GGGG

Lys Ser Met Ala Ala Phe Thr Val Asn Thr Glu Gly Asn Ala Gly Pro Asp Gly Asp Thr Val Gly Lys
Ala Ala Ile Gln Lys Ala Ser Arg Glu Tyr Phe

SEQ ID NO: 197

atgaaaccgtcaaaatcgttttctctgtgcccacgcttcgagcctctccagtagcccaatgctgacgccatttgcatttaactggaag
tactccgacgtcacgcaaacgccctcgcaaatcgccggcggttataaaaaagtgtgatttccagcactgaaatcgagtggaatgaa
tggtgggcacgttatcaaccgaagatctgcgctgacgattccccacttggcaacaaaagtactaaaatccatgattgatctctgaagc
ggctggcggtgatgtgatgccgatgtggtgcttaacatatggccaatgaacatggaagcgtgaagactaaattaccctggcagtgaaatgc
tgcaacaatacgcagctaacaccagttattatggcgacaaacgcttttggcaattaacggaaaacctattctctggtttgacttccaccaga
aggctgtattagcgattggaatgatgccggcaatgttcagtagcgtcttttggcggtgctggtgaccgagggtgcccagacttagatccga
acaactgggtggtgtcacagcaacgtttgtattgaatgcgctaaagggttaggtgtgaaaggctccgattgatgcggttaaacaatgagcc
aatatcaaatcgaccagattttcactgcagagattaccgccgaatgcacgtgttggtaagtgatcaccagtggtggcaaggcgacaccag
ctatgagaacttttagcgcttatctcaacgccaccaaccattcggttacgatttccactgttgcctctattcgaacgccttctctacagcgg
tggtcatgaacatgctcatgatccacaagcctatggccaagggttgaacgcacgttcaattacctttaccatcacgcacgacatcccaacga
acgacgggttccgttatcaaatcatggtaccgaaagatgaagagctggcttacgttatctcctggtaagatggcgccacacctctgattaca
gcgacaacttacctgatacgaagatcgtgataatcgccgttgggaagggtgttgaaccgtgacctgataagaacatgttgcgctccataac
caaatgcaagggaagagatgacgatgctgtacagcgaccaatgtctactgatgttaagcgcggttaacaagggtggtcggcattaataat
gcggtgaagagcgttctacaccgttgacacctatcagcatgagttcaactggtatcagccttacagatacactcactggcgtgactgaaacc
gtgagttcgcttaccacaccttccgaattccagctcgacgcgcgcgatgtacatgctctaa

SEQ ID NO: 198

Met Lys Pro Ser Lys Phe Val Phe Leu Ser Ala Ala Ile Ala Cys Ser Leu Ser Ser Thr Ala Asn Ala
Asp Ala Ile Leu His Ala Phe Asn Trp Lys Tyr Ser Asp Val Thr Gln Asn Ala Ser Gln Ile Ala Ala
Ala Gly Tyr Lys Lys Val Leu Ile Ser Pro Ala Leu Lys Ser Ser Gly Asn Glu Trp Trp Ala Arg Tyr
Gln Pro Gln Asp Leu Arg Val Ile Asp Ser Pro Leu Gly Asn Lys Ser Asp Leu Lys Ser Met Ile Asp
Ala Leu Lys Ala Val Gly Val Asp Val Tyr Ala Asp Val Val Leu Asn His Met Ala Asn Glu Thr Trp
Lys Arg Glu Asp Leu Asn Tyr Pro Gly Ser Glu Val Leu Gln Gln Tyr Ala Ala Asn Thr Ser Tyr Tyr
Ala Asp Gln Thr Leu Phe Gly Asn Leu Thr Glu Asn Leu Phe Ser Gly Phe Asp Phe His Pro Glu
Gly Cys Ile Ser Asp Trp Asn Asp Ala Gly Asn Val Gln Tyr Trp Arg Leu Cys Gly Gly Ala Gly Asp
Arg Gly Leu Pro Asp Leu Asp Pro Asn Asn Trp Val Val Ser Gln Gln Arg Leu Tyr Leu Asn Ala
Leu Lys Gly Leu Gly Val Lys Gly Phe Arg Ile Asp Ala Val Lys His Met Ser Gln Tyr Gln Ile Asp
Gln Ile Phe Thr Ala Glu Ile Thr Ala Gly Met His Val Phe Gly Glu Val Ile Thr Ser Gly Gly Lys Gly
Asp Ser Ser Tyr Glu Asn Phe Leu Ala Pro Tyr Leu Asn Ala Thr Asn His Ser Ala Tyr Asp Phe Pro
Leu Phe Ala Ser Ile Arg Asn Ala Phe Ser Tyr Ser Gly Gly Met Asn Met Leu His Asp Pro Gln Ala
Tyr Gly Gln Gly Leu Glu Asn Ala Arg Ser Ile Thr Phe Thr Ile Thr His Asp Ile Pro Thr Asn Asp
Gly Phe Arg Tyr Gln Ile Met Asp Pro Lys Asp Glu Glu Leu Ala Tyr Ala Tyr Ile Leu Gly Lys Asp
Gly Gly Thr Pro Leu Ile Tyr Ser Asp Asn Leu Pro Asp Asn Glu Asp Arg Asp Asn Arg Arg Trp
Glu Gly Val Trp Asn Arg Asp Leu Met Lys Asn Met Leu Arg Phe His Asn Gln Met Gln Gly Gln
Glu Met Thr Met Leu Tyr Ser Asp Gln Cys Leu Leu Met Phe Lys Arg Gly Lys Gln Gly Val Val
Gly Ile Asn Lys Cys Gly Glu Glu Arg Ser His Thr Val Asp Thr Tyr Gln His Glu Phe Asn Trp Tyr
Gln Pro Tyr Thr Asp Thr Leu Thr Gly Val Thr Glu Thr Val Ser Ser Arg Tyr His Thr Phe Arg Ile
Pro Ala Arg Ser Ala Arg Met Tyr Met Leu

SEQ ID NO: 199

gtgagtttgacaaaaaggctcagtagcaacaaatcggcaccgaaggctcagtagctctctgcaatcaatgccgcgcacacaacaatatcca
aatttacggcgatgtgtgttaccaccgaggtggtgctgatgggaagtcgtgggtcgataccaagcgcgttgattgggacacccgaatattg
aactggcgacaaatgattgaagcttgggttgagtttaatttcttggcgcaacgacaaatactgaacttcattggacttggatcactttgac
gggtgttactgggatgacgccggcaagaaaaagcgtctttaaattcaaggcgaaggaaaagcatgggattgggaagtcagctctgaaaa
aggcaattacgactacctaa

FIGURE
16HHHH

SEQ ID NO: 200

Val Ser Leu Thr Lys Lys Ala Gln Tyr Glu Pro Asn Thr Ala Pro Arg Leu Ser Thr Ser Leu Gln Ser
Met Pro Arg Thr Thr Thr Ile Ser Lys Phe Thr Ala Met Leu Cys Leu Thr Thr Glu Val Val Leu Met
Gly Ser Arg Gly Ser Ile Pro Ser Ala Leu Ile Gly Thr Thr Ala Ile Leu Asn Trp Ala Thr Asn Gly Leu
Lys Leu Gly Leu Ser Leu Ile Phe Leu Ala Ala Thr Thr Asn Thr Arg Thr Ser Ile Gly Leu Gly Ile
Thr Leu Thr Val Leu Thr Gly Met Thr Pro Ala Lys Lys Lys Arg Ser Leu Asn Ser Lys Ala Lys Glu
Lys His Gly Ile Gly Lys Ser Ala Leu Lys Lys Ala Ile Thr Thr Thr

SEQ ID NO: 201

atgacagccaaggctgatgacttacgcattaccagatcatgggtgaaagccttggatggcgataaacaggctcgccatggcaccggctacg
gtaccagccatcacaaaggcgatctgcaagggatcattgactcgcgtggattacattcaatcgctggcgctcaatgccatttggcctaacgccgattt
ttgaatctattccgggtggaggacaagaccattggcgagacaggcttgatgctacaggctacttggcagtgactattcaagatagaccctcgct
ttggcacgttagaacaagcccgtgagctgggtgaaaaggcacacgcgaaggcttgatgtcttcttggatggagtatttggcaccataaaggc
aatgtgggtgccatcaccacaaggtagactgcctgtcgggtgaaaataaccggctcagctacccagagagcctggcgtttacgaagaagtcgcc
agttactgggtgaaagagtaaagattgatggctggcgtctggatcaagcctatcaagtccgaccgatgcattgaaagcgatccgtcagagc
gttgatgaagcgtcacagtcgtaacttatgtgaataacaaagggaaccgtccatccttgggttacatgggtgctgaaatttgaataacgaa
cgttacatcacagaaccgggttacggcaaaagaaggcgatccggcgttgctcggctttgatttccgatgcgttccgagtggtcgaaacctti
ggcgttaacgaaagtggtgtcagccgaaaggcggaatggtgaatgacggcatgtcaetgcacagtcagtatccggatcatgccaagcct
aatttaattgtgggcaaccatgatgtggtgcgttggggtatctgctgaacgtggcggtattgctcaccagaacaaccgcaatactggcagcg
tcataaagcggcgatgtcttcttagcagcgtataccggccaattaccttgtattacggtagaagaattggcgatcaggttgacggccttgcataa
aaaatcaaagaagattgtgcccgttattgtgtgtgatgaccagtgggcgccagcagtgcaagattgatggcggtgacggcgctcactgaatg
cacagcagctgaactcaaatgatgtctctcattgatgacattacgtcgaacatcctgcgttatcacaaagggaacgtactaatgtgatggc
gacagagacagtatacgtagaccataaacaggcagacaatgaagccctgttgatcagtgtagtactgataacgcggagtcagtcacctt
gaaggcgaaagcgattgttcacaagggtgtgctgattgattgttaacgaacgagcgtttatgccaataatgggagtagtgcattccattaac
ggcgttggcgacgattcctcaagattgacactcgcagcggcggggtgtgatggcgcaatctgctgcctcggtatcgctagtaggtgaagg
gatcatggcccaatgtgatacccaaccgttgaaggcaccgggtccggtagcagaaccttgatcgtggttggcgatttggcagtgctggttggga
agcaaaagccgcagcgcgctatcaatacaaaaggcaagcacaatggcagcaacttgatcaagtgtgtgatgaaaagcggcgccctac
aagatgcaatacggcagaaagattggagccacagtttactgcagcggatggcattgaagccgggtaccgcaagtcgctcatagcgggt
ggctacggtaagacaccgcggtgacgttgccggaatccggtaagtgtgtggagcttaacattcagtgatcttggcgagccggagcaaatc
atggtgtctaatgtcagtaa

SEQ ID NO: 202

Met Thr Ala Lys Ala Asp Asp Leu Arg Ile Tyr Gln Ile Met Val Glu Ser Phe Val Asp Gly Asp Lys
Gln Val Gly His Gly Thr Gly Tyr Gly Thr Ser His His Lys Gly Asp Leu Gln Gly Ile Ile Asp Ser
Leu Asp Tyr Ile Gln Ser Leu Gly Val Asn Ala Ile Trp Leu Thr Pro Ile Phe Glu Ser Ile Pro Val Glu
Gly Gln Asp His Trp Ala Asp Arg Leu Asp Ala Thr Gly Tyr Phe Ala Ser Asp Tyr Phe Lys Ile Asp
Pro Arg Phe Gly Thr Leu Glu Gln Ala Arg Glu Leu Val Glu Lys Ala His Ala Lys Gly Leu Tyr Val
Phe Phe Asp Gly Val Phe Gly His His Lys Gly Asn Val Val Pro Ser Pro Gln Gly Arg Leu Pro Val
Gly Glu Asn Asn Pro Val Ser Tyr Pro Glu Ser Leu Ala Phe Tyr Glu Glu Val Ala Ser Tyr Trp Val
Lys Glu Leu Lys Ile Asp Gly Trp Arg Leu Asp Gln Ala Tyr Gln Val Pro Thr Asp Ala Trp Lys Ala
Ile Arg Gln Ser Val Asp Glu Ala Ser Gln Ser Val Thr Tyr Val Asn Asn Lys Gly Glu Thr Val His
Pro Leu Gly Tyr Met Val Ala Glu Ile Trp Asn Asn Glu Arg Tyr Ile Thr Glu Thr Gly Tyr Gly Lys
Glu Gly Asp Pro Ala Leu Cys Ser Ala Phe Asp Phe Pro Met Arg Phe Arg Val Val Glu Thr Phe Ala
Val Asn Glu Ser Gly Val Ser Arg Lys Gly Gly Glu Trp Leu Asn Asp Gly Met Ser Leu His Ser Gln
Tyr Pro Asp His Ala Lys Pro Asn Leu Met Leu Gly Asn His Asp Val Val Arg Phe Gly Asp Leu
Leu Gln Arg Gly Gly Ile Ala Ser Pro Glu Gln Pro Gln Tyr Trp Gln Arg His Lys Ala Ala Met Ser
Phe Leu Ala Ala Tyr Thr Gly Pro Ile Thr Leu Tyr Tyr Gly Glu Glu Ile Gly Asp Gln Val Asp Gly
Phe Ala Lys Lys Ile Lys Glu Asp Cys Ala Val Ile Gly Leu Cys Asp Asp His Val Ala Arg Thr Ser
Ala Lys Ile Asp Gly Val Thr Ala Ser Leu Asn Ala Gln Gln Ser Glu Leu Lys Val Tyr Val Ser Ser
Leu Met Thr Leu Arg Gln Gln His Pro Ala Leu Ser Gln Gly Glu Arg Thr Asn Val Met Ala Thr Glu

FIGURE 16III

Thr Val Tyr Val Asp His Lys Gln Ala Asp Asn Glu Ala Leu Leu Tyr Met Val Ser Thr Thr Asp Asn
 Ala Glu Ser Val Thr Leu Lys Gly Lys Ala Ile Gly Ser Gln Gly Val Leu Ile Asp Leu Leu Thr Asn
 Glu Arg Phe Met Pro Asn Asn Gly Glu Tyr Ala Ile Pro Leu Thr Gly Phe Gly Ala Arg Phe Leu Lys
 Ile Asp Thr Pro Thr Ala Ala Gly Val Met Ala Gln Ser Ala Ala Ser Val Ser Leu Val Gly Glu Gly Ile
 Met Ala Gln Cys Asp Thr Pro Thr Val Glu Gly Thr Gly Pro Val Ala Glu Thr Leu Tyr Val Val Gly
 Asp Phe Ala Asp Ala Gly Trp Lys Gln Lys Pro Gln Arg Ala Tyr Gln Tyr Lys Gly Lys His Asn Gly
 Ser Asn Leu Tyr Gln Val Val Val Asp Glu Lys Ala Gly Ala Tyr Lys Met Gln Tyr Ala Thr Lys Asp
 Trp Ser Pro Gln Phe Thr Ala Asp Gly Met Ala Leu Lys Pro Gly Thr Ala Lys Ser Leu Ile Ala Gly
 Gly Tyr Gly Lys Asp Thr Ala Val Thr Leu Pro Glu Ser Gly Lys Tyr Val Trp Ser Leu Thr Phe Ser
 Asp Leu Gly Glu Pro Glu Gln Ile

SEQ ID NO: 203

atgaagatgaagtcgccggcggtgttaggtagtgagtgccatggcggtggcctcttcggcagccaatgccggtgtcatgggtcacctgtt
 ccagtggaagtacaatgacatcgccaacgagtgcgaaaagggtgctcggtcccaaagggtatgaagcagtgagatcacgcccgtgtgaa
 cacctgcaagggtcctcctggtgggtgtctatcagccgctagctacaagaacttcacttctctggcggttaacgaggccgaactcaaaagca
 tgatcgcccggtgcaaggccgccggggtcaagattacgcgagtgccgtattcaaccagctggctgggtggtatcaggcgtcgggtacagggtgta
 gcagctacaatgccggcagcttcaagctatcccaatttggtacaaacgatttcacacgctgggagcctcaccactatgccagccgaacaa
 tgtgcaaaacggtgccctgtggggtgctcggtatcggtatccggctctgctatgtgcaggatcagctggctacctatatgaagacctgtagt
 ggctggggtgtggcaggttttctgttgatgcagcaaaagcatatgagcgttgccgatctcggccatcgtcagcaaggcgggcaatcctttgt
 ctactccgaggtgattgtggtccacgggtgaaccaatccagccggcggaatataccggcattgggtgctgaccgaatttaataacggcaccga
 tctggcctccaactcaaggggcagatcaagaatcaagagcatggcgagagctggggtctgcttgcgtcgaacaagggtgaagctttgtg
 gtcaacctgaccgtgagcggggacatggcggtggcggtatgctgacctacaaggatgggtccctctacaatctggccaacatcttcatgtg
 gcctggccctatggcgccatcccccaggtgatgtccggctatgatttggcaccataaccgatattgggtggcgagcgtacccttgttcttc
 ggctctagctggaactgcgaacaccgctggagcaacatcgccaacatggtctgctccacaatgccgcccaaggcagctccatgaccaactg
 gtgggataatgtaataaccagatcgctttgtgcggcgccaaggcctttgtggtgatcaacaatgaatcttccactctgagcaagagcctgc
 agacgggtctgccagccggggagtagtgcacatttgcggcggtgatgcctgtgcagcggcagcaccatcaagggtggtgaccagcggtat
 ggccaccttcaacgtggcaggggatgaaggcggcagcgatccatataatgccaaagccgatagcaccagcagtggtgagctcaggctcttct
 ctggctcttctctctgccaccagtaacaagtttgccagcatgaatctgcggggcaccacaatggctggggcagcaccgcatgacagtgga
 tgccaaccgtgtctggcggtgatgtcaccttaccggggcgcggtatgccaatgggtgccagcgttcaagtttgatgtctatggcaactgg
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 tcgctcaaggagagcgacatgagctacacctgaccagctctccagcaatcaggcaccgggtggcgccatcaccccaagacactctccgt
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 cccgaacagctggggcaccacggcgatgaagctgtggcantaacagotggcagggcgaggtgaccttaccgggcaagggcgatgca
 ctggtgcccacgctcaagttcagcgtcaagggtgactggagccagaactacgggtgacagcaacatggacgggactgccgaacggactgg
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SEQ ID NO: 204

Met Lys Met Lys Ser Arg Ala Trp Leu Leu Gly Ser Ala Val Ala Met Ala Leu Ala Ser Ser Ala Ala
 Asn Ala Gly Val Met Val His Leu Phe Gln Trp Lys Tyr Asn Asp Ile Ala Asn Glu Cys Glu Lys Val
 Leu Gly Pro Lys Gly Tyr Glu Ala Val Gln Ile Thr Pro Pro Ala Glu His Leu Gln Gly Ser Ser Trp
 Trp Val Val Tyr Gln Pro Val Ser Tyr Lys Asn Phe Thr Ser Leu Gly Gly Asn Glu Ala Glu Leu Lys
 Ser Met Ile Ala Arg Cys Lys Ala Ala Gly Val Lys Ile Tyr Ala Asp Ala Val Phe Asn Gln Leu Ala
 Gly Gly Ser Gly Val Gly Thr Gly Gly Ser Ser Tyr Asn Ala Gly Ser Phe Ser Tyr Pro Gln Phe Gly
 Tyr Asn Asp Phe His His Ala Gly Ser Leu Thr Asn Tyr Ala Asp Arg Asn Asn Val Gln Asn Gly
 Ala Leu Leu Gly Leu Pro Asp Leu Asp Thr Gly Ser Ala Tyr Val Gln Asp Gln Leu Ala Thr Tyr Met

FIGURE 16JJJ

Lys Thr Leu Ser Gly Trp Gly Val Ala Gly Phe Arg Leu Asp Ala Ala Lys His Met Ser Val Ala Asp
 Leu Ser Ala Ile Val Ser Lys Ala Gly Asn Pro Phe Val Tyr Ser Glu Val Ile Gly Ala Thr Gly Glu Pro
 Ile Gln Pro Gly Glu Tyr Thr Gly Ile Gly Ala Val Thr Glu Phe Lys Tyr Gly Thr Asp Leu Ala Ser
 Asn Phe Lys Gly Gln Ile Lys Asn Leu Lys Ser Met Gly Glu Ser Trp Gly Leu Leu Ala Ser Asn Lys
 Ala Glu Val Phe Val Val Asn His Asp Arg Glu Arg Gly His Gly Gly Gly Met Leu Thr Tyr Lys
 Asp Gly Ala Leu Tyr Asn Leu Ala Asn Ile Phe Met Leu Ala Trp Pro Tyr Gly Ala Tyr Pro Gln Val
 Met Ser Gly Tyr Asp Phe Gly Thr Asn Thr Asp Ile Gly Gly Pro Ser Ala Thr Pro Cys Ser Ser Gly
 Ser Ser Trp Asn Cys Glu His Arg Trp Ser Asn Ile Ala Asn Met Val Ser Phe His Asn Ala Ala Gln
 Gly Thr Ser Met Thr Asn Trp Trp Asp Asn Gly Asn Asn Gln Ile Ala Phe Gly Arg Gly Ala Lys Ala
 Phe Val Val Ile Asn Asn Glu Ser Ser Thr Leu Ser Lys Ser Leu Gln Thr Gly Leu Pro Ala Gly Glu
 Tyr Cys Asn Ile Leu Ala Gly Asp Ala Leu Cys Ser Gly Ser Thr Ile Lys Val Asp Ala Ser Gly Met
 Ala Thr Phe Asn Val Ala Gly Met Lys Ala Ala Ile His Ile Asn Ala Lys Pro Asp Ser Thr Ser
 Ser Gly Ser Ser Gly Ser Ser Ser Gly Ser Ser Ser Ala Thr Ser Asn Lys Phe Ala Ser Met Asn Leu
 Arg Gly Thr Asn Asn Gly Trp Ala Ser Thr Ala Met Thr Val Asp Ala Asn Arg Val Trp Ser Ala Asp
 Val Thr Phe Thr Gly Ala Ala Asp Ala Asn Gly Ala Gln Arg Phe Lys Phe Asp Val Tyr Gly Asn Trp
 Thr Glu Ser Tyr Gly Asp Thr Gln Ala Asp Gly Ile Ala Asp Lys Gly Ser Ala Lys Asp Ile Tyr Phe
 Asn Gly Val Gly Lys Tyr Arg Val Ser Leu Lys Glu Ser Asp Met Ser Tyr Thr Leu Thr Gln Leu Ser
 Ser Asn Gln Ala Pro Val Ala Ala Ile Thr Pro Lys Thr Leu Ser Val Lys Leu Gly Asp Ser Val Val
 Phe Asp Ala Ser Gly Ser Thr Asp Asp Val Gly Val Thr Gly Tyr Ser Trp Ser Thr Gly Gly Ser Ala
 Lys Thr Glu Thr Val Leu Phe Asp Ala Leu Gly Thr Lys Thr Ile Thr Val Thr Val Ala Asp Ala Asp
 Gly Leu Thr Ser Lys Ala Ser Ala Thr Val Thr Val Thr Asp Gly Ser Val Ala Tyr Asn Ser Asn Phe
 Ala Ser Leu Asn Phe Arg Gly Thr Pro Asn Ser Trp Gly Ala Ala Ala Met Thr Leu Val Ala Asp Asn
 Thr Trp Glu Ala Thr Val Asn Phe Asp Gly Gln Ala Asn Gln Arg Phe Lys Phe Asp Ile Lys Gly Asp
 Trp Ser Gln Asn Tyr Gly Asp Ser Asn Lys Asp Gly Val Ala Glu Arg Thr Gly Ala Asp Ile Tyr Thr
 Thr Val Thr Gly Gln Tyr Lys Val Gln Phe Asn Asp Ser Thr Leu Lys Tyr Thr Leu Thr Lys Leu Ala
 Asp Ser Ser Ala Thr Ser Tyr Ser Ala Asn Phe Ala Ser Leu Tyr Leu Arg Gly Thr Pro Asn Ser Trp
 Gly Thr Thr Ala Met Lys Leu Val Ala Asn Asn Ser Trp Gln Ala Glu Val Thr Phe Thr Gly Lys Gly
 Asp Ala Thr Gly Ala Gln Arg Phe Lys Phe Asp Val Lys Gly Asp Trp Ser Gln Asn Tyr Gly Asp Ser
 Asn Met Asp Gly Thr Ala Glu Arg Thr Gly Gly Asp Ile Thr Ser Ala Val Val Gly Thr Tyr Leu Val
 Thr Phe Asn Asp Ser Thr Leu Lys Tyr Thr Leu Thr Ala Lys

SEQ ID NO: 205

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 gactaaagaatcaggaaaactccgcaattttatgggaggcgatatcaagggtatcccaaaaaataatgaggggtatttttagtaactaggc
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 gaccctgccaaagcaaaaaatacttcagcatitggcaaaaactggcgatttcaaggaacaccccgagttgggtgcggaaggcacaacaaac
 ccttggcaaaaagccgttttacaccttttagcagggtttatcaaaaaatgttttatgacaaagtgtgtgtagcattagatgccctaaaggccaaa
 aacaaattaccgttaattgggtttttgatgacggtacaaaacttgtagtgcttattcaggcaagaacccctagttaaaaatggtatcgtttcacttt
 cttctgaatttgataatgtttgttgacacaaataa

FIGURE
16KKKK

SEQ ID NO: 206

Met Tyr Arg Val Ile Pro Ile Ile Leu Ile Met Ser Met Ile Val Ala Cys Glu Ser Pro Lys Lys Lys Thr
Thr Glu Thr Ala Gln Pro Ser Thr Asn Ala Glu Lys Pro Phe Val Trp Glu Ala Ala Asn Val Tyr Phe
Leu Leu Thr Asp Arg Phe Asn Asn Gly Asn Pro Asn Asn Asp Ile Asn Phe Asn Arg Thr Lys Glu
Ser Gly Lys Leu Arg Asn Phe Met Gly Gly Asp Ile Lys Gly Ile Thr Gln Lys Ile Asn Glu Gly Tyr
Phe Ser Lys Leu Gly Val Asn Ala Ile Trp Leu Thr Pro Val Val Glu Gln Ile His Gly Ser Val Asp
Glu Gly Thr Gly Asn Thr Tyr Ala Phe His Gly Tyr Trp Ala Lys Asp Trp Thr Asn Leu Asp Pro Asn
Phe Gly Thr Lys Glu Asp Leu Ala Glu Leu Val Ala Thr Ala His Ala Lys Gly Ile Arg Ile Leu Leu
Asp Val Val Ile Asn His Thr Gly Pro Val Thr Asp Gln Asp Pro Val Trp Gly Glu Asp Trp Val Arg
Thr Gly Pro Gln Cys Thr Tyr Asp Asn Tyr Thr Asn Thr Thr Ser Cys Thr Leu Val Ala Asn Leu Pro
Asp Ile Leu Thr Glu Ser Asn Glu Asn Val Ala Leu Pro Thr Phe Leu Leu Asp Lys Trp Lys Ala Glu
Gly Arg Leu Glu Gln Glu Leu Lys Glu Leu Asp Asp Phe Phe Ser Arg Thr Gly His Pro Arg Ala Pro
Arg Phe Tyr Ile Ile Lys Trp Leu Thr Asp Tyr Ile Arg Glu Phe Gly Val Asp Gly Phe Arg Val Asp
Thr Val Lys His Thr Glu Glu Thr Val Trp Ala Glu Leu Tyr Asp Glu Ala Val Ile Ala Phe Ala Glu
Tyr Lys Lys Ala Asn Pro Asp Lys Val Leu Asp Asp Asn Glu Phe Tyr Met Val Gly Glu Val Tyr
Asn Tyr Gly Ile Ser Gly Gly Arg Phe Tyr Asp Phe Gly Asp Lys Lys Val Asp Tyr Phe Asp His Gly
Phe Lys Ser Leu Ile Asn Phe Glu Met Lys Tyr Asp Ala Asn Phe Thr Tyr Asp Thr Leu Phe Arg Lys
Tyr Asp Thr Leu Leu His Thr Lys Leu Lys Gly Arg Ser Val Leu Asn Tyr Leu Ser Ser His Asp Asp
Gly Ser Pro Phe Asp Lys Met Arg Gln Lys Pro Tyr Glu Ser Ala Thr Lys Leu Leu Leu Thr Pro Gly
Ala Ser Gln Ile Tyr Tyr Gly Asp Glu Thr Ala Arg Ser Leu Asn Ile Glu Gly Ala Gln Gly Asp Ala
Thr Leu Arg Ser Phe Met Asn Trp Glu Glu Leu Ala Glu Asp Pro Ala Lys Gln Lys Ile Leu Gln His
Trp Gln Lys Leu Gly Ser Phe Arg Asn Asn His Pro Ala Val Gly Ala Gly Arg His Lys Thr Leu Gly
Lys Lys Pro Phe Tyr Thr Phe Ser Arg Val Tyr Gln Lys Asn Gly Phe Ile Asp Lys Val Val Val Ala
Leu Asp Ala Pro Lys Gly Gln Lys Gln Ile Thr Val Asn Gly Val Phe Asp Asp Gly Thr Lys Leu Val
Asp Ala Tyr Ser Gly Lys Glu Thr Ser Val Lys Asn Gly Ile Val Ser Leu Ser Ser Glu Phe Asp Ile
Val Leu Leu Glu Gln Lys.

SEO ID NO: 207

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gaggtcaaaagaagtcgtaggggctgtagccatcgaataggcgccgccatgcccttgctcgccgggggaatccaaatggcggaattccc
cctcgtaccactccggtatctgtccttgatggtgtccaccagattctccacctgggacgtccagtagaaggcctgcattataacgccgcct
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SEQ ID NO: 208

Leu Ser Thr Glu Pro Phe Val Leu Gly Ser Arg Leu Thr Leu Ser Pro Pro Arg Ser Ser Ser Arg Arg
Ser Ser Arg Asn Ser Arg Trp Pro Gly Arg Gly Gln Gly Pro Arg Gly Thr Pro Thr Arg Leu Ser Pro
Pro Thr Cys Pro Pro Ser Arg Arg Gly Cys Arg Cys Thr Arg Gly Cys Thr Leu Pro Arg Thr Ser Glu
Arg Arg Pro Thr Phe Arg Leu Cys Leu Arg Arg Gly Cys Met Leu Ser Val Pro Ala Cys Phe Arg

FIGURE 16LLLL

Ser Arg Phe Ser Arg Ile Ser Ala Arg Arg Cys Arg Ser Lys Arg Arg Gln Cys Phe Leu Arg Pro Gly
 Cys His Val Ser Arg Gly Ser Ala Tyr Pro Cys Ala Thr Pro Arg Ser Arg Gly Arg Ile Leu Ser Ala
 Gly Pro Arg Arg Gly Thr Arg Arg Leu Asp Thr Cys Ser Arg Leu Tyr Arg Cys Arg Gly Leu Gln
 Arg Arg Leu Arg Pro Thr Gly Arg Gly Arg Leu Cys Pro Arg Ser Gly Pro Arg Arg Val Arg Glu
 Cys Ser Cys Cys Gln Arg Pro Arg Pro Ser Cys Ser Arg Ala Gly Ser Arg Arg Pro Trp Arg Arg Ser
 Ser Arg Pro Ser Gly Val His Gln Arg Trp Cys Pro Ser Thr Arg Gln Arg Pro Ser Arg Pro Thr Ser
 Ala Ser Pro Arg Pro Thr Leu Arg Gly Pro Ser Arg Ser Gln Ser Ala Arg His Gln Arg Arg Cys Ser
 Leu Gly Arg Arg Arg Ser Ser His Arg Ser Pro Arg Ala Ser Ala Gly Pro Ser Ser Ser Arg Gly Leu
 Cys Leu Gly Ser Leu Gln Met Cys Pro Arg His Ser Thr Pro Arg Trp Gly Gly Ser Arg Gly Ser Trp
 Gln Tyr Ile Cys Pro Arg Pro Pro Leu Arg Ser Pro Ser Arg Cys Ser Pro Gln Arg Thr Gly Ser Thr
 Arg Gly Leu Arg Leu Arg Gly Gly Leu Arg Cys Pro Leu Pro Leu Cys Arg Arg His Gly Pro Cys
 Leu Ser Cys Ser Arg Ala Pro Ala Trp Ser Gln Ser Ala Ser Leu Pro Phe Pro Ser Gly Arg Thr His
 Arg Gly Gln Arg Ser Arg Arg Gly Arg Ser Pro Ser Asn Arg Arg Arg Pro Cys Pro Cys Ser Pro Gly
 Glu Ser Lys Trp Arg Ile Phe Pro Pro Arg Thr Thr Pro Val Ser Cys Ser Trp Cys Pro Thr Arg Phe
 Leu His Leu Gly Arg Pro Ser Arg Arg Pro Ala Leu Arg Arg Pro Leu Pro Ala Arg Ser Thr Trp Pro
 Val Thr Ser Tyr Ile Lys

SEQ ID NO: 209

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 atgtcacccctgcattccttaactggagctatgccgatgtcgtgatcgggcccgttgacatcgtcagcagggtacagtgccgtgctgtggccc
 cggcacttcgatccgaaggcacggcctggtggcgcgataccagccccaggatctccgcttatcgaccatccgtgggcaatacacatgacttc
 gtcaacatgatcgtctcgtatgtggtgtggcggtgtacgcccagacatcgtctcaaccacatggccaatgaggctgcacaaaggcctga
 cctgaactacccggtcaggcagtgcttgacgaatatgttccgatccccggctattcaggggcttgaggctgttcggtaacttgagcttcaatttct
 gtccgacatgatttggaccgccagtgcatcaggattacagcgtatgtttcaggccagaaactggcggctgtgcggaccgccggcggacc
 cgggctgcccgacctggtgccaatgactgggtgatctctcaacagcggcagtatctggaagccatcaaggcgtggtgtggtggtgatgagc
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 cggcgtgcttctggcttgggtggcagcatgagtgaactggtgatctgtcgtctacggcagccctgccaccggaccgcccatcaccctgct
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 ggggtgtcccgcttctgtattccgacaacaatgaaagcggcgatggcgctggatcgtcctggcaacgtccggatcgtgtgcaatgtcggct
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 acaagtgcggccatgcactcagctcctgggtcaacatgaaccagagcgtactgtggtgtacgggactacacagacgtgctcgacagcaacag
 cgtgtcaacatccagtcactcctggcacgagttcatccttcccggccagcagcctgtggttgcga

SEQ ID NO: 210

MIQPMHSREQACRLIPALIMTFALALPLQIRADVTLHAFNWSYADVADRAVDIAAAGYSA
 VLVAPPLRSEGTAWWARYQPQLRLIDHPLGNTHDFVNMIDALDDVGVGVYADIVLNM
 ANEAAQRPDLYTGGQAVLDEYASDPGHFGLRLFONLSNPLSEHDFGPAQCIQDYSDVF
 QVQNWRLCGPPDPGLPDLVANDWVISQQRQYLEAIKALGVAGMRIDAVKHMPMSHINA
 VLTPEIRSLGHVFGVTTSGGAGDTSYDRFLAPYLAQSDHGAYDFPLFETIRRAFVFGGMS
 ELVDPAAYGQALPPDRAITFVITHDIPNNDGFRYQILDVDESLEYAYILGRDGGVPLLYSD
 NNEGSDGRWIDAWQRPDLVAMVGFHNAVHGDMAVLSHDDCHLLFRRGSGLGIVGINKC
 GHALSSWVNMNQSVLWWYADYTDVLDNSVSVNIQSSWHEFILPARQARLWLR

SEQ ID NO: 211

GTGTTTCGTTCTGACACAGTTTCGCGTACCTGCATGTATGGTGCGCTGCGTAATGCCTA
 CCAACCCGATCGGGTGTITACTGGAGTCACGGTGCGGACATGCAACTTAAAAAAGCAT
 GCTCATCGCCAGGCGCTGTTGTTTCATCGTGACGCGGTGCCTGTGCCTGAAATCCAGGC
 AGACCCATAAAAAACAACAACAAACCGATAACAAACGACCCAAGCCTTCTAAGAGGAG
 AAAACGGGATGGCTTTTAAACTACGCAAAAAGGCGCTCGTTGGCCTGTTACGGCCGG

FIGURE
16MMMM

CGCAATGGTATATGCCGGTGCAGCGGCGAGTGGTGAAATCATTCTGCAGGGCTTCCAC
TGGCACTCCAAGTGGGGCGGCAACAATCAGGGTTGGTGGCAGGTGATGGAAGGTCAG
GCCAACACCATCGCCAACGCCGGCTTTACGCACGTGTGGTTCCCGCCGGTCCATAACT
CGGCCGATGCCGAGGGTTACCTACCCCGCGAGCTGAACAACCTCAACTCCAGCTATGG
CTCCGAAGCACAGCTGCGCAGCGCCATCCAGGCACTGAACAATCGCGGCGTGCATGCG
ATTGCCGATGTGGTCATGAACCACCGGGTGGGCTGCTCTGGCTGGGCGGATTTCTGTA
ACCCGGACTGGCCGACCTGGTACATCGTCGCCAATGATTCTGGCCCCGGTGGCCCCGAA
AAGCCAGAACTGGGACACGGGTGAGACGTACCACGCCGCCCGTGACCTCGATCACGC
CAATCCGCAGGTGCGCAACGATATCTCGCACTACCTGAACAGCCGCCTCAAGGACGTC
GGCTTCTCCGGCTGGCGCTGGGACTATGCCAAGGGTTTCTGGCCCCGGCTATGTCGGCG
AGTACAACCTGGAACACCAACCCGAACCTTCTGTGTGGGTGAGGTGTGGGACGATCTCGA
CCCCAACAAATCCCAACCCGCACCGCCAGCAACTGGTGGACTGGGTTGATGCTACCGGT
GGCAGTTGTCACGTCTTCGACTTCACCACCAAGGGGCTGACGAACTATGCGCTGCAGC
ATGGCCAGTACTGGCGCCTGCAGGGTGATAATGGTGGCCCGGCTGGCGGCATCGGCTG
GTGGCCGCAACGCATGGTGACCTTCGTCGACAACCATGACACGGGCCCGAGCAATCAC
TGTGGTGACGGCCAGAACCTCTGGCCCGTGCCCTGTGACAAGGTCATGGAGGCGTATG
CCTACATCCTGACCCATCCGGGCGTGCCGTGCGGTGTACTGGACGCACTTCTTCAACTGG
AATCTTGGTAGCGAGATCAGCCAGTTGATGCAGATCCGCAAGAACCAGGGCGTGCACT
CCGGTTCGACGTCTGGATCGCCGAGGCCCGTCACGGCCTGTACGCCGCCTATATCAA
CGGTAATGTGGCGATGAAGATGGGCTGGGATAACTGGAGCCCCGGGCTGGGGCTGGTC
GCTGGCGGCCTCCGGTAACAACCTGGGCGCTCTGGACACGCTGA

SEQ ID NO: 212

VFRSDTVSRTCMYGALRNA YQPDRVFTGVTVRTCNLKKHAHRQALLFVTRCLCLKSRQT
HKNNNKPIITNDPSLLRGENGMAFKLRKKALVGLFTAGAMVYAGAAASGEILLQGFHWHS
KWGGNNQGWVQVMEGQANTIANAGFTHVWFPPVHNSADAEGYLPRELNNLNSSYGSEA
QLRSAIQALNNRGVHAIAADVVMNHRVGC SGWADFCNPDWPTWYTVANDSWPGGPKSQN
WDTGETYHAARDLDHANPQVRNDISHYLSRLKDVGFSGWRWDYAKGFWP GYVGEYN
WNTNPNFCVGEVWDDLDPNPNPNHRQQLVDWVDATGGSCHVFDFTTKGLTNYALQH GQ
YWRLQGDNGGPAGGIGWWPQRMVTFVDNHD TGPSNHCGDGQNLWPVPCDKVMEAYA
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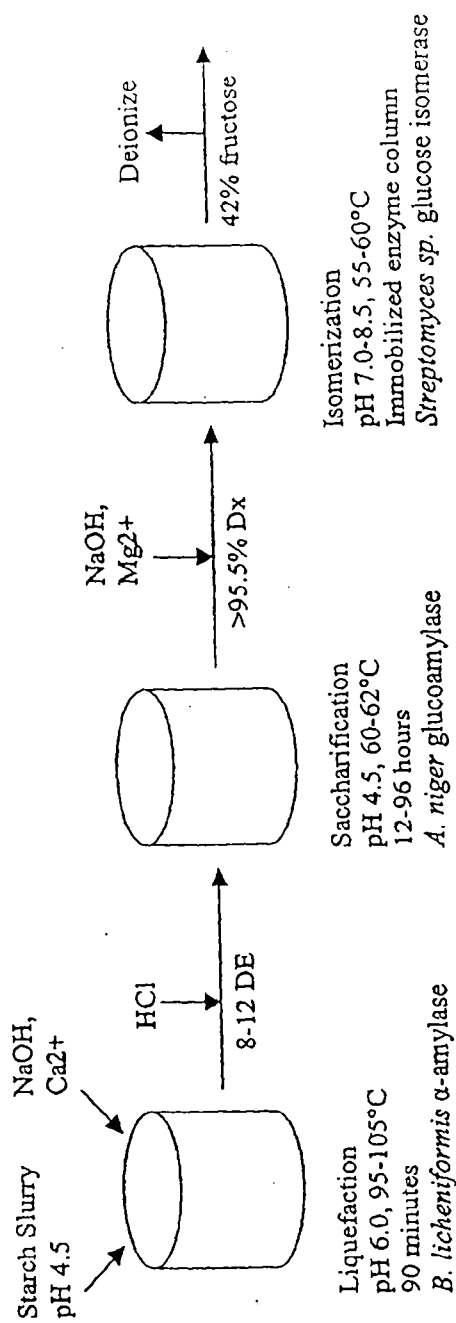


FIGURE 17

	SEQ ID NO.: 81	Pyro	Pyro	thermo	therm2	SEQ ID NO.: 75	SEQ ID NO.: 77	SEQ ID NO.: 83	SEQ ID NO.: 85	SEQ ID NO.: 79	SEQ 437
SEQ ID	100	91.7	75.1	82.1	80.1	82.5	82.6	82.1	82.6	83	77.8
pyro		100	74.8	82.5	80.5	82	82.2	82.9	82.8	84	78.5
Pyro2			100	71.5	71.1	74	74.2	77	77.1	73	70.5
therm				100	81.7	83.5	83.8	82.8	83.2	83.8	76.4
therm2					100	88.9	88.8	84.1	84.7	84	76.3
SEQ ID NO.: 75						100	98.3	84.6	85.2	85.5	77
SEQ ID							100	84.8	84.9	85.4	77.4
SEQ ID								100	96	83.3	78.5
SEQ ID									100	83	78.1
SEQ ID										100	79.8
Clone A											100

FIGURE 18

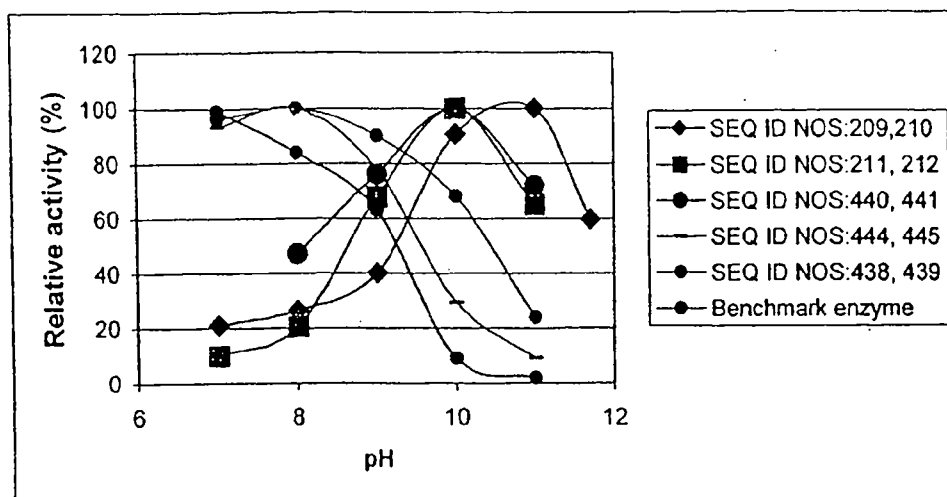


FIGURE 19

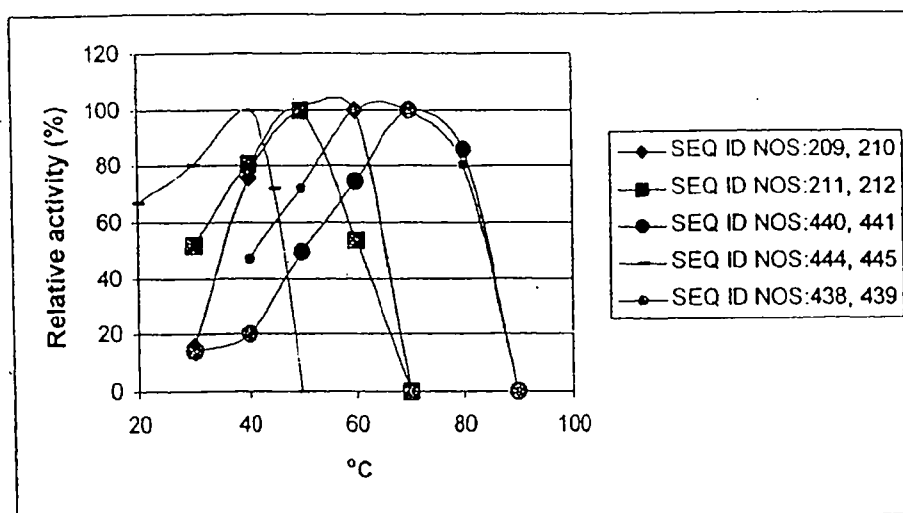


FIGURE 20

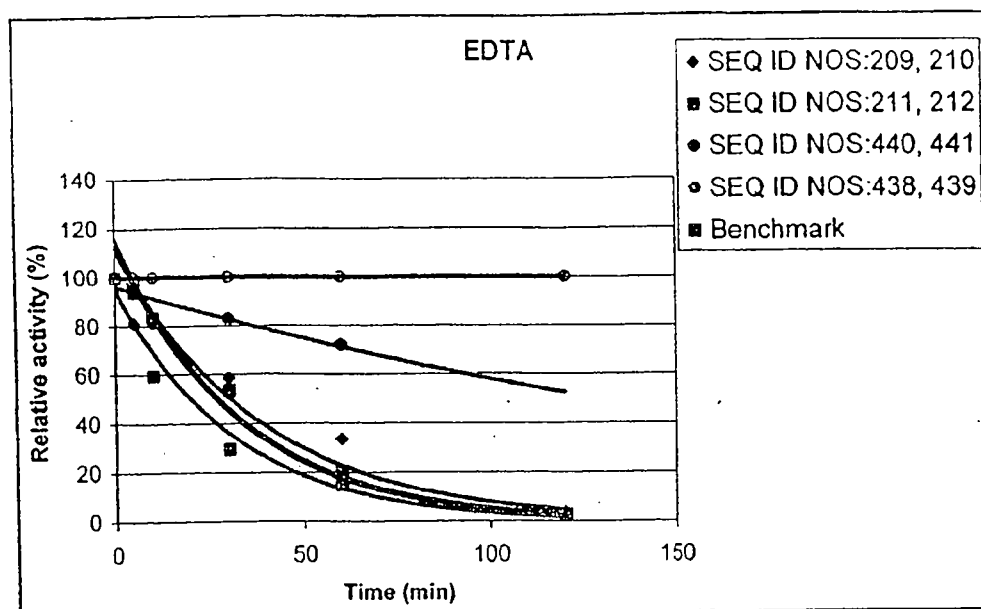


FIGURE 21

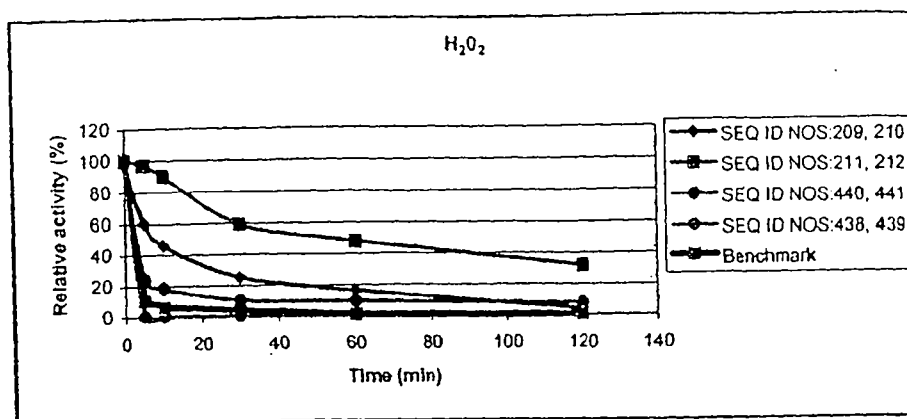


FIGURE 22

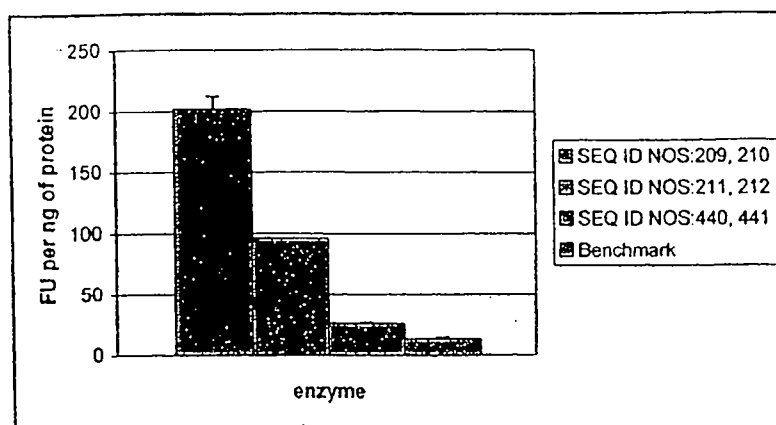


FIGURE 23

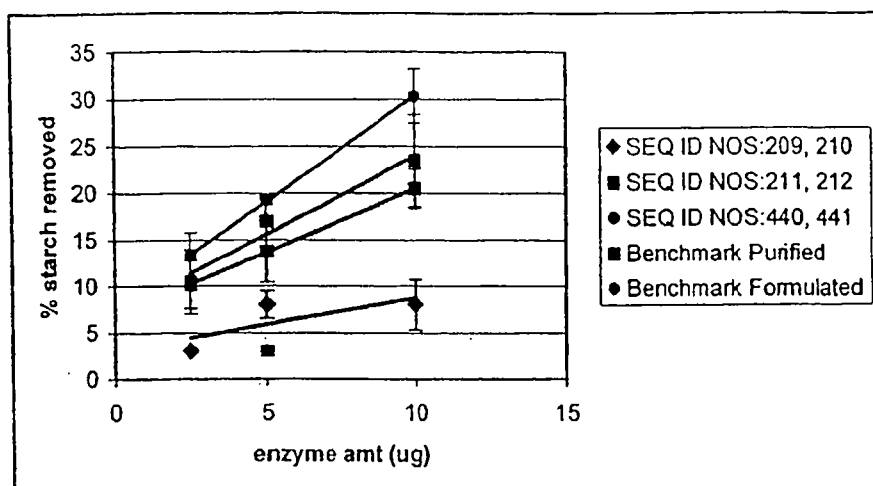


FIGURE 24

FIGURE 25

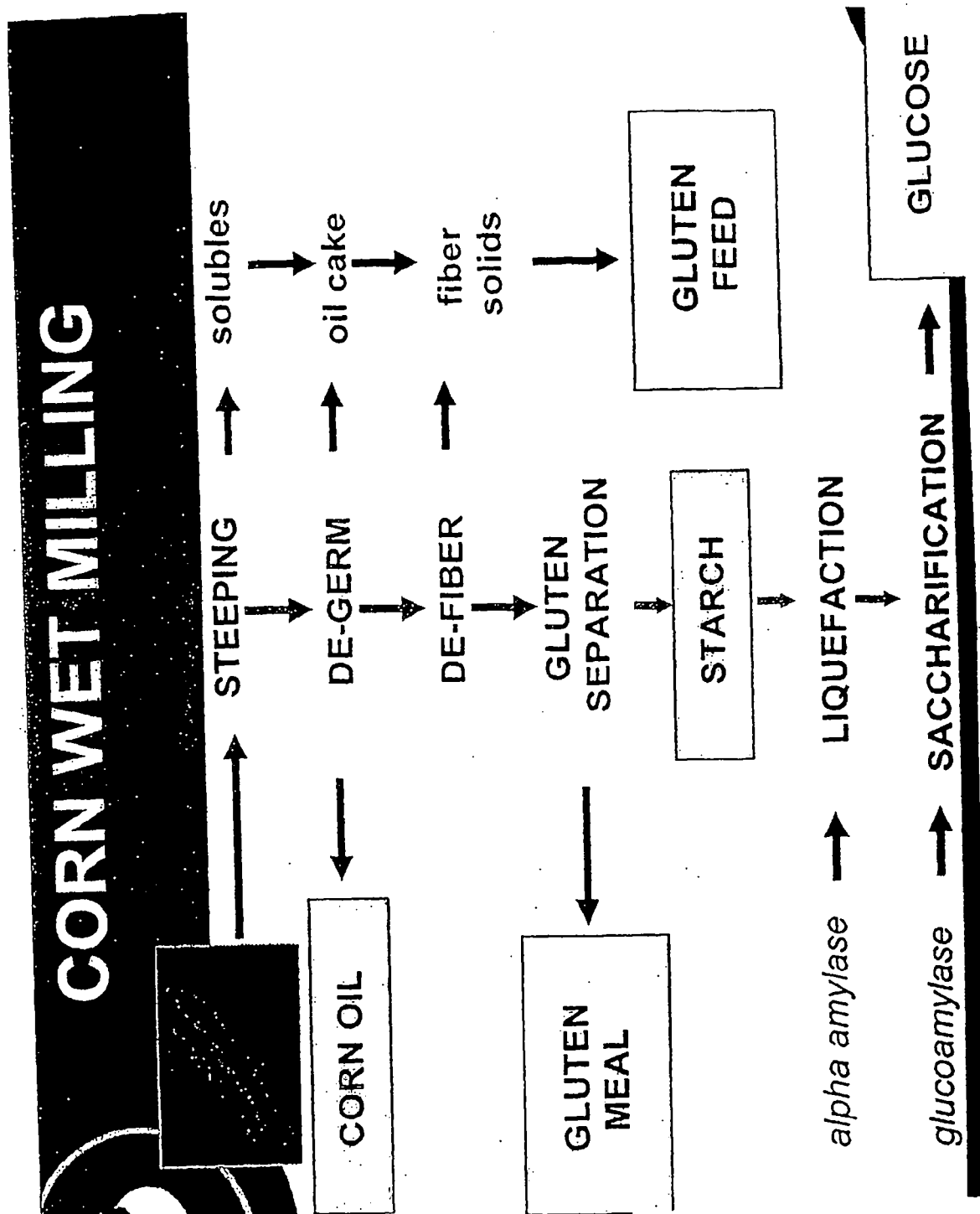


FIGURE 26

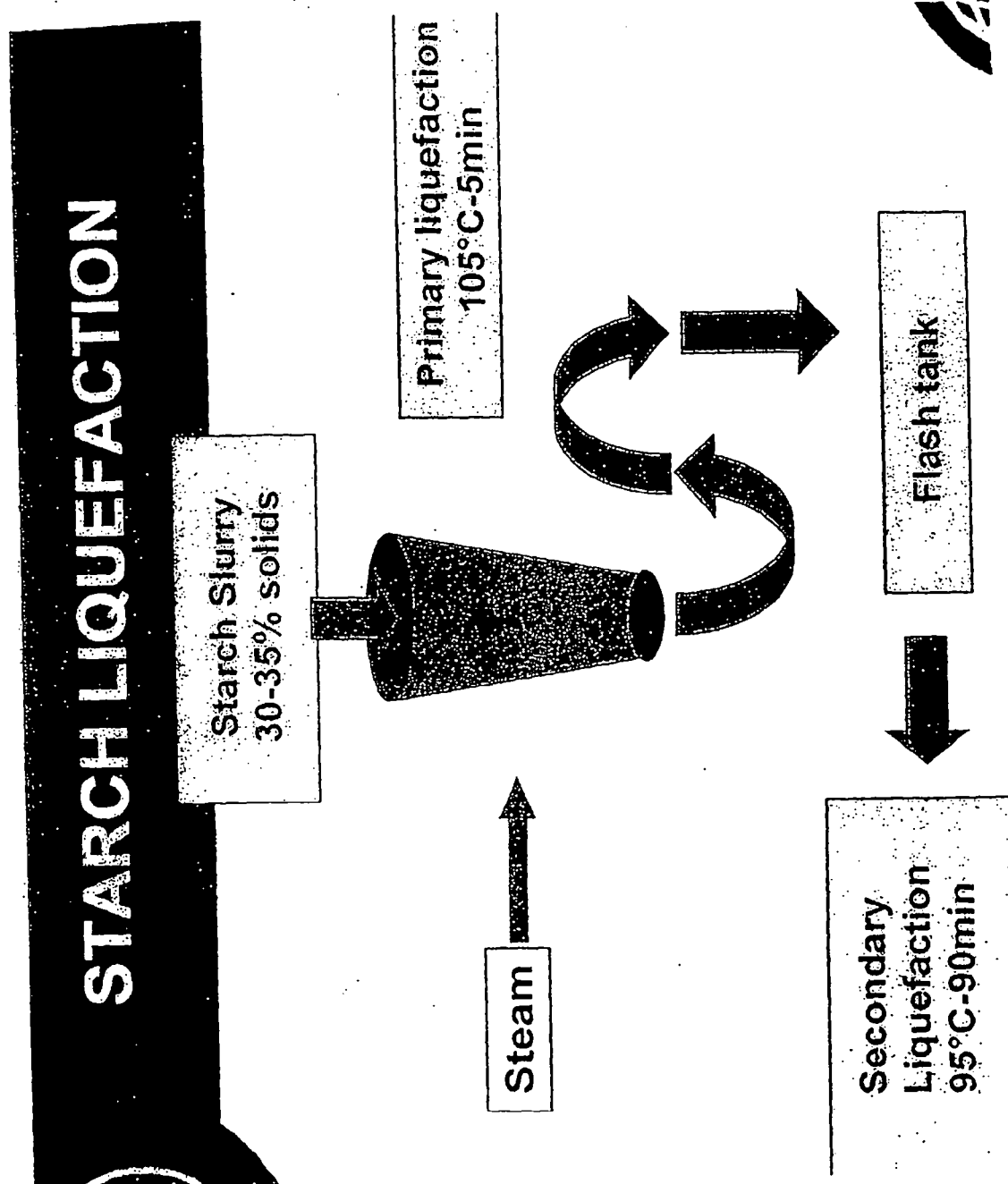


FIGURE 27

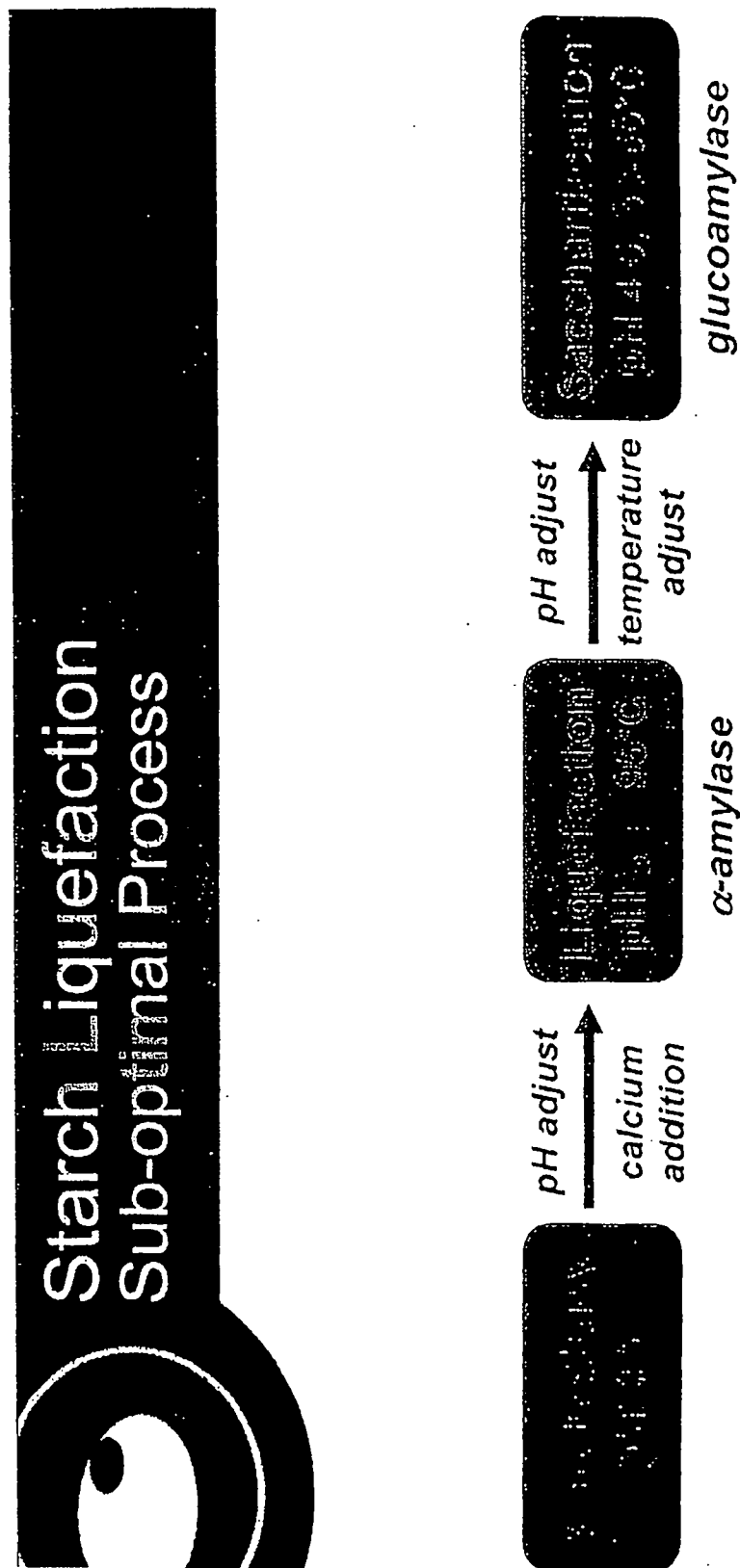
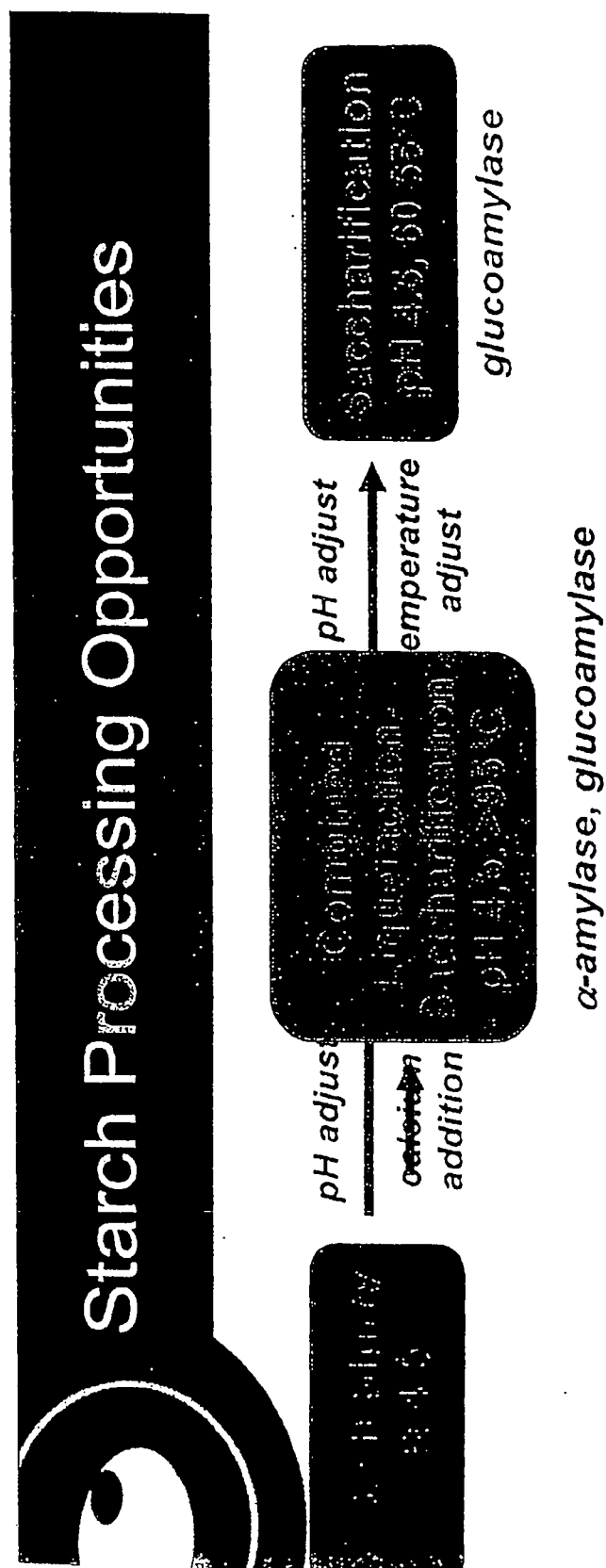


FIGURE 28



Operating Conditions	D45	Termamyl® SC Amylase
Temperature	95°C	85°C
pH range	4.4 – 5.6	5.0 – 6.0 pH optimum 5.7
Ca++	No significant impact on viscosity reduction	Not required
Recycled Backset	Up to 30% well tolerated	Up to 30% well tolerated
Enzyme Dose	0.4 – 0.6 kg/MT starch	0.3 – 0.5 kg/MT starch
DE after liquefaction	6-10	12-14

FIGURE 29

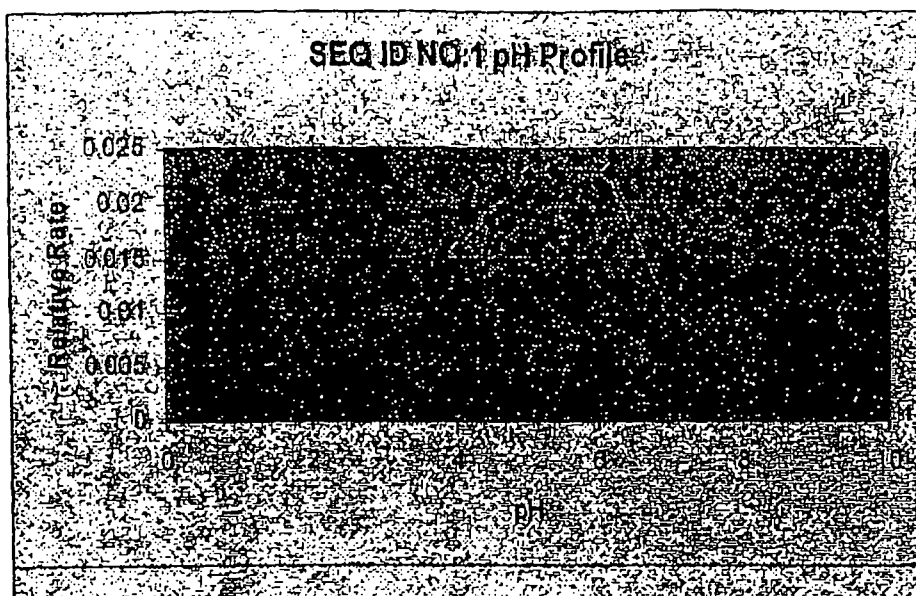


FIGURE 30

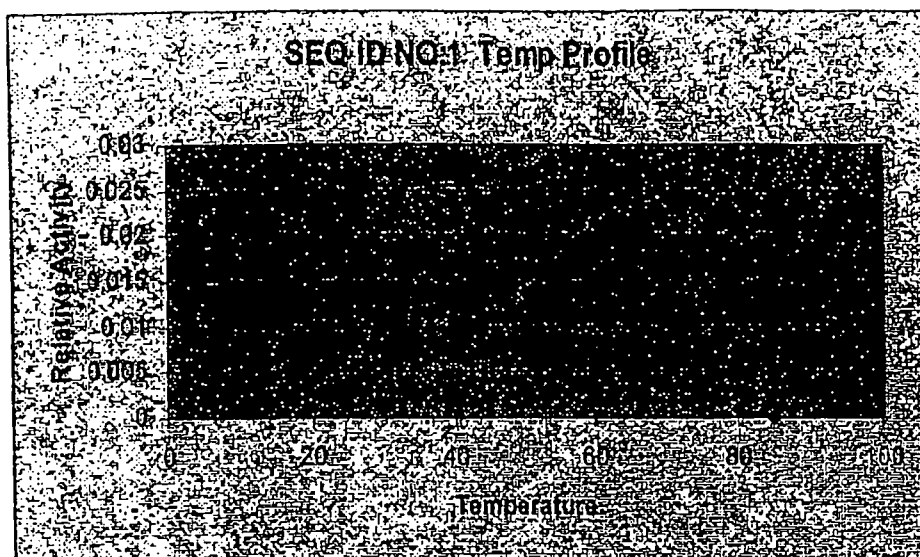


FIGURE 31

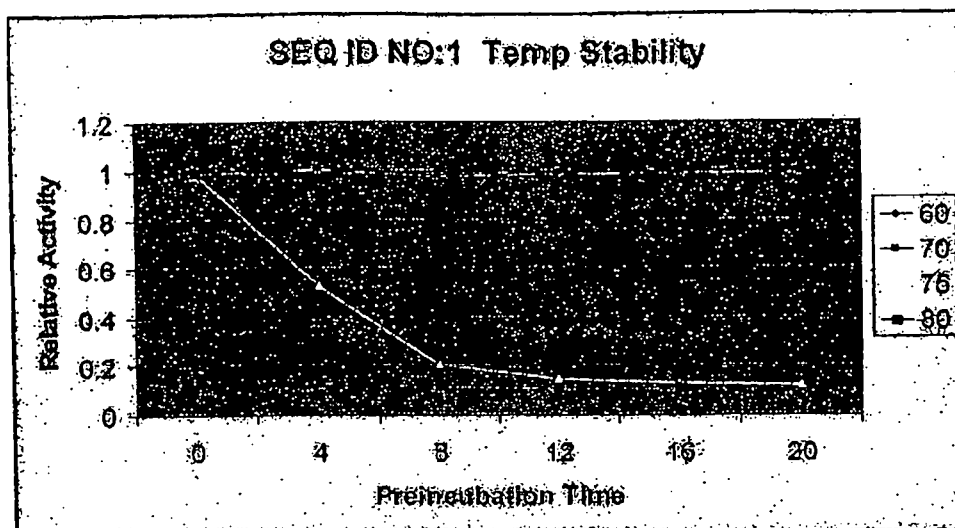


FIGURE 32

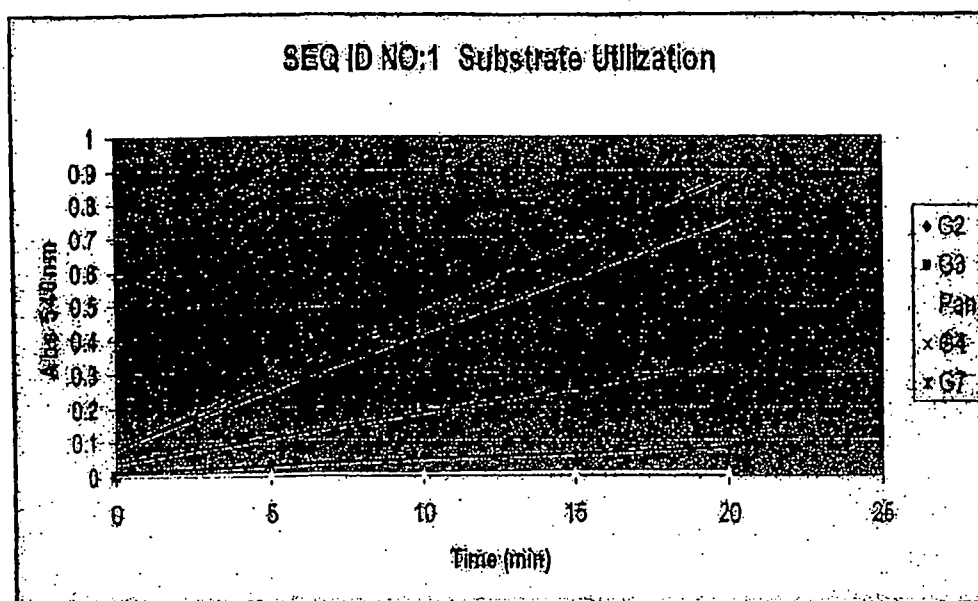


FIGURE 33

	*	*	30	*	*	60
tccaaagaattgaacgctcgtcggatgatcgtgatggtcagttttgaggtatcttccgaag						
	*	*	90	*	*	120
agccaatttggattacccggcagtcgggcagtcggacgggtccagacaaggatcaatcagg						
	*	*	150	*	*	180
gcctggcactattttagcatgtcgggttagtccccaatcgggtctgcatttgacagcggac						
	*	*	210	*	*	240
atctgccgatcaggaacgaaacgggtgcctgccgttggtgccctgtgagctctttatgcac						
	*	*	270	*	*	300
gtcaattcataataatcccgataataggccgcagctattccaccctggggaaaagggttg						
	*	*	330	*	*	360
ttgaatgtcggcgatttgggggtacatctgggggttcgggcacttcagtgctccagggtccg						
	*	*	390	*	*	420
aatttcacatgcattcatgcacgaaaatgcggaggggaacctcgggcactgtcaccatccccgt						
	*	*	450	*	*	480
caaactacgggcttttggcggggtacattgggtacaatggccacaatgggtacaacggcct						
	*	*	510	*	*	540
gtcaagggtatccattaatttctccgaagctcctgcaacgtttcgacaccctaaatttccg						
	*	*	570	*	*	600
tttccgctacggagagcggaggtaccggataactttttggacttttcatgtgcttctgtcgg						
	*	*	630	*	*	660
gcattcaaagaccgggacctgtcctaacgtccaggggatcggttggcagatcttcctcat						
	*	*	690	*	*	720

FIGURE 34A

```

gcgttcacagagegaagtttgtttaattgagtcacaaaacATGTTATTCCAACCGACTT
      *          *          750          *          *          780
      *          *          *          *          *          *
TGTGCGCGGCCCTTGGACTCGCCGCTTGATCGTCCAAGGCGGAGAAGCCAGACCTGAAA
L C A A L G L A A L I V Q G G E A R P E
      *          *          810          *          *          840
CAACCGTCCCACATGCAACGGGCTCGCTCGACGACTTCCTCGCCGCACAGAGTCCGATTG
T T V P H A T G S L D D F L A A Q S P I
      *          *          870          *          *          900
CTTTCCAAGGCATCCTGAACAATATCGGGCCTAGCGGAGCGTACTCGGAAGGTGTCAATC
A F Q G I L N N I G P S G A Y S E G V N
      *          *          930          *          *          960
CGGGTGTGGTCATTGCGAGTCCAAGTAAACAAGATCCCGACTgtatgcctgctctggaaa
P G V V I A S P S K Q D P D
      *          *          990          *          *          1020
tttttcaattctgttggcaggactctctctctaatatggcacatagACTTTTACACCTGG
      *          *          1050          *          *          1080
      *          *          *          *          *          *
GTGCGGACGCTGCTCTCACTGTCCAATATCTGGTGGAGGAGCTGGTTGCAGGAAATGCC
V R D A A L T V Q Y L V E E L V A G N A
      *          *          1110          *          *          1140
AGTCTTCAGTTCCTCATTGAGGACTACATCAGCTCCCAGGCACGACTGCAGACGGTGGAA
S L Q P L I Q D Y I S S Q A R L Q T V E
      *          *          1170          *          *          1200
AATCCATCCGGCTCCCTCTCGTCGGGTGGTCTAGGAGAGCCCAAGTTTCATGTCGACGAG
N P S G S L S S G G L G E P K F H V D E
      *          *          1230          *          *          1260
ACCGCCTTTACGGAATCCTGGGGCCGACCACAGCGGGACGGCCCGCCTCTCCGCGCCATT
T A F T D S W G R P Q R D G P P L R A I
      *          *          1290          *          *          1320
GCCATGATTTGTTTGCCAATTACCTGATTgtaagtcagatttcccatcatgcgagtaaa
A M I S F A N Y L I
      *          *          1350          *          *          1380
ttgacatggatgtgctcagtgtagtttttcagGACAACGGTCATCAATCGACTGTGGAGGA
      *          *          1410          *          *          1440
      *          *          *          *          *          *
CATCATCTGGCCGATTGTTGCAATGACTTGTCTATGTCTCGCAGCATTGGAACGAAAC
I I W P I V R N D L S Y V S Q H W N E T
      *          *          1470          *          *          1500
AACTTTTggtatgtgcttacgccgtactagttgattggagagtttggattataggagagc
T F
      *          *          1530          *          *          1560
      *          *          *          *          *          *
ctcaagctaatacggagtttttccgaaGACATCTGGGAGGAAGTCCATAGCTCATCGTTT
      *          *          1590          *          *          1620
      *          *          *          *          *          *
TTCACCACGGCTGTCCAGTACCGTGCTCTGGTCCAAGGCAGTGCCTTGGCTAGCAAGCTC
F T T A V Q Y R A L V Q G S A L A S K L
      *          *          1650          *          *          1680
GGCCATACCTGCGACAACCTGCGGGTCCCAAGCACCAGATCCTTTGCTTCCTGCAGTCG
G H T C D N C G S Q A P Q I L C F L Q S
      *          *          1710          *          *          1740
TATTGGACCGGGTCCGACATCTTAGCCAACACCGGTGGCGGCCGCTCGGGAAGGACGTC

```

FIGURE 34B

```

Y W T G S H I L A N T G G G R S G K D V
      * 1770 * 1800
AGCACGATCCTCGGCGTCATTGGCTCGTTTGATCCGAACGCCGACTGTGATGACGTTACC
S T I L G V I G S F D P N A D C D D V T
      * 1830 * 1860
TTCCAGCCCTGCTCGGCCCGGGCTCTTGCAAATCACAAGCAGGTCGTTGACAGCTTCCGC
F Q P C S A R A L A N H K Q V V D S F R
      * 1890 * 1920
AGTATCTATGCCATCAACGCTGGCATCCCGTCAGGGTCGGCTGTTGCGGTTGGACGTTAT
S I Y A I N A G I P S G S A V A V G R Y
      * 1950 * 1980
CCCCGAGGATGTCTATCAGGGTGGACACCCCTGGTACCTAACAACGGCTGCGGCGGCGGAG
P E D V Y Q G G H P W Y L T T A A A A E
      * 2010 * 2040
CAGCTTTACGACGCCATTTACCAGTGAACCATGTAGGGCACATCGACATCAATGCTGTC
Q L Y D A I Y Q W N H V G H I D I N A V
      * 2070 * 2100
AATCTGGACTTCTTCAAGAGCATTATCCGTCAGCCGCCGAGGGCACATACACATCAGAC
N L D F F K S I Y P S A A E G T Y T S D
      * 2130 * 2160
TCTTCAACATTTCAAGACATTATATCTGCTGTACGGACCTATGCGGACGGGTTTCTCAGC
S S T F Q D I I S A V R T Y A D G F L S
      * 2190 * 2220
GTAATTgtaagtccaaaccttcgaaaacgaatgcctcaagtottccactgacattttgcg
V I
      * 2250 * 2280
cagGAGAAATACACTCCGCCGGATAACTTGCTTGCCGAGCAGTTCCACCGGGAGACGGGC
E K Y T P P D N L L A E Q F H R E T G
      * 2310 * 2340
ATTCCACTATCGGCAGCTTCTCTGACATGGTCTTACGCCGCGCTCAACACGGCCGCGCAG
I P L S A A S L T W S Y A A L N T A A Q
      * 2370 * 2400
CGGCGAGCGTCAATCGTGCCCTCACCGTGGAACCTTAACAGCACAGATCTCCCGGACAAA
R R A S I V P S P W N S N S T D L P D K
      * 2430 * 2460
TGCTCGGCAACCTCGGCAACAGGGCCGTATGCCACGCCCCACAAACACGGCATGGCCAACC
C S A T S A T G P Y A T P T N T A W P T
      * 2490 * 2520
ACTACGCAGCCACCGAGCGGCCGGCATGCACACCGCCGTCGGAAGTAACACTCACCTTC
T T Q P P E R P A C T P P S E V T L T F
      * 2550 * 2580
AACGCGCTCGTCGACACCGCGTTTGGCCAGAATATTTATCTCGTGGGCTCCATTCCGGAG
N A L V D T A F G Q N I Y L V G S I P E
      * 2610 * 2640
CTCGGATCGTGGGATCCGGCCAACGCCCTCTTGATGAGCGCAAAGAGCTGGACTAGCGGA
L G S W D P A N A L L M S A K S W T S G
      * 2670 * 2700
AATCCGGTCTGGACGCTATCCATTTCCCTTCCAGCAGGAACCTCTTTTGAGTACAAGTTC
N P V W T L S I S L P A G T S F E Y K F
      * 2730 * 2760
ATTCGAAAGGATGATGGTTCCTCGGATGTTGTCTGGGAAAGTGACCCGAATCGTTCGTAC
I R K D D G S S D V V W E S D P N R S Y

```

FIGURE 34C

```

      *      *      2790      *      *      2820
AACGTGCCGATAGGATTGCGGTGCCAACACGGCCACCGTGAATTCTTGGTGCGGATGAacc
N V P K D C G A N T A T V N S W W R *
      *      *      2850      *      *      2880
aacttggtttctgtccacactccgccttggtgtcagttcctggtcgtagatcgataaaata

      *      *      2910      *      *      2940
tgacttggtgacttgacaaagaaatgatgtaaaagcgttctgttatgtagtaggtagca

      *      *      2970      *      *      3000
cttttccttagtagggagtactcgttaggtatgccgataccgaactccgaccggagtaaa

      *      *      3030      *      *      3060
actaacgtgtcgagtatcgcgatggtgcgcgtggggagtaaggacgattagggtgaatg

      *      *      3090      *      *      3120
ctgcagatcctttccttcaccgtctcagagacacggagtcaggtttgaccatggccgcgg

      *      *      3150      *      *      3180
tgtgaagctggtgatgctcattttcctgtccaatatttcaaggacaacgttgccaacatc

      *      *      3210      *      *      3240
aatcccagcaatgagtcctcgtgttgactgtgctgcctgggatcagagacttcgggaat

      *      *      3270      *      *      3300
acattgaagttagcaaaacaatgccgtccactaactatgtgcgctcacgtctaccaagtg

      *      *      3330      *      *      3360
cacgatccgttctgtggaaggggaagttatcctggtacaggcttgcttgcgattctggat

      *      *      3390      *      *      3420
cgactgcgaaaaaagaggggacatgcgtctggaaagttgcgattcaaaacgatgtcatc

      *      *      3450      *      *      3480
agctcattgcctcttgaaatcctgctcgagatcagagattatcttgaaactaaacgatatt

      *      *      3510      *      *      3540
ctccgaagccggatggtgcgtatcgtcttggtgctcgatggtgaaagagacatttttatt

      *      *      3570      *      *      3600
tgacaggagcaaacgtccgaacaagtctcgaaacaatagecgtctattttctcaagcgg

      *      *      3630      *      *      3660
ccagtcatgagaccttttctgcgggaggctcttgcgctcctggacatgaaaggcacggag

      *      *      3690      *      *      3720
atatecgccacggacgttatggcttattttcgatggtacgggtggattgaaatatggaaag

      *      *      3750      *      *      3780
ccagtaaagaaaatgtttctcccatggccagaaggccggataagattggagcaaaggatt

      *      *      3810      *      *      3840

```

FIGURE 34D

gagatttatcgcggcggtatattatcatgaatggaagaagagagcagtcggaatgttg
* * 3870 * * 3900
gatctggaaactagaaagaggcggctttggcccgcaaacgataaacacgtactttgaattc
* * 3930 * * 3960
tttgcacatcgatcgatatcttttgatctcggggtaagttgaggacaggatagtagattcc
* * 3990 * * 4020
tgttggctcgacaggtcgtctaactgcagtgctaggaacgctctgattgcgtgggatata
* * 4050 * * 4080
caggagtcgcggaaagtggggagattagtggtgagccctggcaggtgaacatcgtgcag
* * 4110
gataaggtcgtcatgtttgaacgaaagtcaa (SEQ ID NO: 1)

FIGURE 34E

FIGURE 35A

[illegible]

FIGURE 35B

129, 130	gi 12718271 gb AAE4461.1	5E-68	alpha-amylase [Pseudomonas sp.]	Proteus mirabilis	NO_HITS	26	38	817	480
131, 132	gi 1535792 gb BAA01800.1	0.11	malonate decarboxylase [Pseudomonas sp.]	Pseudomonas sp.	NO_HITS	61	76	828	614
133, 134	gi 1722279 gb AAA63390.1	0.11	alpha-amylase [Bacillus sp.]	Bacillus sp.	NO_HITS	97	84	548	613
135, 136	gi 1722279 gb AAA63390.1	0.11	alpha-amylase [Bacillus sp.]	Bacillus sp.	NO_HITS	84	84	844	613
137, 138	gi 207462 gb AAA4227.1	0.48	topoisomerase [Bacillus sp.]	Bacillus sp.	NO_HITS	19	11	439	307
139, 140	gi 11228329 emb CAG1.1	0.173	stereohydroxylase [Bacillus sp.]	Bacillus sp.	NO_HITS	60	59	507	515
141, 142	gi 15601018 ref NP_232648.1	0	alpha-amylase [Vibrio cholerae]	Vibrio cholerae	NO_HITS	70	67	466	486
143, 144	gi 15601018 ref NP_232648.1	0	alpha-amylase [Vibrio cholerae]	Vibrio cholerae	NO_HITS	69	65	473	486
145, 146	gi 12855802 gb AAK00698.1	0	alpha-amylase [Bacillus megaterium]	Bacillus megaterium	NO_HITS	91	90	513	533
147, 148	gi 19031818 gb AAAF829.1	1E-77	beta-glucanase [Pseudomonas sp.]	Pseudomonas sp.	NO_HITS	28	36	781	842
149, 150	gi 19031818 gb AAAF829.1	1E-77	beta-glucanase [Pseudomonas sp.]	Pseudomonas sp.	NO_HITS	24	36	834	842
151, 152	gi 14587798 emb CAC4.1	4E-32	PHYSICOMITRIN [Physicomitella patens]	Physicomitella patens	NO_HITS	16	28	704	430
153, 154	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	76	73	539	520
155, 156	gi 1502675 gb AAE443.1	0	alpha-amylase [Bacillus subtilis]	Bacillus subtilis	NO_HITS	97	98	591	659
157, 158	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	85	91	535	520
159, 160	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	96	93	526	520
161, 162	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	85	88	540	520
163, 164	gi 12855802 gb AAK00698.1	0	alpha-amylase [Bacillus megaterium]	Bacillus megaterium	NO_HITS	69	64	473	533
165, 166	gi 12855802 gb AAK00698.1	0	alpha-amylase [Bacillus megaterium]	Bacillus megaterium	NO_HITS	89	88	478	533
167, 168	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	86	82	531	520
169, 170	gi 11228329 emb CAG1.1	0.166	stereohydroxylase [Bacillus sp.]	Bacillus sp.	NO_HITS	60	59	508	515
171, 172	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	88	94	476	520
173, 174	gi 113814 sp P20846A.1	0	ALPHA-AMYLASE [Bacillus subtilis]	Bacillus subtilis	NO_HITS	88	83	531	520

FIGURE 35C

176, 178	gi115601018 NP_232848.1	0	alpha-amylase (Vibrio cholerae)	Vibrio cholerae Bacteria	NO HITS	73	68	465	468
177, 178	gi11228220 emb CAC16485.1	e-187	unnamed protein product (Geobacillus stearothermophilus)	Geobacillus stearothermophilus Bacteria	NO HITS	80	69	607	515
179, 180	gi17905890 p AAE91102.1	e-166	Sequence 8 from patent US 6287828	Unknown	NO HITS	69	N/A	507	614
181, 182	gi113760 p P22883A.MT4.PSES	0	GLUCAN 1,4-ALPHA-MALTOSE-6-DEHYDROLYASE PRECURSOR (G)	Pseudomonas saccharophila Bacteria	NO HITS	89	N/A	809	651
183, 184	gi113814 p P20845A.MT.BACNE	0	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN-GLUCANOHYDROLASE)	Bacillus megaterium Bacteria	NO HITS	95	92	631	520
185, 186	gi113814 p P20845A.MT.BACNE	0	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN-GLUCANOHYDROLASE)	Bacillus megaterium Bacteria	NO HITS	97	94	620	520
187, 188	gi15228763 NP_181616.1	3E-68	4-alpha-glucanotransferase (Arabidopsis thaliana)	Arabidopsis thaliana Eukarya	NO HITS	29	28	683	738
189, 190	gi113814 p P20845A.MT.BACNE	0	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN-GLUCANOHYDROLASE)	Bacillus megaterium Bacteria	NO HITS	95	92	631	520
191, 192	gi113814 p P20845A.MT.BACNE	0	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN-GLUCANOHYDROLASE)	Bacillus megaterium Bacteria	NO HITS	96	92	631	520
193, 194	gi13274586 p AAK17984.1 AF33076.1	0	alpha-amylase 4 (Pseudomonas sp. KFCC10818)	Pseudomonas sp. KFCC10818 Bacteria	NO HITS	77	66	853	765
195, 196	gi1482670 p JAS0999	0	3,2,1,1 precursor 4 (Micrococcus sp. 13720)	Micrococcus sp. Bacteria	3,2,1,1	59	66	929	1104
197, 198	gi115601018 NP_232848.1	0	alpha-amylase (Vibrio cholerae)	Vibrio cholerae Bacteria	NO HITS	72	69	486	486
199, 200	gi114029135 p JAX51132.1	1.9	orphan seven transmembrane receptor (Rattus norvegicus)	Rattus norvegicus Eukarya	NO HITS	27	44	132	730
201, 202	gi13182951 p JAAK15003.1 AF23372.1	1E-53	neopulvinase (Bacillus stearothermophilus)	Bacillus stearothermophilus Bacteria	NO HITS	12	40	630	688
203, 204	gi1728848 p P41131A.MYA.AERHY	e-148	ALPHA-D-GLUCAN-GLUCANOHYDROLASE	Aeromonas hydrophila Bacteria	NO HITS	32	35	856	443
205, 206	gi197759 p S10789	e-101	amylase A-180 - alkaliphilic eubacterium 183-26	alkaliphilic eubacterium 183-26 Bacteria	NO HITS	42	64	657	1694

FIGURE 35D

207, 208	g[116216810]refNP_174202.1	0.94	plasma fibronectin protein [Arabidopsis thaliana]	Arabidopsis thaliana Eukaryota	NO HITS	11	10	439	137
209, 210	g[477015]p[AA7874.a]	6-176	alpha-amylase [Xanthomonas campestris]	Xanthomonas campestris	NO HITS	61	66	472	475
211, 212	g[15222859]e[NP_177_210E-69]		alpha-amylase, putative [Arabidopsis thaliana]	Arabidopsis thaliana Eukaryota	NO HITS	42	30	498	413

FIGURE 35E

322, 323	alpha-amylase precursor [Thermococcus sp. GU5L5] hypothetical protein [Burkholderia fungorum]	Thermococcus sp. GU5L5	3.2.1.1	21326995	0	2251107	1392	483	461	83
324, 325		Burkholderia fungorum	3.2.1.1	22986674	1E-28	7379424	1296	431	1146	25
326, 327	alpha-amylase (EC 3.2.1.1) precursor - Bacillus megaterium hypothetical protein [Chloroflexus aurantiacus] hypothetical protein [Chloroflexus aurantiacus]	Bacillus megaterium	3.2.1.1	80110	7E-98	11344494	1359	452	520	42
328, 329		Chloroflexus aurantiacus	3.2.1.1	22970588	1E-155	535791	1677	558	575	53
330, 331		Chloroflexus aurantiacus	2.4.1.1	22971468	0	4633806	3129	1042	851	55
332, 333	[Microbulifer degradans 2-40] hypothetical protein [Microbulifer degradans 2-40]	Microbulifer degradans 2-40	3.2.1.1	23027235	1E-117	166984	1707	569	643	49
334, 335		Microbulifer degradans 2-40	3.2.1.1	23027235	7E-91	62191	2061	686	643	43

FIGURE 35F

336, 337	alpha-amylase (EC 3.2.1.1) precursor - Streptomyces violaceus	Streptomyces violaceus	3.2.1.1	80864	0	153156	1731	576	569	65
338, 339	unnamed protein, product	Bacillus sp.	3.2.1.1	13539158	0	14774986	1704	568	587	90
	outer membrane protein									
340, 341	[Bacteroides thetaiotaomicron] hypothetical protein	Bacteroides thetaiotaomicron	3.2.1.10	1478030	1E-137	153158	1848	615	692	46
342, 343	[Microbulbifer degradans 2-40] hypothetical protein	Microbulbifer degradans 2-40	3.2.1.1	23027631	0	13274585	2061	686	563	64
344, 345	[Microbulbifer degradans 2-40] hypothetical protein	Microbulbifer degradans 2-40	3.2.1.1	23027631	1E-179	8247214	1980	659	563	62
346, 347	amylase precursor alpha-amylase [Nostoc sp. PCC 7120]	Aeromonas hydrophila	3.2.1.1	141870	0	141869	1398	465	464	92
348, 349		Nostoc sp. PCC 7120	3.2.1.1	17229682	0	450848	1488	495	492	61
	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN GLUCANOHYDROLASE) hypothetical protein									
350, 351		Pseudoalteromonas haloplanktis	3.2.1.1	6226551	0	2467084	2001	666	669	67
352, 353	[Microbulbifer degradans 2-40] hypothetical protein	Microbulbifer degradans 2-40	3.2.1.1	23027235	1E-135	3549647	1263	420	643	55

FIGURE 35G

354, 355	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D- GLUCAN GLUCANOHYDRO LASE)	Aeromonas hydrophila	3.2.1.1	728848	0	304014	2577	858	443	44
356, 357	hypothetical protein [Microbulbifer degradans 2-40]									
358, 359	Glycosidase [Vibrio vulnificus CMCP6]	Microbulbifer degradans 2-40	3.2.1.1	23027235	7E-95	5442101	4875	1625	643	47
360, 361	amylase	Vibrio vulnificus	3.2.1.1	27366839	1E-147	155351	1422	473	466	57
	hypothetical protein [Microbulbifer degradans 2-40]	Bacillus thuringiensis	2.4.1.18	580662	0	2635411	1938	645	648	98
362, 363	Sequence 6 from patent US 5753460	Microbulbifer degradans 2-40	3.2.1.1	23027235	1E-65	13362592	2094	697	643	33
364, 365			3.2.1.1	3994289	0	722278	1536	511	549	69
366, 367	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D- GLUCAN GLUCANOHYDRO LASE)	Pseudoalteromonas as haloplanktis	3.2.1.1	6226551	0	2487084	1992	663	669	70
368, 369	hypothetical protein [Burkholderia fungorum]	Burkholderia fungorum	3.2.1.1	22986674	7E-28	14547281	1257	418	1146	27

FIGURE 35H

370, 371	alpha-amylase A [Halothermothrix orenil]	Halothermothrix orenil	24306106	1E-107	216309	1614	537	515	41
	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40							
372, 373			3.2.1.1	23027235	6E-77	166984	1437	478	42
374, 375	amylase precursor hypothetical protein [Microbulbifer degradans 2-40]	Aeromonas hydrophila	3.2.1.1	141870	0	141869	1398	465	91
376, 377		Microbulbifer degradans 2-40	3.2.1.1	23027235	6E-52	13702782	1551	516	32
	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40							
378, 379			3.2.1.1	23027235	1E-124	20334	1269	422	53
	periplasmic alpha- amylase precursor [Xanthomonas campestris]	Xanthomonas campestris	3.2.1.1	1166403	1E-140	1166402	1644	547	49
380, 381									
	putative bi- functional protein (secreted alpha- amylase/dextrinase)[Streptomyces coelicolor A3(2)]	Streptomyces coelicolor A3(2)	3.2.1.1	21220698	0	288182	4176	1391	50
382, 383									
	alpha-amylase [Xanthomonas campestris pv. campestris]	Xanthomonas campestris pv. campestris	3.2.1.1	19224331	0	155351	1434	477	65
384, 385									
	hypothetical protein [Chloroflexus aurantiacus]	Chloroflexus aurantiacus	3.2.1.1	22970588	1E-162	1771460	1458	485	59
386, 387									

FIGURE 35I

388, 389	GLUCAN 1,4-ALPHA- MALTOTETRAHY DROLASE PRECURSOR (G4 AMYLAZE) (MALTOTETRAOS E-FORMING AMYLASE) (EXO- MALTOTETRAOH YDROLASE) (MALTOTETRAOS E-FORMING EXO- AMYLASE)	Pseudomonas stutzeri	3.2.1.1	2506188	0	45821	1662	553	548	92
390, 391	alpha-amylase [Xanthomonas axonopodis pv. citri str. 306]	Xanthomonas axonopodis pv. citri str. 306	3.2.1.1	21106921	1E-177	155351	1497	498	475	61
392, 393	hypothetical protein [Nostoc punctiforme]	Nostoc punctiforme	3.2.1.1	23126762	8E-28	7799230	2100	699	552	29
394, 395	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40	3.2.1.1	23027235	1E-147	14023709	1347	448	643	60
396, 397	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40	3.2.1.1	23027244	1E-151	11433676	1644	547	566	48
398, 399	hypothetical protein [Nostoc punctiforme]	Nostoc punctiforme	2.4.1.18	23126762	5E-28	13276803	2040	679	552	32
400, 401	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40	3.2.1.1	23027235	1E-140	13507463	1245	414	643	57

FIGURE 35J

402, 403	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D- GLUCAN GLUCANOHYDRO- LASE)	Pseudoalteromonas as haloplanktis	3.2.1.1	6226551	0	2467084	1995	664	669	66
	hypothetical protein [Microbulbifer degradans 2-40]									
404, 405		Microbulbifer degradans 2-40	3.2.1.1	23027244	1E-155	2337886	1653	550	566	48
406, 407	amylase precursor alpha-amylase [Xanthomonas axonopodis pv. citri str. 306]	Aeromonas hydrophila	3.2.1.1	141870	0	141869	1398	465	464	94
408, 409		Xanthomonas axonopodis pv. citri str. 306	3.2.1.1	21106921	0	155351	1476	491	475	64
	hypothetical protein [Microbulbifer degradans 2-40]									
410, 411		Microbulbifer degradans 2-40	3.2.1.1	23027235	0	14861204	1875	624	643	59
	hypothetical protein [Chloroflexus aurantiacus]									
412, 413		Chloroflexus aurantiacus	3.2.1.1	22971473	0	8250619	2088	695	627	53
414, 415	hypothetical protein [Nostoc punctiforme] beta-agarase	Nostoc punctiforme	3.2.1.1	23126762	3E-38	14091925	2262	753	552	25
416, 417	[Pseudomonas sp. W7] alpha-amylase [Xanthomonas campestris pv. campestris]	Pseudomonas sp. W7	3.2.1.1	9081816	5E-31	14518450	1344	447	642	27
418, 419		Xanthomonas campestris pv. campestris	3.2.1.1	19224331	1E-169	155351	1455	484	475	60

FIGURE 35K

420, 421	hypothetical protein [Microbulbifer degradans 2-40]	Microbulbifer degradans 2-40	3.2.1.1	23027235	1E-149	18899	1308	435	643	57
422, 423	secreted alpha- amylase; [Streptomyces coelicolor A3(2)]	Streptomyces coelicolor A3(2)	3.2.1.1	21225304	4E-87	6855158	2751	916	993	14
	cyclomaltodextrin glucanotransferase (EC 2.4.1.19) precursor [validated] - Bacillus circulans (strain 8)									
424, 425	alpha-amylase, Alpha amylase, catalytic domain [Bacillus anthracis A2012]	Bacillus circulans	2.4.1.19	278549	0	39565	2169	722	718	80
426, 427	alpha-amylase [Xanthomonas campestris pv. campestris]	Bacillus anthracis str. A2012	3.2.1.41	21400626	1E-151	10728478	2139	712	724	42
428, 429	alpha-amylase [Bacillus sp. TS-23]	Xanthomonas campestris pv. campestris	3.2.1.1	19224331	1E-142	9789644	1425	474	475	52
430, 431	alpha-amylase (EC 3.2.1.1) precursor - Streptomyces violaceus	Bacillus sp. TS- 23	3.2.1.1	722279	0	722278	1650	550	613	85
432, 433		Streptomyces violaceus	3.2.1.1	80864	0	7619766	1752	583	569	64

FIGURE 35L

434, 435	alpha-amylase (EC 3.2.1.1) Isozyme III	-	rice	Oryza sativa	3.2.1.1	11263719	1E-110	3769329	1302	433	437	48
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FIGURE 35M

438, 439	hypotheical protein [Chloroflexus aurantiacus]	22970478	1E-166	Chloroflexus aurantiacus	Thermus flavus amyloamylase encoding DNA	AAW83330	1E-157	Thermus rubens glucanotransferase gene amplifying Xba PCR primer	AAD04867	4E-04	5	2.4.1.2	1539	512	1806	601	54	58
440, 441	Crystal Structure Of Amyk38 N289h Mutant	34811325	6E-19		Alpha-amylase K38AMY mutagenic PCR primer Q209	AAO21008	8E-20	Shewanella sp. SCRC-21406 (FERM BP-5979) ORFb DNA SEQ ID NO:4	AAA92470	0.87	3.2.1.1	3447	1148	7875	480	13		
442, 443	GLUCAN 1,4-ALPHA-MALTOTETRAHYDROLASE PRECURSOR, MALTOTETRAOSE-FORMING AMYLASE, MALTOTETRAOSE-FORMING EXO-AMYLASE	2506188	0	Pseudomonas	Maltotetraose	AAR07282	0	Maltotetraose	AAQ05095	0	3.2.1.1	1668	555		548	75	78	

FIGURE 35N

444, 445	alpha-amylase Thermoactinomyces vulgaris	322327	1E-107	Thermoactinomyces vulgaris	Alpha amylase DNA PCR primer #1.	ABU030 92	1E-121	Human cDNA encoding secreted/transmembrane protein ABX16 #76.	0.38	2.4.1.1 9	1521	506	2196	502	44	43
446, 447	alpha-amylase Thermoactinomyces vulgaris	322327	5E-74	Thermoactinomyces vulgaris	Alpha amylase DNA PCR primer #1.	ABU030 92	3E-56	NOVX related reverse PCR primer SEQ ID No 149.	0.89	2.4.1.1 8	3537	1178		482	14	35
448, 449	AmyM [uncultured bacterium]	37183425	1E-158	uncultured bacterium	Alpha amylase DNA PCR primer #1.	ABU031 33	1E-126	Human cDNA SEQ ID NO 201.	0.099	2.4.1.2 5	1575	524	1554	517	51	57
450, 451	alpha-amylase 3 [Bacteroides thetaotaomicro n VPI-5482]	29346183	1E-151	Bacteroides thetaotaomicro n VPI-5482	Alpha amylase DNA PCR primer #1.	ABU031 34	4E-16	Human cDNA encoding human transporter polypeptide AAS16 905.	0.47		1890	629		565	43	49
452, 453	AmyA [uncultured bacterium]	37222142	1E-135	uncultured bacterium	Alpha amylase DNA PCR primer #1.	ABU031 31	1E-72	Alpha amylase DNA PCR primer #1.	0.007	3.2.1.1	1734	577		608	43	56

FIGURE 350

454. 455	alpha-amylase [Hordeum vulgare]	295804	1E-116	Hordeum vulgare	Endoxylol glucan transferase sequence #165	AAM001 08	1E-118	Partial sequence of tomato Ca2+ ATPase.	AAZ29 771	0.33	3.2.1.1	1341	446	6263	430	47	53
456. 457	AmyA [uncultured bacterium]	37222142	1E-136	uncultured bacterium	Alpha amylase DNA PCR primer #1.	ABU031 31	2E-73	Alpha amylase DNA PCR primer #1.	ABX08 502	0.028	3.2.1.1	1734	577		608	44	56
458. 459	ALPHA- AMYLASE PRECURSOR (1,4-ALPHA-D- GLUCAN GLUCANOHYD ROLASE).	113814	3E-81	Bacillus megaterium	Alpha amylase DNA PCR primer #1.	ABU031 15	7E-85	Alpha amylase DNA PCR primer #1.	ABX08 486	0.002	3.2.1.1	1698	565	1620	539	30	47
460. 461	maltase [Aspergillus oryzae]	14278921	1E-133	Aspergillus oryzae	Vibrio harveyi endoglucanase DNA.	AAW34 990	1E-108	Chromosome 13q31- q33 biallelic marker containing amplicon SEQ ID #182.	AAH51 601	0.11	3.2.1.2	1752	583	1725	574	43	53

FIGURE 35P

462-466	related to glucosylase precursor [Neurospora crassa]	38524238	1E-143	Neurospora crassa	Thielavia terrestris glucosylase DNA PCR primer #4	AAM515	96	2E-39	NO 3150	Aspergillus oryzae polynucleotide	ABZ54	259	0.001	1206	401	405	60	63
467-474	hypothetical protein MG03287.4 [Magnaporthe grisea 70-15]	38105244	1E-177	Magnaporthe grisea 70-15	Sequence of amylose gene and upstream regulator YDNA	AAR093	59	1E-125	#1	Alpha amylase DNA PCR primer	ABX08	477	0.61	3.2.1.1	806	518	36	39
475-479	putative alpha-1,3-glucan synthase [Aspergillus fumigatus]	16418019	0	Aspergillus fumigatus	Maize Starch synthase IIb (SSIIb) LINKR domain related protein #42	ABU065	20	3E-13	NO 3150	Aspergillus oryzae polynucleotide	ABZ53	554	5E-04	2.4.1.1	2376	7281	59	59
480-485	hypothetical protein MG10209.4 [Magnaporthe grisea 70-15]	38101134	1E-175	Magnaporthe grisea 70-15	Amino acid sequence of a fungamyl-like alpha-amylase	AAB842	08	1E-103	NO:1176	Fusarium venenatum EST	AAF07	664	5E-65	3.2.1.1	460	600	62	62

FIGURE 35Q

486-493	GLUCOAMYLASE PRECURSOR (GLUCAN 1,4-ALPHA-GLUCOSIDASE) (1,4-ALPHA-D GLUCAN GLUCOHYDROLASE).	461509	Amorpha hotheca resinosa	cDNA encoding glucanase P.	AAW30155	0	cDNA encoding glucanase P.	AAT90830	5E-04	3.2.1.3	1932	643	616	61	60
494-499	alpha-amylase AmyA [Emeticella nidulans].	6561867	Emeticella nidulans	Mutant alpha-amylase.	AAR46065	1E-142	Fusarium venenatum EST	AAF12832	4E-04	3.2.1.1	1479	492	1473	490	49
500-510	alpha-glucosidase [Schizosaccharomyces pombe].	19111855	Schizosaccharomyces pombe	PCR primer B1 from J092340	AAW27300	1E-123	Drosophila melanogaster polypeptide of SEQ ID NO 24465.	ABL18081	0.002	3.2.1.2	1833	610	1740	579	42
511-516	acid-stable alpha-amylase [Aspergillus kawachii].	2570150	Aspergillus kawachii	Amino acid sequence of a fungal alpha-amylase.	AAB84206	1E-122	Arabidopsis thaliana stress regulated gene	ABZ16178	1.6	3.2.1.1	1650	549	1923	640	43

FIGURE 35R

517-518	alpha-amylase [AmyA [<i>Emericella nidulans</i>]	6561867	1E-110	Emeric ella nidulan s	Alpha- amylase variant with leucine at position 84.	AAR241 36	1E-110	S. pneumon iae type 4 strain protein from coding region #1864.	ABS56 454	0.44	3.2.1.1	1758	585	1473	490	37	42
519-523	alpha- glucosidase [<i>Aspergillus oryzae</i>]	23503475	1E-179	Asperg illus oryzae	PCR primer B1 from J092340	AAW27 300	1E-129	Fusarium venenatu m EST SEQ ID NO:1176.	AAF08 465	0.002	3.2.1.2	1788	595	1809	602	50	56
524-528	OLIGO-1,6- GLUCOSIDASE (SUCRASE- ISOMALTASE) (LIMIT DEXTRINASE) (ISOMALTASE) (DEXTRIN 6- ALPHA-D- GLUCANOHYD ROLASE)	129007	1E-145	Geoba cillus thermo glucosi gasius	PCR primer B1 from J092340	AAW27 300	1E-138	Bacillus lichenfor mis genomic sequence tag (GST) #933.	ABK73 353	0.007	3.2.1.1	1755	584		562	46	51
529-533	alpha- glucosidase [<i>Aspergillus oryzae</i>]	23503475	1E-165	Asperg illus oryzae	PCR primer B1 from J092340	AAW27 300	1E-141	Aspergillu s oryzae polynucle otide SEQ ID NO 3150.	ABZ54 241	5E-07	3.2.1.2	1797	598	1809	602	48	55

FIGURE 35S

534-540	hypothetical protein MG10209.4 [Magnaporthe grisea 70-15]	38101134	0	Magnaporthe grisea 70-15	Amino acid sequence of a fungamyl-like alpha-amylase	AAB84206	1E-105	Fusarium venenatum EST SEQ ID NO:1176	AAF07684	5E-13	3.2.1.1	1770	589	600	65	68
541-545	hypothetical protein [Neurospora crassa]	32411795	0	Neurospora crassa	HIV multifunctional fusion polypeptide	AAR13230	1E-137	HIV multifunctional fusion polypeptide	AAQ12770	1E-04	3.2.1.2	1992	663	608	60	
546-553	glucan 1,4-alpha-glucosidase (EC 3.2.1.3) precursor - Neurospora crassa	486943	0	Neurospora crassa	Thielavia terrestris: glucanase DNA PCR primer #4	AAM51596	.0	Thielavia terrestris: glucanase DNA PCR primer #4	ABA01139	5E-07	3.2.1.3	2019	672	626	58	36
554-559	maltase [Aspergillus oryzae]	14278921	0	Aspergillus oryzae	HIV multifunctional fusion polypeptide	AAR13230	1E-147	Fusarium venenatum EST SEQ ID NO:1176	AAF13291	5E-04	3.2.1.2	1797	598	574	57	59

FIGURE 35T

550-566	ALPHA-AMYLASE A PRECURSOR (1,4-ALPHA-D-GLUCAN GLUCANOHYDROLASE A)	1703298	1E-125	Aspergillus niger var. awamori	Amino acid sequence of alpha-amylase	AAB84206	1E-125	Vector: pPR70-4 xlnB expression element AAV61 459	0.38	3.2.1.1	1524	507	498	45
567-568	beta-agarase [Pseudomonas sp. W7]	9081816	1E-126	Pseudomonas sp. W7	Alpha amylase DNA PCR primer #1	ABU03140	1E-113	Human secreted protein SEQ ID NO 792	1.4	3.2.1.1	1446	481	1926	49
569-570		40189607	0								1095	364	1095	100
571-572	alpha-amylase [Bacillus sp. TS-23]	722279	0	Bacillus sp. TS-23	Alpha amylase DNA PCR primer #1	ABU03102	0	Alpha amylase DNA PCR primer #1	3E-11	3.2.1.1	1839	612	1842	59
573-574	CYCLOMALTO DEXTRIN GLUCANOTRANSFERASE PRECURSOR (CYCLODEXTRIN-GLYCOSYLTRANSFERASE) (CGTASE)	1351937	0	Thermooanaerobacterium thermosulfurigenes	Thermooanaerobacterium thermosulfurigenes CGTase variant G180S	AAW06772	0	Thermooanaerobacterium thermosulfurigenes CGTase variant G180S	2.4.1.1	2.4.1.1	2133	710	710	100

FIGURE 35U

575, [Bacillus sp. TS-23], 576	722279	7E-77	Bacillus sp. TS-23	Nucleotide sequence of alpha-amylase of Bacillus NCIMB 40916	AAB676	55	0	0	3.2.1.1	1677	558	1764	587	99	97
577, cyclodextrin glucanotransferase [Bacillus circulans], 578	11139208	0	Bacillus circulans	Plasmid pTN603 encoding novel alpha-amylase gene	AAV025	99	0	0	2.4.1.1	2160	719	2142	713	67	66
579, AMYLOPULLULANASE PRECURSOR (ALPHA-AMYLASE/PULLULANASE) 580	114076	0	Thermobacter hydrophilus	Alpha amylase pullulanae gene	AAR082	21	0	0	3.2.1.1	5010	1669		1475	87	90

FIGURE 35V

581, 582	CYCLOMALTO DEXTRIN GLUCANOTRANSFERASE PRECURSOR (CYCLODEXTRIN- IN- GLYCOSYLTRANSFERASE) (CGTASE)	399222	0	Bacillus sp. 17-1	Cyclomaltodextrin glucotransferase 17-1 AAR100 gene.	52	0	AAQ01810	2.4.1.1.9	2142	713	713	95	89
583, 584	amylolipullanase [Geobacillus stearothermophilus]	12006232	0	Geobacillus stearothermophilus	Alpha amylase pullulanase gene.	AAR08221	0	AA199683	3.2.1.2	6618	2205	6057	2018	45
585, 586	alpha-amylase [Bacillus sp. TS-23]	722279	0	Bacillus sp. TS-23	Alpha amylase DNA PCR primer #1.	ABU03091	0	ABX08462	0 3.2.1.1	1860	619	1842	613	75
603, 604	alpha-amylase [Bacillus sp. TS-23]	722279	1E-77	Bacillus sp. TS-23	Nucleotide sequence of an alpha-amylase of Bacillus NCIMB 40916.	AAB67655	0	AAF55662	0 3.2.1.1	1773	590	1764	587	95

FIGURE 35W

605, 606	CYCLOMALTOSE DEXTRIN GLUCANOTRANSFERASE PRECURSOR (CYCLODEXTRIN- IN- GLYCOSYLTRANSFERASE) (CGTASE)	399219	0	Bacillus circulans	AA025 variant	99	0	Plasmid pTN603 encoding novel amylase gene.	AAN60 705	2.4.1.1 9	2160	719	6057	2018	718	74	70
607, 608	amylolipulanase [Geobacillus stearothermophilus]	12006232	0	Geobacillus stearothermophilus	ABU030 primer #1	83	0	Alpha amylase DNA PCR primer #1.	ABX08 454	3.2.1.1 35	3708	1235	6057	2018	2018	77	71
609, 610	alpha-amylase [Bacillus sp. TS-23]	722279	7E-77	Bacillus sp. TS-23	NCIMB 40916	55	0	Nucleotide sequence of an alpha- amylase of Bacillus NCIMB 40916.	AAF55 662	0 3.2.1.1	1764	587	1764	587	587	98	97
611, 612	hypothetical protein - Bacillus sp.	2126830	0	Bacillus sp.	ABU030 primer #1.	83	0	Alpha amylase DNA PCR primer #1.	ABX08 454	3.2.1.1 35	3807	1268		2032	2032	75	

FIGURE 35X

613, 614	alpha-amylase [Bacillus sp. TS-23]	722279	3E-77	Bacillus sp. TS-23	Nucleotide sequence of an alpha-amylase of Bacillus NCIMB 40916	AAB676	55	0	Nucleotide sequence of an alpha-amylase of Bacillus NCIMB 40916	AAF55	682	0	3.2.1.1	1764	587	1764	587	99	97
615, 616	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN GLUCANOHYDROLASE)	113822	0	Streptomyces violaceus	Amylase from Streptomyces griseus IMRU 3570	AAR082	63	0	Amylase from Streptomyces griseus IMRU 3570	AAN80	309	2E-61	3.2.1.1	1752	583		569	64	72
617, 618	GLUCAN 1,4-ALPHA-MALTOTETRAHYDROLASE PRECURSOR (G4-AMYLASE) (MALTOTETRAOSE-FORMING AMYLASE) (EXO-MALTOTETRAHYDROLASE) (MALTOTETRAOSE-FORMING EXO-AMYLASE)	113760	0	Pseudomonas saccharophila	Maltotetraose	AAR072	82	0	Maltotetraose	AAQ06	095	0	3.2.1.1	1668	555		551	74	78

FIGURE 35Y

619, 620	ALPHA-AMYLASE PRECURSOR (1,4-ALPHA-D-GLUCAN GLUCANOHYDROLASE).	113822	0	Streptomyces violaceus	Amylase from Streptomyces griseus IMRU 3570.	AAR082 63	0	Amylase from Streptomyces griseus IMRU 3570.	AAQ06 844	0	3.2.1.1	1716	571	569	91	81
621, 622	alpha-amylase [Bacillus sp. TS-23].	722279	3E-77	Bacillus sp. TS-23	Nucleotide sequence of an alpha-amylase of Bacillus NCIMB 40916.	AAB676 55	0	Nucleotide sequence of an alpha-amylase of Bacillus NCIMB 40916.	AAF55 662	0	3.2.1.1	1773	590	1764	587	98
																97

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gtcggggagt	actgggacac	aaacgttgat	gcactgctca	actgggccta	ctcgagcgat	720
gcaaaaagtct	tcgacttccc	gctctactac	aagatggacg	cggcctttga	caacaagaac	780
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aaggccgtaa	ccttcgttgc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga	1020
agcactgaca	tcgtttacta	cgacaacgac	gagctgatat	tcgtgagaaa	cggctacgga	1080
agcaagccgg	gactgataac	atacatcaac	ctcgcctcaa	gcaaagccgg	aagggtgggtt	1140
tacgttccga	agttcgcagg	ctcgtgcata	cacgagtaca	ccggcaatct	cggcggctgg	1200
gtggacaagt	gggtggactc	aagcggctgg	gtctacctcg	aggctcctgc	ccacgaccgc	1260
gccaacggcc	agtacggcta	ctccgtctgg	agctactgcg	gtgttggttg	a	1311

<210> 2

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

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<400> 2
Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Leu Ile Met Gln Ala
1 10 15
Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala
20 25 30
Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
35 40 45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
50 55 60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
65 70 75 80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
85 90 95
Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
100 105 110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
115 120 125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130 135 140
Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
145 150 155 160
Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
165 170 175
Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180 185 190
Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
195 200 205
Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210 215 220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
225 230 235 240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala Ala Phe
245 250 255
Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly
260 265 270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr phe Val Ala Asn
275 280 285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290 295 300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305 310 315 320
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
325 330 335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
340 345 350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355 360 365
Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
370 375 380
Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385 390 395 400
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405 410 415
Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420 425 430
Cys Gly Val Gly
435

```

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<210> 3
<211> 1311
<212> DNA
<213> Artificial Sequence

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<220>
<223> synthetically generated oligonucleotide

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<400> 3
atggccaagt acctggagct cgaagagggc gggctcataa tgcaggcctt ctactgggac

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60

1001827087_1.txt

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gtcccatgg gaggaatctg gtgggacacg atagcccaga agatacccga ctgggcaagc 120
gccgggattt cggcgatatg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctac 300
ggcatcaagg tcatcgcaga catagtaatc aaccaccgcg ccggaggaga ccttgagtgg 360
aacccttcg tcaatgacta cacctggacg gactttctcg aggtcgcttc cggcaagtac 420
acggccaatt acctcgactt ccacccgaac gagctccatg cgggcgattc cggcaattt 480
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caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac 600
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gttgagagt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggt 720
gccaaggctt ttgacttcgc cctctactac aagatggatg aggcctttga caacaaaaac 780
attccagcgc tcgtctctgc ccttcagaac ggccagactg ttgtctcccg cgacccttc 840
aaggccgtaa cctttgtagc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
tacgcgttca tcctcaccta cgagggccag ccgacaatat tctaccgcga ctacgaggag 960
tggtcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
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tacgttccga agttcgcggg agcgtgcatc cagagtaca ccggcaacct cggcggctgg 1200
gtggacaagt ggggtggactc aagcgggtgg gtgtacctcg aggccctgc ccacgaccg 1260
gccaacggct attacggcta ctccgtctgag agctattgcg gtgttgggtg a 1311

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<210> 4

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 4

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Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Leu Ile Met Gln Ala
 1          5          10          15
Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala
          20          25          30
Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
 35          40          45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50          55          60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65          70          75          80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Val Asn Met Ile Asn Thr
 85          90          95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100          105          110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
115          120          125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130          135          140
Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
145          150          155          160
Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
165          170          175
Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180          185          190
Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
195          200          205
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210          215          220
Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
225          230          235          240
Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
245          250          255
Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
260          265          270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275          280          285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile

```

1001827087_1.txt

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      290      295      300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305      310      315      320
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
      325      330      335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
      340      345      350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
      355      360      365
Ile Asn Leu Ala Ser Ser Glu Ala Gly Arg Trp Val Tyr Val Pro Lys
      370      375      380
Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385      390      395      400
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
      405      410      415
Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr
      420      425      430
Cys Gly Val Gly
      435

```

<210> 5
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

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<400> 5
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gtcccagggtg gaggaatctg gtgggacacc atcaggagca agataccgga gtggtacgag      120
gcgggaatat ccgccatttg gattcccccg gcaagcaagg gcatgggcgg cgcctattcg      180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta      240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctat      300
ggcatgaagg taatagccga tatagtcatc aaccaccgcg ccggcgggtga cctggagtgg      360
aacccttctg tgaacgacta tacctggacc gacttctcaa aggtcgcgtc gggtaaatac      420
acggccaact acctcgactt ccaccgaaac gagctccatg cgggcgattc cgggaacattt      480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggccagc      540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac      600
gtcaagggct atgtccctg ggtcgtcaag gactggctga actggtgggg aggctgggcg      660
gttggagagt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggt      720
gccaaggtct ttgacttcgc cctctactac aagatggatg aggcctttga caacaaaaac      780
attccagcgc tcgtctctgc ctttcagaac ggccagactg ttgtctcccg cgacccttc      840
aaggccgtaa ctttgtagc aaaccacgac accgatataa tctggaacaa gtaccttgct      900
tatgctttca tcctcaccta cgaaggccag cccgtcatat tctaccgca ccacgaggag      960
tggctcaaca aggacaggtt gaacaacctc atatggatac acgaccacct cgcagggtgga      1020
agcaccgaca tagtctacta cgataacgat gaactcatct tcgtcaggaa cggctacggg      1080
gacaagccgg ggcttataac ctacatcaac ctaggctcga gcaaggccgg aagggtgggtt      1140
tatgtgccga agttcgcggg cgcggtgcatc cagagtataa ctggtaacct cggaggctgg      1200
gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgacct      1260
gccaacgggc agtatggcta ctccgtgtgg agctactgcg ggggtgggctg a      1311

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<210> 6
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

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<400> 6
Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala
 1      5      10      15
Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg
      20      25      30
Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
      35      40      45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp

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50	Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	Glu	Tyr	Asp	Gln	Lys	Gly	Thr	Val
65	Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr
	Ala	His	Ala	Tyr	Gly	Met	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
	Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr	Thr
	Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
	Leu	Asp	Phe	His	Pro	Asn	Glu	Leu	His	Ala	Gly	Asp	Ser	Gly	Thr	Phe
	Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	Asp	Lys	Ser	Trp	Asp	Gln	Tyr	Trp
	Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
	Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Ala	Pro	Trp	Val
	Val	Lys	Asp	Trp	Leu	Asn	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	
	Trp	Asp	Thr	Asn	Val	Asp	Ala	Val	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Gly
	Ala	Lys	Val	Phe	Asp	Phe	Ala	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
	Asp	Asn	Lys	Asn	Ile	Pro	Ala	Leu	Val	Ser	Ala	Leu	Gln	Asn	Gly	Gln
	Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
	His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Leu	Ala	Tyr	Ala	Phe	Ile
	Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Val	Ile	Phe	Tyr	Arg	Asp	His	Glu	Glu
	Trp	Leu	Asn	Lys	Asp	Arg	Leu	Asn	Asn	Leu	Ile	Trp	Ile	His	Asp	His
	Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu
	Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	Asp	Lys	Pro	Gly	Leu	Ile	Thr	Tyr
	Ile	Asn	Leu	Gly	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys
	Phe	Ala	Gly	Ala	Cys	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp
	Val	Asp	Lys	Tyr	Val	Tyr	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro
	Ala	Tyr	Asp	Pro	Ala	Asn	Gly	Gln	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr
	Cys	Gly	Val	Gly												

<210> 7
 <211> 16
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 7
 Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp
 1 5 10 15

<210> 8
 <211> 26
 <212> PRT
 <213> Bacterial

<400> 8

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Met Lys Pro Ala Lys Leu Leu Val Phe Val Leu Val Val Ser Ile Leu
 1 5 10 15
 Ala Gly Leu Tyr Ala Gln Pro Ala Gly Ala
 20 25

<210> 9
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> synthetically generated oligonucleotide

<400> 9
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 gtcccatg gaggaatctg gtgggacacg atagcccaga agatacccga ctgggcaagc 120
 gccgggattt cggcgatatg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaaacggta 240
 gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
 aaccggttcg ttggggacta cacctggacg gactttctca aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
 ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggccagc 540
 caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac 600
 gtcaagggct atgctccctg ggtcgtcaag gactggctga actggtgggg aggctgggcg 660
 gttggagagt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggg 720
 gccaaaggct ttgacttcgc cctctactac aagatggacg aggccttcga taacaacaac 780
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 tacgttccga agttcgcagg ctcgtgcata cagagtaca ccggcaatct cggcggctgg 1200
 gtggacaagt ggggtggactc aagcggctgg gtctacctg aggcctctgc ccacgaccgc 1260
 gccaacggcc agtacggcta ctccgtctgg agctactgcg gtgttgggtg a 1311

<210> 10
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> synthetically generated peptide

<400> 10
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 1 5 10 15
 Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala
 20 25 30
 Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175

1001827087_1.txt

Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Ala	Pro	Trp	Val
Val	Lys	Asp	Trp	Leu	Asn	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr
Trp	Asp	Thr	Asn	Val	Asp	Ala	Val	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Gly
Ala	Lys	Val	Phe	Asp	Phe	Ala	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
Asp	Asn	Asn	Asn	Ile	Pro	Ala	Leu	Val	Asp	Ala	Leu	Arg	Tyr	Gly	Gln
Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	Asn
Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu
Ile	Phe	Ala	Arg	Asn	Gly	Tyr	Gly	Ser	Lys	Pro	Gly	Leu	Ile	Thr	Tyr
Ile	Asn	Leu	Ala	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys
Phe	Ala	Gly	Ser	Cys	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp
Val	Asp	Lys	Trp	Val	Asp	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro
Ala	His	Asp	Pro	Ala	Asn	Gly	Gln	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr
Cys	Gly	Val	Gly												

<210> 11
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 11	
atggccaagt acctggagct cgaggagggc gggctcataa tgcaggcctt ctactgggac	60
gtcccatg gaggaatctg gtgggacacg atagcccaga agatacccga ctgggcaagc	120
gccgggattt cggcgatatg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg	180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta	240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctat	300
ggcatgaagg taatagccga tatagtcatc aaccaccgcg ccggcggtga cctggagtgg	360
aacccttcg tgaacgacta tacctggacc gacttctcaa aggtcgcgtc gggtaaatac	420
acggccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt	480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggccagc	540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac	600
gtcaagggct atgctccctg ggtcgtcaag gactggctga actgggtggg aggctgggcg	660
gttggaaggt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggg	720
gccaaagtct ttgacttcgc cctctactac aagatggacg aggccttcga taacaacaac	780
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agcaagcctg gccttataac ttacatcaac ctcggctcga gcaaggttgg aagggtgggtc	1140
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gtggacaagt ggggtggactc aagcgggtgg gtgtacctcg aggccctgc ccacgacctg	1260
gccaacggct attacggcta ctccgtctgg agctactgcg gtgttggtcg a	1311

<210> 12

<211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 12
 Met Ala Lys Tyr Leu Glu Leu Glu Glu Gly Gly Leu Ile Met Gln Ala
 1 5 10 15
 Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala
 20 25 30
 Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Thr Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 13
 <211> 1311
 <212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 13

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gccggaatct	ccgcaatatg	gattcccccg	gcgagcaagg	gcatgggdcg	cgcctattcg	180
atgggctacg	acccctacga	cttctttgac	ctcggtagtg	atgaccagaa	gggaacggta	240
gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacacggc	acatgcctac	300
ggcataaagg	tcatagcgga	catcgtcata	aaccaccgcg	caggcggaga	cctcgaagtgg	360
aacccgttcg	ttggggacta	cacctggacg	gactttctcaa	aggtggcctc	gggcaaatat	420
actgccaact	acctcgactt	ccaccccaac	gaggtcaagt	gctgtgacga	gggcacattt	480
ggaggcttcc	cagacatagc	ccacgagaag	agctgggacc	agcactggct	ctgggcgagc	540
gatgagagct	acgccgccta	cctaaggagc	atcggcggtg	atgcctggcg	cttcgactac	600
gtcaagggct	acggagcgtg	ggtcgtcaag	gactggctgg	actggtgggg	aggctgggcc	660
gtcggggagt	actgggacac	aaacgttgat	gcactgctca	actgggccta	ctcgaagcat	720
gcaaaagtct	tcgacttccc	gctctactac	aagatggatg	aggcctttga	caacaaaaac	780
attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgacccttc	840
aagggccgtaa	cccttgtagc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggctcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga	1020
agcactgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg	1080
gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aaggtggggt	1140
tatgtgcga	agttcgcggg	cgcgtgcctc	cacgagtata	ctggtaacct	cggaggctgg	1200
gtagacaagt	acgtctactc	aagcggctgg	gtctatctcg	aagctccagc	ttacgaccct	1260
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<210> 14

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 14

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Phe	Tyr	Trp	Asp	Val	Pro	Ser	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg
			20					25					30		
Gln	Lys	Ile	Pro	Glu	Trp	Tyr	Asp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile
		35					40					45			
Pro	Pro	Ala	Ser	Lys	Gly	Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr	Asp
	50					55					60				
Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	Glu	Tyr	Asp	Gln	Lys	Gly	Thr	Val
65					70					75				80	
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr
			85					90					95		
Ala	His	Ala	Tyr	Gly	Ile	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
			100					105					110		
Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Gly	Asp	Tyr	Thr
		115					120					125			
Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
	130					135					140				
Leu	Asp	Phe	His	Pro	Asn	Glu	Val	Lys	Cys	Cys	Asp	Glu	Gly	Thr	Phe
145					150					155					160
Gly	Gly	Phe	Pro	Asp	Ile	Ala	His	Glu	Lys	Ser	Trp	Asp	Gln	His	Trp
			165					170						175	
Leu	Trp	Ala	Ser	Asp	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
			180					185					190		
Val	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val
		195					200					205			
Val	Lys	Asp	Trp	Leu	Asp	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr
	210					215					220				
Trp	Asp	Thr	Asn	Val	Asp	Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Asp
225					230				235						240

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Ala	Lys	Val	Phe	Asp	Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
				245					250					255	
Asp	Asn	Lys	Asn	Ile	Pro	Ala	Leu	Val	Ser	Ala	Leu	Gln	Asn	Gly	Gln
			260					265					270		
Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
		275					280					285			
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
	290					295					300				
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
	305				310					315					320
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	Asn
				325					330					335	
Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu
			340					345					350		
Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	Asp	Lys	Pro	Gly	Leu	Ile	Thr	Tyr
		355					360					365			
Ile	Asn	Leu	Gly	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys
	370					375					380				
Phe	Ala	Gly	Ala	Cys	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp
	385				390					395					400
Val	Asp	Lys	Tyr	Val	Tyr	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro
				405					410					415	
Ala	Tyr	Asp	Pro	Ala	Asn	Gly	Gln	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr
			420					425					430		
Cys	Gly	Val	Gly												
			435												

<210> 15
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 15	
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gccgggattt cggcgatatg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg	180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta	240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac	300
ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg	360
aaccggttcg ttggggacta cacctggacg gactttctca aggtggcctc gggcaaatat	420
actgccaact acctcgactt ccaccggaac gagctccatg cgggcgattc cggaacattt	480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctggggccagc	540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac	600
gtcaagggct acggagcgtg ggtcgtcaag gactggctgg actggtgggg aggcctggcc	660
gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat	720
gcaaaagtct tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac	780
attccagcgc tcgtctctgc ctttcagaac ggccagactg ttgtctcccg cgaccggttc	840
aaggccgtaa cttttgtagc aaaccacgac accgatataa tttggaacaa gtaccgggcc	900
tacgccttca tcctcaccta cgaggggccag ccgacgatat tctaccgca ctacgaggag	960
tggctcaaca aggacaggct caagaacctc atctggatac acgaccacct tgccggtgga	1020
agcactgaca tcgtttacta cgacaacgac gagctgatat tcgtgagaaa cggctacgga	1080
agcaagccgg gactgataac atacatcaac ctcgcctcaa gcaaagccgg aaggctgggtt	1140
tatgtgccga agttcgcggg cgctgcatc cagcagatata ctggtaacct cgaggctgg	1200
gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgacctt	1260
gccaacgggc agtatggcta ctccgtgtgg agctattgca gtgttggtg a	1311

<210> 16
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 16

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Phe	Tyr	Trp	Asp 20	Val	Pro	Met	Gly	Gly 25	Ile	Trp	Trp	Asp	Thr 30	Ile	Ala
Gln	Lys	Ile 35	Pro	Asp	Trp	Ala	Ser 40	Ala	Gly	Ile	Ser	Ala 45	Ile	Trp	Ile
Pro	Pro 50	Ala	Ser	Lys	Gly	Met 55	Gly	Gly	Ala	Tyr	Ser 60	Met	Gly	Tyr	Asp
Pro 65	Tyr	Asp	Phe	Phe	Asp 70	Leu	Gly	Glu	Tyr	Asp 75	Gln	Lys	Gly	Thr	Val 80
Glu	Thr	Arg	Phe	Gly 85	Ser	Lys	Gln	Glu	Leu 90	Val	Asn	Met	Ile	Asn 95	Thr
Ala	His	Ala	Tyr 100	Gly	Ile	Lys	Val	Ile 105	Ala	Asp	Ile	Val	Ile 110	Asn	His
Arg	Ala	Gly 115	Gly	Asp	Leu	Glu	Trp 120	Asn	Pro	Phe	Val	Gly 125	Asp	Tyr	Thr
Trp	Thr 130	Asp	Phe	Ser	Lys	Val 135	Ala	Ser	Gly	Lys	Tyr 140	Thr	Ala	Asn	Tyr
Leu 145	Asp	Phe	His	Pro	Asn 150	Glu	Leu	His	Ala	Gly 155	Asp	Ser	Gly	Thr	Phe 160
Gly	Gly	Tyr	Pro	Asp 165	Ile	Cys	His	Asp	Lys 170	Ser	Trp	Asp	Gln	Tyr 175	Trp
Leu	Trp	Ala	Ser 180	Gln	Glu	Ser	Tyr	Ala 185	Ala	Tyr	Leu	Arg	Ser 190	Ile	Gly
Ile	Asp	Ala 195	Trp	Arg	Phe	Asp	Tyr 200	Val	Lys	Gly	Tyr	Gly 205	Ala	Trp	Val
Val	Lys 210	Asp	Trp	Leu	Asp	Trp 215	Trp	Gly	Gly	Trp	Ala 220	Val	Gly	Glu	Tyr
Trp 225	Asp	Thr	Asn	Val	Asp 230	Ala	Leu	Leu	Asn	Trp 235	Ala	Tyr	Ser	Ser	Asp 240
Ala	Lys	Val	Phe	Asp 245	Phe	Pro	Leu	Tyr	Tyr 250	Lys	Met	Asp	Glu	Ala 255	Phe
Asp	Asn	Lys	Asn 260	Ile	Pro	Ala	Leu	Val 265	Ser	Ala	Leu	Gln	Asn 270	Gly	Gln
Thr	Val	Val 275	Ser	Arg	Asp	Pro	Phe 280	Lys	Ala	Val	Thr	Phe 285	Val	Ala	Asn
His 290	Asp	Thr	Asp	Ile	Ile	Trp 295	Asn	Lys	Tyr	Pro	Ala 300	Tyr	Ala	Phe	Ile
Leu 305	Thr	Tyr	Glu	Gly	Gln 310	Pro	Thr	Ile	Phe	Tyr 315	Arg	Asp	Tyr	Glu	Glu 320
Trp	Leu	Asn	Lys	Asp 325	Arg	Leu	Lys	Asn	Leu 330	Ile	Trp	Ile	His	Asp 335	His
Leu	Ala	Gly	Gly 340	Ser	Thr	Asp	Ile	Val 345	Tyr	Tyr	Asp	Asn	Asp 350	Glu	Leu
Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly 360	Ser	Lys	Pro	Gly	Leu	Ile 365	Thr	Tyr
Ile	Asn 370	Leu	Ala	Ser	Ser	Lys 375	Ala	Gly	Arg	Trp	Val 380	Tyr	Val	Pro	Lys
Phe 385	Ala	Gly	Ala	Cys	Ile 390	His	Glu	Tyr	Thr	Gly 395	Asn	Leu	Gly	Gly	Trp 400
Val	Asp	Lys	Tyr	Val 405	Tyr	Ser	Ser	Gly	Trp 410	Val	Tyr	Leu	Glu	Ala 415	Pro
Ala	Tyr	Asp	Pro 420	Ala	Asn	Gly	Gln	Tyr 425	Gly	Tyr	Ser	Val	Trp 430	Ser	Tyr
Cys	Gly	Val 435	Gly												

<210> 17
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

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 gtcccatg gaggaatctg gtgggacacg atagcccaga agatacccga ctgggcaagc

60
 120

1001827087_1.txt

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atgggctacg acccctacga cttcttttgac ctcggtgagt acgaccagga gggaacggta 240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
aaccgcgttcg ttggggacta cacctggacg gactttctcaa aggtggcctc gggcaaatat 420
actgccaaact acctcgactt ccaccccaac gaggtcaagt gctgtgacga gggcacattt 480
ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggcgagc 540
gatgagagct acgccgccta cctaaggagc atcggcgttg atgcctggcg cttcgactac 600
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gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat 720
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attcccgcac tcgtcgaggc cctcaagaac gggggcacag tcgtcagccg cgaccctgtt 840
aaggccgtaa ccttcgttgc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
tacgcgttca tcctcaccta cgagggccag ccgacaatat tctaccgca ctacgaggag 960
tggtcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
agcacgagca tagtttacta cgacagcgac gagatgatct tcgtgaggaa cggctatgga 1080
agcaagcctg gccttataac ttacatcaac ctcggctcga gcaaggttgg aaggtggggt 1140
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gtggacaagt ggggtggactc aagcggctgg gtctacctcg aggcctcctgc ccacgacccg 1260
gccaacggcc agtacggcta ctccgtctgg agctactgcg gtgttggttg a 1311

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<210> 18

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 18

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Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Gly Leu Ile Met Gln Ala
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Phe Tyr Trp Asp Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala
20     25     30
Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
35     40     45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
50     55     60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Glu Gly Thr Val
65     70     75     80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
85     90     95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100    105    110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
115    120    125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130    135    140
Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
145    150    155    160
Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
165    170    175
Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180    185    190
Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
195    200    205
Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210    215    220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
225    230    235    240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala Ala Phe
245    250    255
Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly
260    265    270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275    280    285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290    295    300

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Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 19
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

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 gccgggattt cggcgatatg gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
 atgggctacg accctacga ttattttgac cttggtgagt actaccagaa gggaacgggtg 240
 gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgag caggcggaga cctcgagtgg 360
 aaccggttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
 ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggccagc 540
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 gtcaagggtc atgctccctg ggtcgtcaag gactggctga actgggtggg gggctgggag 660
 gttggagagt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggg 720
 gccaaaggtc ttgacttcgc cctctactac aagatggatg aggcctttga caacaaaaac 780
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 aaggccgtaa cttttgtagc aaaccacgac accgatataa tttggaacaa gtaccgggac 900
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 tggctcaaca aggacaggct caagaacctc atctggatac acgaccacct cgccggtgga 1020
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 agcaagccgg gactgataac atacatcaac ctgcctcaa gcaaagccgg aaggtgggtt 1140
 tatgtgccga agttcgggg cgctgcatc cagagcata ctggtaacct cggaggctgg 1200
 gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgaccct 1260
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<210> 20
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 20
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 20 25 30
 Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60

1001827087_1.txt

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65 70 75 80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
85 90 95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100 105 110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
115 120 125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130 135 140
Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
145 150 155
Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
160 165 170 175
Leu Trp Ala Ser Gln Glu Ser Tyr Ala Tyr Leu Arg Ser Ile Gly
180 185 190
Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
195 200 205
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210 215 220
Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
225 230 235 240
Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
245 250 255
Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
260 265 270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275 280 285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290 295 300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305 310 315 320
Trp Leu Asn Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His
325 330 335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
340 345 350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355 360 365
Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
370 375 380
Phe Ala Gly Ala Cys Ile His Glu His Thr Gly Asn Leu Gly Gly Trp
385 390 395 400
Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405 410 415
Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420 425 430
Cys Gly Val Gly
435

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<210> 21

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 21

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gtcccagggtg gaggaatctg gtgggacacc atcaggagca agataccgga gtggtacgag 120
gcgggaatat ccgccatttg gattcctccc gggagcaagg gtatgagcgg cggctattcg 180
atgggctacg acccctacga tgatttgagc ctgggtgagt actaccagaa gggaacggtg 240
gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
aaccggttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggcagc 540
caggagagct acgcggtata tctcaggagc atcggcatcg atgcctggcg cttcgactac 600

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gtcaaggggct	acggagcgtg	ggtcgtcaag	gactggctgg	actggtgggg	aggctggggc	660
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gcaaaagtct	tcgacttccc	gctctactac	aagatggatg	aggcctttga	caacaaaaac	780
attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgacccgttc	840
aaggccgtaa	ccittgtagc	aaaccacgac	accgatataa	tttgaacaa	gtacccggcc	900
tacgccttca	tcctcaccta	cgagggccag	ccgacgatat	tctaccgca	ctacgaggag	960
tggtcaaca	aggacaggct	caagaacctc	atctggatac	acgactacct	cgccggtgga	1020
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agcaagccgg	gactgataac	atacatcaac	ctcgcctcaa	gcaaagccgg	aagggtgggtt	1140
tatgtgccga	agttcgcggg	cgcgtgcatc	cacgagtata	ctggtaacct	cggaggctgg	1200
gtagacaagt	acgtctactc	aagcggctgg	gtctatctcg	aagctccagc	ttacgaccct	1260
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<210> 22
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 22
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 20 25 30
 Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Gly Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Asp Leu Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Val Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
 225 230 235 240
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp Tyr
 325 330 335
 Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365

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Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 23
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> synthetically generated oligonucleotide

<400> 23
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 gcgggaatat ccgccatttg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
 gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
 ggcataaaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
 aacccgttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
 ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggcccagc 540
 caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac 600
 gtcaagggct acggagcgtg ggtcgtcaag gactggctgg actggtgggg aggctgggcc 660
 gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat 720
 gcaaaaagtct tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac 780
 attccagcgc tcgtctctgc cttcagaac ggccagactg ttgtctcccg cgaccggttc 840
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 agcaagcctg gccttataac ttacatcaac ctcggtctga gcaaggttgg aagggtgggtc 1140
 tacgttccga agttcgcggg agcgtgcac cagagtaca ccggcaacct cggcggctgg 1200
 gtggacaagt ggggtggactc aagcgggtgg gtgtacctg aggccctgc ccacgaccg 1260
 gccaacggct attacggcta ctccgtctgg agctattgcy gtgttggtcg a 1311

<210> 24
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> synthetically generated peptide

<400> 24
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 Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125

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Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
 225 230 235
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315
 Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Met Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 25

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 25

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gccgggattt	cggcgatatg	gattcctccc	gcgagcaagg	gtatgagcgg	cggctattcg	180
atgggctacg	acccctacga	ttattttgac	ctcggtgagt	actaccagaa	gggaacgggtg	240
gaaacgaggt	tcggctcaaa	gcaggagctc	ataaacatga	taaacaccgc	ccacgcctat	300
ggcatgaagg	taatagccga	tatagtcatc	aaccaccgcg	cggcggtga	cctggagtgg	360
aaccccttcg	tgaacgacta	tacctggacc	gactttctca	aggctcgcgtc	gggtaaatac	420
acggccaact	acctcgactt	ccacccgaac	gagctccatg	cgggcgattc	cggaacattt	480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctgggccagc	540
caggagagct	acgcggcata	tctcaggagc	atcggcatcg	atgcctggcg	cttcgactac	600
gtcaagggct	atgctccctg	ggtcgtcaag	gactggctga	actggtgggg	aggctgggcg	660
gttggaaggt	actgggacac	caacgtcgac	gctgtttctc	actgggcata	ctcgagcggg	720
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attcccgcgc	tgggtggcgc	cctcagatac	ggtcagacag	tggtcagccg	cgaccggttc	840
aaggctgtga	cgtttgtagc	caaccacgat	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggctcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga	1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggca	cggctacggg	1080

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gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aaggtggggtt	1140
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gtggacaagt	gggtggactc	aagcggctgg	gtctacctcg	aggctcctgc	ccacgaccgg	1260
gccaacggcc	agtacggcta	ctccgtctgg	agctattgca	gtgttgggtg	a	1311

<210> 26
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 26
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 20 25 30
 Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Asn Asn Ile Pro Ala Leu Val Gly Ala Leu Arg Tyr Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
 Ile Phe Val Arg His Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430

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1001827087_1.txt

Cys Gly Val Gly
435

<210> 27
<211> 1311
<212> DNA
<213> Artificial Sequence

<220>
<223> Synthetically generated oligonucleotide

<400> 27
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gtcccagggt gaggaatctg gtgggacacc atcaggagca agataccgga gtggtacgag 120
gcgggaatat ccgccatttg gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
atgggctacg acccctacga ttattttgac ctcggtgagt actaccagaa gggaacgggtg 240
gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
ggcataaaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
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actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
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gccaaaggtct ttgacttcgc cctctactac aagatggacg cggcctttga caacaagaac 780
attcccgcac tcgtcgaggc cctcaagaac gggggcacag tcgtcagccg cgaccggtt 840
aaggccgtaa ctttcgttgc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
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gccaacggcc agtacggcta ctccgtctgg agctactgag gtgttgggtg a 1311

<210> 28
<211> 436
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetically generated peptide

<400> 28
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Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg
20 25 30
Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
35 40 45
Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
50 55 60
Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
65 70 75 80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
85 90 95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100 105 110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
115 120 125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130 135 140
Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
145 150 155 160
Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
165 170 175
Leu Trp Ala Ser Gln Glu Ser Tyr Ala Tyr Leu Arg Ser Ile Gly
180 185 190

1001827087_1.txt

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Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
195 200 205
Val Lys Asp Trp Leu Asn Trp Gly Gly Trp Ala Val Gly Glu Tyr
210 215 220
Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
225 230 235 240
Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Ala Ala Phe
245 250 255
Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly
260 265 270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275 280 285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290 295 300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305 310 315 320
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
325 330 335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
340 345 350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355 360 365
Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
370 375 380
Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385 390 395 400
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405 410 415
Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420 425 430
Cys Gly Val Gly
435

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<210> 29

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 29

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gccgggattt cggcgatatg gattcccccg gcgagcaagg gcatgggagg cgcctattcg 180
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aaccggttcg ttggggacta cacctggacg gacttctcaa aggtggtctc gggcaaatat 420
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gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgacctt 1260
gccaacgggc agtatggcta ctccgtgtgg agctactgag gtgttggttg a 1311

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<210> 30

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 30

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Phe Tyr Trp Asp 20 Val Pro Met Gly 25 Gly Ile Trp Trp Asp 30 Thr Val Ala
Gln Lys 35 Ile Pro Asp Trp Ala Ser 40 Ala Gly Ile Ser 45 Ala Ile Trp Ile
Pro 50 Pro Ala Ser Lys Gly 55 Met Gly Gly Ala Tyr Ser 60 Met Gly Tyr Asp
Pro 65 Tyr Asp Phe Phe Asp 70 Leu Gly Glu Tyr Asp 75 Gln Lys Gly Thr Val
Glu Thr Arg Phe 85 Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
Ala His Ala Tyr 100 Gly Ile Lys Val Ile 105 Ala Asp Ile Val Ile Asn His
Arg Ala Gly 115 Gly Asp Leu Glu Trp 120 Asn Pro Phe Val Gly 125 Asp Tyr Thr
Trp Thr 130 Asp Phe Ser Lys Val 135 Val Ser Gly Lys Tyr 140 Thr Ala Asn Tyr
Leu 145 Asp Phe His Pro Asn 150 Glu Leu His Ala Gly 155 Asp Ser Gly Thr Phe
Gly Gly Tyr Pro Asp 165 Ile Cys His Asp Lys 170 Ser Trp Asp Gln Tyr Trp
Leu Trp Ala Ser 180 Gln Glu Ser Tyr Ala 185 Ala Tyr Leu Arg Ser Ile Gly
Ile Asp Ala 195 Trp Arg Phe Asp Tyr 200 Val Lys Gly Tyr Ala 205 Pro Trp Val
Val Lys 210 Asp Trp Leu Asn Trp 215 Trp Gly Gly Trp Ala 220 Val Gly Glu Tyr
Trp 225 Asp Thr Asn Val Asp 230 Ala Val Leu Asn Trp 235 Ala Tyr Ser Ser Gly
Ala Lys Val Phe Asp 245 Phe Ala Leu Tyr Tyr 250 Lys Met Asp Glu Ala Phe
Asp Asn Lys Asn 260 Ile Pro Ala Leu Val Ser 265 Ala Leu Gln Asn Gly Gln
Thr Val Val 275 Ser Arg Asp Pro Phe 280 Lys Ala Val Thr Phe Val Ala Asn
His Asp 290 Thr Asp Ile Ile Trp 295 Asn Lys Tyr Leu Ala Tyr Ala Phe Ile
Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr 300 Arg Asp Tyr Glu Glu
305 Trp Leu Asn Lys Asp 310 Arg Leu Asn Asn Leu 315 Ile Trp Ile His Asp His
Leu Ala Gly Gly 325 Ser Thr Asp Ile Val 330 Tyr Tyr Asp Asn Asp Glu Leu
Ile Phe Val 340 Arg Asn Gly Tyr Gly 345 Asp Lys Pro Gly Leu Ile Thr Tyr
Ile Asn 355 Leu Gly Ser Ser Lys 360 Ala Gly Arg Trp Val Tyr Val Pro Lys
370 Phe Ala Gly Ala Cys Ile 375 His Glu Tyr Thr Gly 380 Asn Leu Gly Gly Trp
385 Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp 395 Val Tyr Leu Glu Ala Pro
Ala Tyr Asp Pro 405 Ala Asn Gly Gln Tyr 410 Gly Tyr Ser Val Trp Ser Tyr
425 Cys Gly Val Gly 435

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<210> 31

<211> 1311

<212> DNA

<213> Artificial Sequence

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<220>

<223> Synthetically generated oligonucleotide

<400> 31

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gtcccagggtg	gaggaatctg	gtgggacacc	atcaggagca	ggataccgga	gtggtacgag	120
gcgggaatat	ccgccatttg	gattcccccg	gcgagcaagg	gcatgggcgg	cgcctattcg	180
atgggctacg	acccctacga	cttctttgac	ctcggtgagt	acgaccagaa	gggaacggta	240
gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacacggc	ccatgcctac	300
ggcataaagg	tcatagcgga	catcgtcata	aaccaccgcg	caggcggaga	cctcgagtgg	360
aacccgttcg	ttggggacta	cacctggacg	gactttctcaa	aggtggcctc	gggcaaatat	420
actgccaaact	acctcgactt	ccacccgaac	gagctccatg	cgggcgattc	cggaacattt	480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctggggccagc	540
caggagagct	acgcggcata	tctcaggagc	atcggcatcg	atgcctggcg	ctttgactac	600
gtgaagggtc	acggagcgtg	ggtcgtcaag	gactggctca	actggtgggg	cggctggggc	660
gttggtcgagt	actgggacac	caacgttgat	gcactcctca	actgggccta	ctcgagcggc	720
gccaaaggtct	tcgacttccc	gctctactac	aagatggacg	aggccttcga	taacaacaac	780
attcccgcgc	tggtggacgc	cctcagatac	ggtcagacag	tggtcagccg	cgaccgcgtt	840
aaggctgtga	cgtttgtagc	caaccacgat	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtccaaca	aggaataagct	caagaacctc	atctggatac	atgacaacct	ggccggagga	1020
agcacgagca	tagtttacta	cgacagcgac	gagatgatct	tcgtgaggac	cggctatgga	1080
agcaagcctg	gccttataac	ttacatcaac	ctcggctcga	gcaaggttgg	aaggtgggtt	1140
tatgtgccga	agttcgcggt	cgctgcatc	cacgagtata	ctggtaacct	cggaggctgg	1200
gtagacaagt	acgtctactc	aagcggctgg	gtctatctcg	aagctccagc	ttacgaccct	1260
gccaacgggc	agtatggcta	ctccgtgtgg	agctatttgcg	gtgttggctg	a	1311

<210> 32

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 32

Met	Ala	Lys	Tyr	Ser	Glu	Leu	Glu	Gly	Gly	Val	Ile	Met	Gln	Ala
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Phe	Tyr	Trp	Asp	Val	Pro	Gly	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile
			20					25				30		Arg
Ser	Arg	Ile	Pro	Glu	Trp	Tyr	Glu	Ala	Gly	Ile	Ser	Ala	Ile	Trp
		35					40				45			Ile
Pro	Pro	Ala	Ser	Lys	Gly	Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr
	50					55					60			Asp
Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	Glu	Tyr	Asp	Gln	Lys	Gly	Thr
65					70				75					80
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn
			85					90					95	Thr
Ala	His	Ala	Tyr	Gly	Ile	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn
			100					105					110	His
Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Gly	Asp	Tyr
		115					120					125		Thr
Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn
	130					135					140			Tyr
Leu	Asp	Phe	His	Pro	Asn	Glu	Leu	His	Ala	Gly	Asp	Ser	Gly	Thr
145					150					155				Phe
Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	Asp	Lys	Ser	Trp	Asp	Gln	Tyr
			165						170					175
Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile
			180					185					190	Gly
Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp
		195					200					205		Val
Val	Lys	Asp	Trp	Leu	Asn	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu
	210					215					220			Tyr
Trp	Asp	Thr	Asn	Val	Asp	Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser
225					230					235				Gly
Ala	Lys	Val	Phe	Asp	Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala
			245						250					255

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Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln
260
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
325
Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
340
Ile Phe Val Arg Thr Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355
Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
370
Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385
Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405
Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420
Cys Gly Val Gly
435

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<210> 33
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

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<400> 33
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gtgccttcag gaggaatatg gtgggacaca atacggcaga agataccgga gtggtacgat 120
gccggaatct ccgcaatatg gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
atgggctacg accctacga ttattttgac ctcggtgagt actaccagaa gggaacgggtg 240
gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
ggcataaagg tcatagcgga catcgtcata aaccaccgag caggcggaga cctcgagtgg 360
aaccggttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggccagc 540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg ctttgactac 600
gtgaagggct acggagcgtg ggtcgtcaag gactggctca actggtgggg cggctgggcc 660
gttggcgagt actgggacac caacgttgat gcactcctca actgggccta ctcgagcggc 720
gccaaggtct tcgactttcc gctctactac aagatggacg cggcctttga caacaagaac 780
attccgcac tcgtcgaggc cctcaagaac gggggcacag tcgtcagccg cgaccggtt 840
aaggccgtaa ccttcgttgc aaaccacgac accgatataa tctggaccaa gtaccttgct 900
tatgctttca tcctcaccta cgaaggccag cccgtcatat tctaccgca ctacgaggag 960
tggtcaaca aggacaggtt gaacaacctc atatggatac acgaccacct cgcagggtgga 1020
agcaccgaca tagctacta cgataacgat gaactcatct tcgtcaggaa cggctacggg 1080
gacaagccgg ggcttataac ctacatcaac ctaggctcga gcaaggccgg aaggtgggtt 1140
tacgttccga agttcgcagg ctcgtgcata cagagtaca ccggcaatct cggcggctgg 1200
gtggacaagt ggtgggactc aagcggctgg gtctacctg aggtcctgc ccacgaccg 1260
gccaacggcc agtacggcta ctccgtctgg agctactgag gtgttggtg a 1311

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<210> 34
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

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<400> 34
Met Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala
1 5 10 15

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Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Thr Lys Tyr Leu Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His
 325 330 335
 Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 35
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> synthetically generated oligonucleotide

<400> 35
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 gtcccaggtg gaggaatctg gtgggacacc atcaggagca agataccgga gtggtacgag 120
 gcgggaatat ccgccatttg gattcccccg gcgagcaagg gcatgggagg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240

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gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacaccgc	ccacgcctac	300
ggcatcaagg	tcatcgaga	catagtaatc	aaccaccgcg	cggaggaga	ccttgagtgg	360
aacccttcg	tcaatgacta	cacctggacg	gacttctcga	aggtcgcttc	cggcaagtac	420
acggccaact	acctcgactt	ccacccaac	gaggtcaagt	gctgtgacga	gggcacattt	480
ggaggcttcc	cagacatagc	ccacgagaag	agctgggacc	agcactggct	ctgggcgagc	540
gatgagagct	acgccgccta	cctaaggagc	atcggcgttg	atgcctggcg	cttcgactac	600
gtcaagggct	atgctccctg	ggtcgtcaag	gactggctga	actggtgggg	aggctgggcg	660
gttgagagct	actgggacac	caacgtcgac	gctgttctca	actgggcata	ctcgagcggg	720
gccaaggctc	ttgacttcgc	cctctactac	aagatggacg	cggcctttga	caacaagaac	780
attcccgcac	tcgtcgaggc	cctcaagaac	gggggcacag	tcgtcagccg	cgaccctgtt	840
aaggccgtaa	ccttcgttgc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgaggggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtcaaca	aggataagct	caagaacctc	atctggatac	atgacaacgt	cggcggagga	1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg	1080
gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aagggtgggt	1140
tacgttccga	agttcgcagg	ctcgtgcata	cacgagtaca	cgggcaatct	cggcggctgg	1200
gtggacaagt	gggtggactc	aagcggctgg	gtctacctcg	aggctcctgc	ccacgaccgc	1260
gccaacggcc	agtacggcta	ctccgtctgg	agctactgcg	gtgttggtg	a	1311

<210> 36

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 36

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Phe	Tyr	Trp	Asp	Val	Pro	Gly	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg
			20					25					30		
Ser	Lys	Ile	Pro	Glu	Trp	Tyr	Glu	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile
		35					40					45			
Pro	Pro	Ala	Ser	Lys	Gly	Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr	Asp
	50					55					60				
Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	Glu	Tyr	Asp	Gln	Lys	Gly	Thr	Val
65					70					75					80
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr
				85					90					95	
Ala	His	Ala	Tyr	Gly	Ile	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
			100					105					110		
Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr	Thr
		115					120					125			
Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
	130					135					140				
Leu	Asp	Phe	His	Pro	Asn	Glu	Val	Lys	Cys	Cys	Asp	Glu	Gly	Thr	Phe
145					150					155					160
Gly	Gly	Phe	Pro	Asp	Ile	Ala	His	Glu	Lys	Ser	Trp	Asp	Gln	His	Trp
				165					170					175	
Leu	Trp	Ala	Ser	Asp	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
			180					185					190		
Val	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Ala	Pro	Trp	Val
		195					200					205			
Val	Lys	Asp	Trp	Leu	Asn	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	
	210					215				220					
Trp	Asp	Thr	Asn	Val	Asp	Ala	Val	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Gly
225					230					235					240
Ala	Lys	Val	Phe	Asp	Phe	Ala	Leu	Tyr	Tyr	Lys	Met	Asp	Ala	Ala	Phe
				245					250					255	
Asp	Asn	Lys	Asn	Ile	Pro	Ala	Leu	Val	Glu	Ala	Leu	Lys	Asn	Gly	Gly
			260					265					270		
Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
			275				280					285			
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
	290					295					300				
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
305					310					315					320

1001827087_1.txt

Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Val Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 37
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 37
 atggccaagt acctggagct cgaagagggc ggggtcataa tgcaggcggt ctactgggac 60
 gtgcccttcag gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgat 120
 gccggaatct ccgcaatat gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
 gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctat 300
 ggcatagaagg taatagccga tatagtcac aaccaccgcg ccggcggtga cctggagtgg 360
 aacccttcg tgaacgacta tacctggacc gacttctcaa aggtcgcgtc gggtaaatac 420
 acggccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
 ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctgggcccagc 540
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 gtgaagggtc acggagcgcg ggtcgtcaag gactggctca actggtgggg cggctgggccc 660
 gttggcgagt actgggacac caacgttgat gcactcctca actgggccta ctcgagcggc 720
 gccaaaggtct tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac 780
 attccagcgc tcgtctctgc cttcagaac ggccagactg ttgtctcccg cgacccttc 840
 aaggccgtaa cttttagtag aaaccacgac accgatataa tctggaacaa gtatccagcc 900
 tacgcgttca tcctcaccta cgagggccag ccgacaatat tctatcgcg ctagcaggag 960
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 agcaagccgg gactgataac atacatcaac ctgcctcaa gcaaagccgg aaggtgggtt 1140
 tacgttccga agttcgcagg ctcgtgcata cagagtaca ccggcaatct cggcggctgg 1200
 gtggacaagt gggtagactc aagcggctgg gtctacctg aggtcctgc ccacgaccg 1260
 gccaacggcc agtacggcta ctccgtctgg agctactgcg gggtaggggtg a 1311

<210> 38
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 38
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 Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80

1001827087_1.txt

Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr
Ala	His	Ala	Tyr	85	Gly	Met	Lys	Val	Ile	90	Ala	Asp	Ile	Val	110
Arg	Ala	Gly	Gly	100	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr
Trp	Thr	Asp	Phe	115	Ser	Lys	Val	120	Ala	Ser	Gly	Lys	Tyr	125	Thr
Leu	Asp	Phe	His	130	Pro	Asn	Glu	135	Leu	His	Ala	Gly	Asp	Ser	Thr
Gly	Gly	Tyr	Pro	145	Asp	Ile	Cys	150	His	Asp	Lys	Ser	Trp	Asp	160
Leu	Trp	Ala	Ser	165	Gln	Glu	Ser	170	Tyr	Ala	Tyr	Leu	Arg	Ser	175
Ile	Asp	Ala	Trp	180	Arg	Phe	Asp	185	Val	Lys	Gly	Tyr	Gly	Ala	190
Val	Lys	Asp	Trp	195	Leu	Asn	Trp	200	Gly	Gly	Trp	Ala	Val	Gly	205
Trp	Asp	Thr	Asn	210	Val	Asp	Ala	215	Leu	Leu	Asn	Trp	Ala	Tyr	220
Ala	Lys	Val	Phe	225	Asp	Phe	Pro	230	Leu	Tyr	Tyr	Lys	Met	Asp	235
Asp	Asn	Lys	Asn	245	Ile	Pro	Ala	250	Leu	Val	Ser	Ala	Leu	Gln	255
Thr	Val	Val	Ser	260	Arg	Asp	Pro	265	Phe	Lys	Ala	Val	Thr	Phe	270
His	Asp	Thr	Asp	275	Ile	Ile	Trp	280	Asn	Lys	Tyr	Pro	Ala	Tyr	285
Leu	Thr	Tyr	Glu	290	Gly	Gln	Pro	295	Thr	Ile	Phe	Tyr	Arg	Asp	300
Trp	Leu	Asn	Lys	305	Asp	Lys	Leu	310	Lys	Asn	Leu	Ile	Trp	Ile	315
Leu	Ala	Gly	Gly	325	Ser	Thr	Asp	330	Ile	Val	Tyr	Tyr	Asp	Asn	335
Ile	Phe	Val	Arg	340	Asn	Gly	Tyr	345	Gly	Ser	Lys	Pro	Gly	Leu	350
Ile	Asn	Leu	Ala	355	Ser	Ser	Lys	360	Ala	Gly	Arg	Trp	Val	Tyr	365
Phe	Ala	Gly	Ser	370	Cys	Ile	His	375	Glu	Tyr	Thr	Gly	Asn	Leu	380
Val	Asp	Lys	Trp	385	Val	Asp	Ser	390	Ser	Gly	Trp	Val	Tyr	Leu	395
Ala	His	Asp	Pro	405	Ala	Asn	Gly	410	Gln	Tyr	Gly	Tyr	Ser	Val	415
Cys	Gly	Val	Gly	420				425						Trp	430
				435											

<210> 39

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 39

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gccggaatct	ccgcaatatg	gattcctccc	gcgagcaggg	gtatgagcgg	cggctattcg	180
atgggctacg	acccctacga	ttattttgac	ctcggtagt	actaccagaa	gggaacgggtg	240
gaaacgaggt	tcggctcaaa	gcaggagctc	ataaacatga	taaaccaccg	ccacgcctat	300
ggcatgaagg	taatagccga	tatagtcatc	aaccaccgcg	cggcggtga	cctggagtgg	360
aacccttcg	tgaacgacta	tacctggacc	gacttctcaa	aggctcgcgc	gggtaaatac	420
acggccaact	acctcgactt	ccaccggaac	gagctccatg	cgggcgattc	cggaacattt	480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctgggccagc	540
caggagagct	acgcggcata	tctcaggagc	atcggtatcg	atgcctggcg	ctttgactac	600
gtgaagggct	acggagcgtg	ggtcgtcaag	gactggctca	actggtgggg	cggctgggccc	660
gttggcgagt	actgggaccc	caacgttgat	gccctcctcc	cctgggccta	ctcgagcggc	720

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aaggccgtaa	cctttgtagc	caaccacgat	accgatataa	tctggaacaa	gtatccagcc	900
tacgcggtca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggctcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga	1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg	1080
gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aaggtgggtc	1140
tacgttccga	agttcgcggg	agcgtgcac	cacgagtaca	ccggcaacct	cggcggctgg	1200
gtggacaagt	gggtggactc	aagcgggtgg	gtgtacctcg	aggcccctgc	ccacgaccgc	1260
gccaacggct	attacggcta	ctccgtctgg	agctactgcg	gggtgggctg	a	1311

<210> 40

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 40

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Phe	Tyr	Trp	Asp	Val	Pro	Ser	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg
			20					25					30		
Gln	Lys	Ile	Pro	Glu	Trp	Tyr	Asp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile
		35					40					45			
Pro	Pro	Ala	Ser	Arg	Gly	Met	Ser	Gly	Gly	Tyr	Ser	Met	Gly	Tyr	Asp
	50					55					60				
Pro	Tyr	Asp	Tyr	Phe	Asp	Leu	Gly	Glu	Tyr	Tyr	Gln	Lys	Gly	Thr	Val
65					70					75				80	
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Ile	Asn	Met	Ile	Asn	Thr
				85					90					95	
Ala	His	Ala	Tyr	Gly	Met	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
			100					105					110		
Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr	Thr
		115					120					125			
Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
	130					135					140				
Leu	Asp	Phe	His	Pro	Asn	Glu	Leu	His	Ala	Gly	Asp	Ser	Gly	Thr	Phe
145					150					155					160
Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	Asp	Lys	Ser	Trp	Asp	Gln	Tyr	Trp
			165						170					175	
Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
			180					185					190		
Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val
		195					200					205			
Val	Lys	Asp	Trp	Leu	Asn	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr
	210					215					220				
Trp	Asp	Pro	Asn	Val	Asp	Ala	Leu	Leu	Pro	Trp	Ala	Tyr	Ser	Ser	Gly
225					230					235					240
Ala	Lys	Val	Phe	Asp	Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
				245					250					255	
Asp	Asn	Lys	Asn	Ile	Pro	Ala	Leu	Val	Ser	Ala	Leu	Gln	Asn	Gly	Gln
			260					265					270		
Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
		275					280					285			
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
	290					295				300					
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
305					310					315					320
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	Asn
				325					330					335	
Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu
			340					345					350		
Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	Asp	Lys	Pro	Gly	Leu	Ile	Thr	Tyr
		355					360					365			
Ile	Asn	Leu	Gly	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys
	370					375					380				

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Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 41
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 41
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 gccggaatct ccgcaatat gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
 atgggctacg acccctacga ttattttgac ctcggtgagt actaccagaa gggaacgggtg 240
 gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
 aacccgttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
 ggaggctatc ccgacatat ccacgacaag agctgggacc agtactggct ctggggccagc 540
 caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg ctttgactac 600
 gtgaagggct acggagcgtg ggtcgtcaag gactggctca actggtgggg cggctgggcc 660
 gttggcgagt actgggacac caacgttgat gcactcctca actgggccta ctcgagcggc 720
 gccaaagttc tcgacttccc gctctactac aagatggacg cggcctttga caacaagaac 780
 attcccgcac tcgtcgaggc cctcaagaac gggggcacag tcgtcagccg cgaccggtt 840
 aaggccgtaa ccttcgttgc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
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 agcacgagca tagtttacta cgacagcgac gagatgatct tcgtgaggaa cggctatgga 1080
 agcaagcctg gccttataac ttacatcaac ctcggtcga gcaagggttg aaggtgggtt 1140
 tatgtgcga agttcgcggg cgctgcatc cacgagtata ctggtaacct cggaggctgg 1200
 gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgacct 1260
 gccaacgggc agtatggcta ctccgtgtgg agctactgcg gtgttgggtg a 1311

<210> 42
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 42
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 Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140

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Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Ala Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Glu Ala Leu Lys Asn Gly Gly
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 43

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 43

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gcgggaatat	ccgccatttg	gattcccccg	gcgagcaagg	gcatgggcgg	cgctatttcg	180
atgggctacg	acccctacga	cttctttgac	ctcggtgagt	acgaccagaa	gggaacggta	240
gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacacggc	ccatgcctac	300
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aaccgttcg	ttggggacta	cacctggacg	gacttctcaa	aggtggcctc	gggcaaatat	420
actgccaaat	acctcgactt	ccaccccaac	gaggtcaagt	gctgtgacga	gggcacattt	480
ggaggcttcc	cagacatagc	ccacgagaag	agctgggacc	agcactggct	ctgggcgagc	540
gatgagagct	acgccgccta	cctaaggagc	atcggcggtg	atgcctggcg	cttcgactac	600
gtcaagggct	acggagcgtg	ggtcgtcaag	gactggctgg	actggtgggg	aggctgggcc	660
gtcggggagt	actgggacac	aaacgttgat	gcactgctca	actgggccta	ctcgagcgat	720
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attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgacccttcc	840
aaggccgtaa	cctttgtagc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtctcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgtcggagga	1020
agcacgagca	tagtttacta	cgacagcgac	gagatgatct	tcgtgaggaa	cggctatgga	1080
agcaagcctg	gccttataac	ttacatcaac	ctcggctcga	gcaaggttgg	aaggtggggt	1140
tacgttccga	agttcgcagg	ctcgtgcata	cacgagtaca	cgggcaatct	cggcggctgg	1200

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gtggacaagt ggggtggactc aagcggctgg gtctacctcg aggctcctgc ccacgacccg 1260
gccaacggcc agtacggcta ctccgtctgg agctactgcg gtgttggtg a 1311

<210> 44
<211> 436
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetically generated peptide

<400> 44
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20 25 30
Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
35 40 45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
50 55 60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
65 70 75 80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
85 90 95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100 105 110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
115 120 125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130 135 140
Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
145 150 155 160
Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
165 170 175
Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180 185 190
Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
195 200 205
Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210 215 220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
225 230 235 240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
245 250 255
Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
260 265 270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275 280 285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
290 295 300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
305 310 315 320
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
325 330 335
Leu Val Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
340 345 350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355 360 365
Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
370 375 380
phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385 390 395 400
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405 410 415
Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420 425 430
Cys Gly Val Gly
435

<210> 45
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 45
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 gcgggaatat ccgccatttg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
 gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgcg caggcggaga cctcgagtgg 360
 aaccgcgttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaaact acctcgactt ccaccccaac gaggtcaagt gctgtgacga gggcacattt 480
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 aaggccgtaa cctttgtagc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
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 tggctcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
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 gacaagccgg ggcttataac ctacatcaac ctaggctcga gcaaggccgg aagggtgggt 1140
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 gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgaccct 1260
 gccaacgggc agtatggcta ctccgtgtgg agctattgcg gtgttggtg a 1311

<210> 46
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 46
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 20 25 30
 Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
 145 150 155 160
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
 165 170 175
 Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205

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Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
 260 265 270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
Ala Tyr Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
Cys Gly Val Gly
 435

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<210> 47

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 47

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gcgggaatat ccgccatttg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
atgggctacg accctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctat 300
ggcatgaagg taatagccga tatagtcatc aaccaccgcg ccggcggtga cctggagtgg 360
aacccttctg tgaacgacta tacctggacc gacttctcaa aggtcgcgtc gggtaaatac 420
acggccaact acctcgactt ccacccaac gaggtcaagt gctgtgacga gggcacattt 480
ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggcgagc 540
gatgagagct acgccgcta cctaaggagc atcggcgttg atgcctggcg ctttgactac 600
gtgaagggct acggagcgtg ggtcgtcaag gactggctca actggtgggg cggttgggcc 660
gttggcgagt actgggacac caacgttgat gcactcctca actgggccta ctcgagcggc 720
gccaaggctc tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac 780
attccagcgc tcgtctctgc cttcagaac ggccagactg ttgtctcccg cgaccggttc 840
aaggccgtaa cttttgtagc aaaccacgta accgatataa tctggaacaa gtaccttgct 900
tatgctttca tcctcaccta cgaaggccag cccgtcatat tctaccgca ctacgaggag 960
tggctcaaca aggacaggtt gaacaacctc atatggatac acgaccacct cgcagggtgga 1020
agcacgagca tagtttacta cgacagcgac gagatgatct tcgtgaggaa cggctatgga 1080
agcaagcctg gccttataac ttacatcaac ctcggtctga gcaaggttgg aaggtggggt 1140
tacgttccga agttcgcagg cccgtgcata cagagtaca ccggcaatct cggcggctgg 1200
gtggacaagt gggtggactc aagcggctgg gtctacctcg aggtctctgc ccacgaccgc 1260
gccaacggcc agtacggcta ctccgtctgg agctactgcg gtgttgggta g 1311

```

<210> 48

<211> 436

<212> PRT

<213> Artificial Sequence

1001827087_1.txt

<220>

<223> Synthetically generated peptide

<400> 48

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Met Ala Lys Tyr Thr Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala
1      5      10      15
Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg
20      25      30
Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
35      40      45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
50      55      60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
65      70      75      80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
85      90      95
Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
100     105     110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
115     120     125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130     135     140
Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
145     150     155     160
Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
165     170     175
Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180     185     190
Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
195     200     205
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210     215     220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
225     230     235     240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
245     250     255
Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
260     265     270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
275     280     285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu Ala Tyr Ala Phe Ile
290     295     300
Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu
305     310     315     320
Trp Leu Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His
325     330     335
Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
340     345     350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
355     360     365
Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
370     375     380
Phe Ala Gly Pro Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
385     390     395     400
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
405     410     415
Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
420     425     430
Cys Gly Val Gly
435

```

<210> 49

<211> 387

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

1001827087_1.txt

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<400> 49
gtggtttatg acgatgtccg ctatgacctt tatgccgtag gcatggggcg tgtttatcat      60
gttcacgagc tcctgcttgg agccaaagcg cgtctctacc gttcccttct ggtcgtactc      120
accgaggtca aagaagtcgt aggggtcgta gcccatcgaa taggcgccgc ccatgccctt      180
gctcgccggg ggaatccata tcgccgaaat cccggcgctt gccagtcgg gtatcttctg      240
ggctatcgtg tcccaccaga ttcttcccat ggggacgtcc cagtagaagg cctgcattat      300
gagcccgccc tcttcgagcc cggaatactt tgccataagt tacctcctac tagtagatta      360
aaattctgtt tcctgtgtga aattgtt                                387

```

```

<210> 50
<211> 129
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Synthetically generated peptide

```

```

<400> 50
Val Val Tyr Asp Asp Val Arg Tyr Asp Leu Tyr Ala Val Gly Met Gly
1      5      10      15
Arg Val Tyr His Val His Glu Leu Leu Leu Gly Ala Lys Ala Arg Leu
20     25     30
Tyr Arg Ser Leu Leu Val Val Leu Thr Glu Val Lys Glu Val Val Gly
35     40     45
Val Val Ala His Arg Ile Gly Ala Ala His Ala Leu Ala Arg Arg Gly
50     55     60
Asn Pro Tyr Arg Arg Asn Pro Gly Ala Cys Pro Val Gly Tyr Leu Leu
65     70     75     80
Gly Tyr Arg Val Pro Pro Asp Ser Ser His Gly Asp Val Pro Val Glu
85     90     95
Gly Leu His Tyr Glu Pro Ala Leu Phe Glu Pro Gly Ile Leu Cys His
100    105    110
Lys Leu Pro Pro Thr Ser Arg Leu Lys Phe Cys Phe Leu Cys Glu Ile
115    120    125
Val

```

```

<210> 51
<211> 1311
<212> DNA
<213> Artificial Sequence

```

```

<220>
<223> Synthetically generated oligonucleotide

```

```

<400> 51
atggccaagt acctggagct cgaagagggc ggggtcataa tgcaggcggt ctactgggac      60
gtgccttcag gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgat      120
gccggaatct ccgcaatatg gattcccccg gcgagcaagg gcatggggcg cgcctattcg      180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta      240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacaccgc ccacgcctat      300
ggcatgaagg taatagccga tatagtcatc aaccaccgcg ccggcggtga cctggagtgg      360
aacccttcg tgaacgacta tacctggacc gacttctcaa aggtcgcgtc gggtaaatac      420
acggccaact acctgactt ccacccaac gaggtcaagt gctgtgacga gggcacattt      480
ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggcgagc      540
gatgagagct acgccgccta cctaaggagc atcggcggtg atgcctggcg ctttgactac      600
gtgaagggct acggagcgtg ggtcgtcaag gactggctca actggtgggg cggctgggcc      660
gttggcgagt actgggacac caacgttgat gcactcctca actgggccta ctcgagcggc      720
gccaaagttc tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac      780
attccagcgc tcgtctctgc cttcagaac ggccagactg ttgtctcccg cgaccggtt      840
aaggccgtaa ctttgtagc aaaccacgac accgatataa tctggaacaa gtatccagcc      900
tacgcgttca tcctcaccta cgagggccag ccgacaatat tctaccgca ctacgaggag      960
tggctcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga      1020
agcactgaca tcgtttacta cgacaacgac gagctgatat tcgtgagaaa cggctacgga      1080
agcaagccgg gactgataac atacatcaac ctgcctcaa gcaaagccgg aaggtgggtt      1140
tacgttcga agttcgagg ctctgcgata cacgagtaca ccggcaatct cggcggctgg      1200
gtggacaagt ggggtggactc aagcggctgg gtctacctcg aggtcctgc ccacgacctg      1260

```

gccaacggcc agtacggcta ctccgtctgg agctattgcg gtgttggtg a

<210> 52
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> synthetically generated peptide

<400> 52
 Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala
 1 5 10 15
 Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
 145 150 155 160
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
 165 170 175
 Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn
 325 330 335
 Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 53
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 53
 atggccaagt actccgagct ggaagagggc ggcgttataa tgcaggcctt ctactgggac 60
 gtcccagggt gaggaatctg gtgggacacc atcaggagca agataccgga gtggtacgag 120
 gcgggaatat ccgccatttg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
 atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
 gagacgcgtt ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgca caggcggaga cctcgagtgg 360
 aaccgcgttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccacccaac gaggtcaagt gctgtgacga gggcacattt 480
 ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggcgagc 540
 gatgagagct acgccgccta cctaaggagc atcggcgctt atgcctggcg cttcgactac 600
 gtcaagggtc acggagcgtg ggtcgtcaag gactggctgg actggtgggg aggctgggcc 660
 gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat 720
 gcaaaagtct tcgacttccc gctctactac aagatggatg aggcctttga caacaaaaac 780
 attccagcgc tcgtctctgc ccttcagaac ggccagactg ttgtctcccg cgaccgcttc 840
 aaggccgtaa cttttgtagc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
 tacgcgttca tcctcaccta cgagggccag ccgacaatat tctaccgca ctacgaggag 960
 tggctcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
 agcactgaca tcgtttacta cgacaacgac gagctgatat tcgtgagaaa cggctacgga 1080
 agcaagccgg gactgataac atacatcaac ctgcctcaa gcaaagccgg aagggtgggtc 1140
 tacgttccga agttcgcggg agcgtgcata cagagtaca ccggcaacct cggcggctgg 1200
 gtggacaagt ggggtggactc aagcgggtgg gtgtacctcg aggccctgc ccacgacctg 1260
 gccaacggct attacggcta ctccgtctgg agctactgca gtgttggtg a 1311

<210> 54
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 54
 Met Ala Lys Tyr Ser Glu Leu Glu Gly Gly Val Ile Met Gln Ala
 1 5 10 15
 Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Thr Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
 145 150 155 160
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
 165 170 175
 Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asp Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr

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[illegible]

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<210> 55
<211> 1311
<212> DNA
<213> Artificial Sequence
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<220>
<223> synthetically generated oligonucleotide

<400>	55						
atggccaagt	acctggagct	cgaggagggc	ggggtcataa	tgcaggcggt	ctactggggac		60
gtgccttcag	gaggaatatg	gtgggacaca	atacggcaga	agataccgga	gtggtacgat		120
gccggaatct	ccgcaatatg	gattcccccg	gcgagcaagg	gcatgggcgg	cgccatttcg		180
atgggctacg	accctctacga	ctcttttgac	ctcggtgagt	acgaccagaa	gggaacggtat		240
gagacgcgct	tgtggctccaa	cgaggagctc	gtgaacatga	taaacaccgc	ccacgcctat		300
ggcatgaagg	taatagccga	tatagtcatc	aaccaccgcg	ccggcgggtga	cctggagtggt		360
aaccccttcg	tgaacgacta	tacctggacc	gacttctcaa	aggctcgcgtc	gggtaaatac		420
acggccaact	acctcgactt	ccacccgaac	gagctccatg	cgggcgattc	cggaacattt		480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctggggccagc		540
caggagagct	acgcggcata	tctcaggagc	atcggcatcg	atgcctggcg	ctttgactac		600
gtgaaggggct	acggagcgtg	ggtcgtcaag	gacttggtca	actggtgggg	cggctggggcc		660
gttggcgagt	actgggacac	caacgttgat	gcactcctca	actgggccta	ctcgagcggc		720
gccaaaggct	tcgacttccc	gctctactac	aagatggatg	aggcctttga	caacaaaaac		780
attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgacccggtt		840
aaggccgtaa	cccttgtagc	aaaccacgac	accgataata	tctggaacaa	gtaccttgct		900
tatgctttca	tctctaccata	cgaaggccag	cccgtcatat	tctaccgcga	ctacgaggag		960
tggctcaaca	aggacagggt	gaacaacctc	atatggatac	acgaccacct	cgcagggtgga		1020
agcacgagca	tagtttacta	cgacagcgac	gagatgatct	tcgtgaggaa	cggctatgga		1080
agcaagcctg	gccttataac	ttacatcaac	ctcggctcga	gcaagggttg	aagggtgggt		1140
tacgttccga	agtttcgagg	ctcgtgcata	cacgagtaca	cgggcaatct	cggcggctgg		1200
gtggacaagt	gggtggactc	aagcggctgg	gtctacctcg	aggctctctg	ccacgaccgc		1260
gtcaacggcc	agttacggcta	ctccgctcgg	agctatttcg	gtgttggtcg	a		1311

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<210> 56
<211> 436
<212> PRT
<213> Artificial Sequence
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$\langle 220 \rangle$

1001827087_1.txt

<223> Synthetically generated peptide

<400> 56

Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala
 1 5 10 15
 Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Lys Asn Ile Pro Ala Leu Val Ser Ala Leu Gln Asn Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His
 325 330 335
 Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
 405 410 415
 Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
 420 425 430
 Cys Gly Val Gly
 435

<210> 57

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

1001827087_1.txt

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<400> 57
atggccaagt acctggagct cgaagagagc ggggtcataa tgcaggcggt ctactgggac 60
gtgccttcag gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgat 120
gccggaatct ccgcaatatg gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
atgggctacg acccctacga ttattttgac ctcggtgagt actaccagaa gggaacgggtg 240
gaaacgaggt tcggtcctaa gcaggagctc ataaacatga taaacaccgc ccacgcctac 300
ggcatcaagg tcatcgacga catagtaatc aaccaccgcg ccggaggaga ctttgagtgg 360
aacccttcg tcaatgacta cacctggacg gacttctcga aggtcgcttc cggcaagtac 420
acggccaact acctcgactt ccacccaac gaggtcaagt gctgtgacga gggcacattt 480
ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggcgagc 540
gatgagagct acgccgccta cctaaggagc atcggcggtg atgcctggcg ctttgactac 600
gtgaagggtc acggagcgtg ggtcgtcaag gactggctca actggtgggg tggctgggcc 660
gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat 720
gcaaaagtct tcgacttccc gctctactac aagatggacg aggccttcga taacaacaac 780
attcccgcgc tgggtggacg cctcagatac ggtcagacag tggtcagccg cgaccggttc 840
aaggctgtga cgtttgtagc caaccacgat accgatataa tctggaacaa gtaccttgct 900
tatgctttca tcctcaccta cgaaggccag cccgtcatat tctaccgga ctacgaggag 960
tggctcaaca aggacaggtt gaacaacctc atatggatac acgaccacct cgcagggtga 1020
agcactgaca tcgtttacta cgacaacgac gagctgatat tcgtgagaaa cggctacgga 1080
agcaagccgg gactgataac atacatcaac ctcgcctcaa gcaaagccgg aaggtgggtc 1140
tacgttccga agttcgcggt agcgtgcatc cacgagtaca ccggcaacct cggcggctgg 1200
gtggacaagt ggggtggactc aagcgggtgg gtgtacctcg aggccctgc ccacgacctg 1260
gccaacggct attacggcta ctccgtctgg agctattgct gtgttggtcg a 1311

```

<210> 58

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 58

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Met Ala Lys Tyr Leu Glu Leu Glu Glu Ser Gly Val Ile Met Gln Ala
1      5      10      15
Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
20     25     30
Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
35     40     45
Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
50     55     60
Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
65     70     75     80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
85     90     95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100    105    110
Arg Ala Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
115    120    125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130    135    140
Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
145    150    155    160
Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp
165    170    175
Leu Trp Ala Ser Asp Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
180    185    190
Val Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
195    200    205
Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
210    215    220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Asp
225    230    235    240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
245    250    255
Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln
260    265    270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn

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      275      280      285
His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Leu Ala Tyr Ala Phe Ile
 290      300
Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu
 305      310      315
Trp Leu Asn Lys Asp Arg Leu Asn Asn Leu Ile Trp Ile His Asp His
      325      330      335
Leu Ala Gly Gly Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu
      340      345      350
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
      355      360      365
Ile Asn Leu Ala Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys
      370      375      380
Phe Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385      390      395
Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
      400      405      410      415
Ala His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val Trp Ser Tyr
      420      425      430
Cys Gly Val Gly
      435

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<210> 59
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

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<400> 59
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gtgccttcag gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgat      120
gccggaatct ccgcaatat gattcctccc gcgagcaagg gtatgagcgg cggctattcg      180
atgggctacg acccctacga ttattttgac ctcggtgagt actaccagaa gggaacggtg      240
gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacaccgc ccacgcctac      300
ggcatcaagg tcatcgaga catagtaatc aaccaccgcg ccggaggaga ccttgagtgg      360
aacccttcg tcaatgacta cacctggacg gacttctcga aggtcgcttc cggcaagtac      420
acggccaact acctcgactt ccaccgaac gagctccatg cgggcgattc cggaacattt      480
ggaggctatc ccgacatat ccacgacaag agctgggacc agtactggct ctgggccagc      540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac      600
gtcaagggct atgctccctg ggtcgtcaag gactggctga actggtgggg aggctgggcg      660
gttgagaggt actgggacac caacgtcgac gctgttctca actgggcata ctcgagcggg      720
gccaaggtct ttgacttcgc cctctactac aagatggacg aggccttcga taacaacaac      780
attcccgccc tgggtggacgc cctcagatac ggtcagacag tggtcagccg cgaccgcttc      840
aaggctgtga cgttttagc caaccacgat accgatataa tttggaacaa gtacccggcc      900
tacgccttca tcctcaccta cgagggccag ccgacgatat tctaccgcga ctacgaggag      960
tggtcaaca aggacaggct caagaacctc atctggatac acgaccacct cgccggtgga      1020
agcactgaca tcgtttacta cgacaacgac gactgtgatc tcgtgagaaa cggctacgga      1080
agcaagccgg gactgataac atacatcaac ctcgcgtcaa gcaaagccgg aagggtgggtt      1140
tatgtgccga agttcgcggt cgcggtgcac cagcagatata ctggtaacct cggaggctgg      1200
gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgaccct      1260
gccaacgggc agtatggcta ctccgtgtgg agctattgcg gtgttggttg a      1311

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<210> 60
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

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<400> 60
Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala
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Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
      20      25      30
Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile

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Pro 35 40 45
Pro 50 Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
65 Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
Glu Thr Arg Phe Gly 70 Ser Lys Gln Glu Leu 90 Ile Asn Met Ile Asn Thr
Ala His Ala Tyr 85 Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
Arg Ala Gly Gly Asp Leu Glu Trp 105 Asn Pro Phe Val Asn Asp Tyr Thr
Trp Thr Asp Phe Ser Lys Val 120 Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
130 Asp Phe His Pro Asn 135 Glu Leu His Ala Gly Asp Ser Gly Thr Phe
145 Gly Gly Tyr Pro Asp 150 Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
Leu Trp Ala Ser 165 Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
Ile Asp Ala Trp Arg Phe Asp Tyr 185 Val Lys Gly Tyr Ala Pro Trp Val
Val Lys Asp Trp Leu Asn Trp 200 Trp Gly Gly Trp Ala Val Gly Glu Tyr
Trp 210 Asp Thr Asn Val Asp 215 Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
225 Ala Lys Val Phe Asp 230 Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln
Thr Val Val Ser Arg Asp Pro Phe 265 Lys Ala Val Thr Phe Val Ala Asn
His Asp Thr Asp Ile Ile Trp 280 Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
Trp Leu Asn Lys Asp 310 Arg Leu Lys Asn Leu Ile Trp Ile His Asp His
Leu Ala Gly Gly Ser Thr Asp Ile Val 330 Tyr Tyr Asp Asn Asp Glu Leu
Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
Ile Asn Leu Ala Ser Ser Lys 360 Ala Gly Arg Trp Val Tyr Val Pro Lys
Phe 370 Ala Gly Ala Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
Val Asp Lys Tyr Val Tyr Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro
Ala Tyr Asp Pro 405 Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr
Cys Gly Val Gly 425 430

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<210> 61

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 61

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gtgccctcag gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgag 120
gcggaatat ccgccatttg gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
atgggctacg acccctacga ttattttgac ctcggtgagt actaccagaa gggaacggtg 240
gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacaccgc ccacgcctac 300
ggcatcaagg tcatcgcaga catagtaatc aaccaccgcg ccggaggaga ccttgagtgg 360
aacccttcg tcaatgacta cacctggacg gacttctcga aggtcgcttc cggcaagtac 420

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acggccaact	acctcaactt	ccacccgaac	gagctccatg	cgggcgattc	cggaacattt	480
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caggagagct	acgcggcata	tctcaggagc	atcggcatcg	atgcctggcg	cttcgactac	600
gtcaagggct	acggagcgtg	ggctgtcaag	gactggctgg	actgggtggg	aggctgggccc	660
gtcggggagt	actgggacac	aaacgttgat	gcactgctca	actgggccta	ctcgagcgat	720
gcaaaagtct	tcgacttccc	gctctactac	aagatggatg	aggcctttga	caacaaaaaac	780
attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgaccctgttc	840
aaggccgtaa	cctttgtagc	aaaccatgac	accgatataa	tctggaacaa	gtatccagcc	900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga	1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg	1080
gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aaggtggggtc	1140
tacgttccga	agttcgcggg	agcgtgcatc	cacgagtaca	ccggcaacct	cggcggctgg	1200
gtggacaagt	gggtggactc	aagcgggtgg	gtgtacctcg	aggccccctgc	ccacgaccgc	1260
gccaacggct	attacggcta	ctccgtctgg	agctactgcg	gggtgggctg	a	1311

<210> 62

<211> 436

<212> PRT

<213> Artificial Sequence

<220>

<223> Synthetically generated peptide

<400> 62

Met	Ala	Lys	Tyr	Ser	Glu	Leu	Lys	Lys	Gly	Gly	Val	Ile	Met	Gln	Ala
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Phe	Tyr	Trp	Asp	Val	Pro	Ser	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg
			20					25					30		
Gln	Lys	Ile	Pro	Glu	Trp	Tyr	Glu	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile
		35					40					45			
Pro	Pro	Ala	Ser	Lys	Gly	Met	Ser	Gly	Gly	Tyr	Ser	Met	Gly	Tyr	Asp
	50					55					60				
Pro	Tyr	Asp	Tyr	Phe	Asp	Leu	Gly	Glu	Tyr	Tyr	Gln	Lys	Gly	Thr	Val
65					70					75					80
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Ile	Asn	Met	Ile	Asn	Thr
				85					90					95	
Ala	His	Ala	Tyr	Gly	Ile	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
			100					105					110		
Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr	Thr
		115					120					125			
Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
	130					135					140				
Leu	Asn	Phe	His	Pro	Asn	Glu	Leu	His	Ala	Gly	Asp	Ser	Gly	Thr	Phe
145					150					155					160
Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	Asp	Lys	Ser	Trp	Asp	Gln	Tyr	Trp
				165					170					175	
Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Tyr	Leu	Arg	Ser	Ile	Gly	
			180					185					190		
Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val
		195					200					205			
Val	Lys	Asp	Trp	Leu	Asp	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr
	210					215					220				
Trp	Asp	Thr	Asn	Val	Asp	Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Asp
225					230					235					240
Ala	Lys	Val	Phe	Asp	Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
				245					250					255	
Asp	Asn	Lys	Asn	Ile	Pro	Ala	Leu	Val	Ser	Ala	Leu	Gln	Asn	Gly	Gln
			260					265					270		
Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
		275					280					285			
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
	290					295					300				
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
305					310					315					320
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	Asn
				325					330					335	
Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu

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Ile	Phe	Val	340	Asn	Gly	Tyr	Gly	345	Asp	Lys	Pro	Gly	Leu	350	Ile	Thr	Tyr
		355					360						365				
Ile	Asn	Leu	Gly	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys		
	370					375					380						
Phe	Ala	Gly	Ala	Cys	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp		
385				390						395					400		
Val	Asp	Lys	Trp	Val	Asp	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro		
				405					410					415			
Ala	His	Asp	Pro	Ala	Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr		
			420					425					430				
Cys	Gly	Val	Gly														
		435															

<210> 63
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 63																	
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gtgcccttcag	gaggaatatg	gtgggacaca	atacggcaga	agataccgga	gtggtacgat												120
gccggaatct	ccgcaatatg	gattcccccg	gcgagcaagg	gcatgggdcg	cgccatttcg												180
atgggctacg	acccctacga	cttctttgac	ctcgggtgagt	acgaccagaa	gggaacggta												240
gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacacggc	ccatgcctac												300
ggcataaagg	ccatagcgga	catcgtcata	aaccaccgcg	caggcggaga	cctcgagtgg												360
aacccgttcg	ttggggacta	cacctggacg	gacttctcaa	aggtggcctc	gggcaaatat												420
actgccaact	acctcgactt	ccaccccaac	gaggtcaagt	gctgtgacga	gggcacattt												480
ggaggcttcc	cagacatagc	ccacgagaag	agctgggacc	agcactggct	ctgggdcgagc												540
gatgagagct	acgccgccta	cctaaggagc	atcggcggtg	atgcctggcg	ctttgactac												600
gtgaagggct	acggagcggt	ggtcgtcaag	gactggctca	actggtgggg	cggctgggccc												660
gttggcgagt	actgggacac	caacgttgat	gcactcctca	actgggccta	ctcgagcggc												720
gccaaggtct	tcgacttccc	gctctactac	aagatggacg	cggcctttga	caacaagaac												780
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aaggccgtaa	cttctgttgc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc												900
tacgcgttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag												960
tggtcaaca	aggataagct	caagaacctc	atctggatac	atgacaacct	cgccggagga												1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg												1080
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gtagacaagt	acgtctactc	aagcggctgg	gtctatctcg	aagctccagc	ttacgacctt												1260
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<210> 64
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

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			20				25					30					
Gln	Lys	Ile	Pro	Glu	Trp	Tyr	Asp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile		
		35				40					45						
Pro	Pro	Ala	Ser	Lys	Gly	Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr	Asp		
		50				55					60						
Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	Glu	Tyr	Asp	Gln	Lys	Gly	Thr	Val		
65				70					75					80			
Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr		
			85					90					95				
Ala	His	Ala	Tyr	Gly	Ile	Lys	Ala	Ile	Ala	Asp	Ile	Val	Ile	Asn	His		

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Arg	Ala	Gly	100	Gly	Asp	Leu	Glu	Trp	105	Asn	Pro	Phe	Val	Gly	110	Asp	Tyr	Thr
Trp	Thr	Asp	115	Phe	Ser	Lys	Val	Ala	120	Ser	Gly	Lys	Tyr	Thr	125	Ala	Asn	Tyr
Leu	Asp	Phe	130	His	Pro	Asn	Glu	Val	135	Lys	Cys	Cys	Asp	Glu	140	Gly	Thr	Phe
Gly	Gly	Phe	145	Pro	Asp	Ile	Ala	His	150	Glu	Lys	Ser	Trp	Asp	155	Gln	His	Trp
Leu	Trp	Ala	160	Ser	Asp	Glu	Ser	Tyr	165	Ala	Ala	Tyr	Leu	Arg	170	Ser	Ile	Gly
Val	Asp	Ala	175	Trp	Arg	Phe	Asp	Tyr	180	Val	Lys	Gly	Tyr	Gly	185	Ala	Trp	Val
Val	Lys	Asp	190	Trp	Leu	Asn	Trp	Gly	195	Gly	Gly	Trp	Ala	Val	200	Gly	Glu	Tyr
Trp	Asp	Thr	205	Asn	Val	Asp	Ala	Leu	210	Leu	Asn	Trp	Ala	Tyr	215	Ser	Ser	Gly
Ala	Lys	Val	220	Phe	Asp	Phe	Pro	Leu	225	Tyr	Tyr	Lys	Met	Asp	230	Ala	Ala	Phe
Asp	Asn	Lys	235	Asn	Ile	Pro	Ala	Leu	240	Val	Glu	Ala	Leu	Lys	245	Asn	Gly	Gly
Thr	Val	Val	250	Ser	Arg	Asp	Pro	Phe	255	Lys	Ala	Val	Thr	Phe	260	Val	Ala	Asn
His	Asp	Thr	265	Asp	Ile	Ile	Trp	Asn	270	Lys	Tyr	Pro	Ala	Tyr	275	Ala	Phe	Ile
Leu	Thr	Tyr	280	Glu	Gly	Gln	Pro	Thr	285	Ile	Phe	Tyr	Arg	Asp	290	Tyr	Glu	Glu
Trp	Leu	Asn	295	Lys	Asp	Lys	Leu	Lys	300	Asn	Leu	Ile	Trp	Ile	305	His	Asp	Asn
Leu	Ala	Gly	310	Gly	Ser	Thr	Asp	Ile	315	Val	Tyr	Tyr	Asp	Asn	320	Asp	Glu	Leu
Ile	Phe	Val	325	Arg	Asn	Gly	Tyr	Gly	330	Asp	Lys	Pro	Gly	Leu	335	Ile	Thr	Tyr
Ile	Asn	Leu	340	Gly	Trp	Ser	Lys	Ala	345	Gly	Arg	Trp	Val	Tyr	350	Val	Pro	Lys
Phe	Ala	Gly	355	Ala	Cys	Ile	His	Glu	360	Tyr	Thr	Gly	Asn	Leu	365	Gly	Gly	Trp
Val	Asp	Lys	370	Tyr	Val	Tyr	Ser	Ser	375	Gly	Trp	Val	Tyr	Leu	380	Glu	Ala	Pro
Ala	Tyr	Asp	385	Pro	Ala	Asn	Gly	Gln	390	Tyr	Gly	Tyr	Ser	Val	395	Trp	Ser	Tyr
Cys	Gly	Val	400	Gly					405						410			
			415						420						425			
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<210> 65

<211> 1311

<212> DNA

<213> Artificial Sequence

<220>

<223> Synthetically generated oligonucleotide

<400> 65

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gcgggaatat	ccgccatttg	gattcctccc	gcgagcaagg	gtatgagcgg	cggctatttcg	180
atgggctacg	acccctacga	ttattttgac	ctcggtagt	actaccagaa	gggaacgggtg	240
gaaacgaggt	tcggctcaaa	gcaggagctc	ataaacatga	taaacaccgc	ccacgcctat	300
ggcatgaagg	taatagccga	tatagtcatc	aaccaccgcg	cggcggtga	cctggagtg	360
aaccccttcg	tgaacgacta	tacctggacc	gacttctcaa	aggctcgcgc	gggtaaatac	420
acggccaact	acctcgactt	ccaccggaac	gagctccatg	cgggcgattc	cggaacattt	480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctgggccagc	540
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gtcaagggt	atgctccctg	ggctgtcaag	gactggctga	actggtgggg	aggctggg	660
gttgagaggt	actgggacac	caacgtcgac	gctgttctca	actgggcata	ctcgagcgg	720
gccaaaggtct	ttgacttcgc	cctctactac	aagatggacg	aggccttcga	taacaacaac	780
attccgccc	tggtggacgc	cctcagatac	ggtcagacag	tggtcagccg	cgaccggttc	840
aaggctgtga	cgttttagtc	caaccacgat	accgatataa	tttgaacaa	gtaccggcc	900

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tacgccttca	tcctcaccta	cgagggccag	ccgacgatat	tctaccgcga	ctacgaggag	960
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agcacgagca	tagtttacta	cgacagcgac	gagatgatct	tcgtgaggaa	cggctatgga	1080
agcaagcctg	gccttataac	ttacatcaac	ctcggctcga	gcaagggttg	aaggtgggtt	1140
tacgttccga	agttcgcagg	ctcgtgcata	cacgagtaca	ccggcaatct	cggcggctgg	1200
gtggacaagt	gggtggactc	aagcggctgg	gtctacctcg	aggctcctgc	ccacgaccgc	1260
gccaacggcc	agtacggcta	ctccgtctgg	agctattgca	gtgttggtcg	a	1311

<210> 66
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> synthetically generated peptide

<400> 66
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 Phe Tyr Trp Asp Val Pro Gly Gly Ile Trp Trp Gly Thr Ile Arg
 20 25 30
 Ser Lys Ile Pro Glu Trp Tyr Glu Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe
 145 150 155 160
 Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp
 165 170 175
 Leu Trp Ala Ser Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
 180 185 190
 Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val
 195 200 205
 Val Lys Asp Trp Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr
 210 215 220
 Trp Asp Thr Asn Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly
 225 230 235 240
 Ala Lys Val Phe Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe
 245 250 255
 Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln
 260 265 270
 Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
 275 280 285
 His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile
 290 295 300
 Leu Thr Tyr Glu Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu
 305 310 315 320
 Trp Leu Asn Lys Asp Arg Leu Lys Asn Leu Ile Trp Ile His Asp His
 325 330 335
 Leu Ala Gly Gly Ser Thr Ser Ile Val Tyr Tyr Asp Ser Asp Glu Met
 340 345 350
 Ile Phe Val Arg Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr
 355 360 365
 Ile Asn Leu Gly Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys
 370 375 380
 phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp
 385 390 395 400
 Val Asp Lys Trp Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro

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Ala	His	Asp	Pro	405	Ala	Asn	Gly	Gln	Tyr	410	Gly	Tyr	Ser	Val	Trp	415	Ser	Tyr
			420						425						430			
Cys	Gly	Val	Gly	435														

<210> 67
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated oligonucleotide

<400> 67
 atggccaagt acctggagct cgaagagggc ggggtcataa tgcaggcggt ctactgggac 60
 gtgccttcgg gaggaatat gtgggacaca atacggcaga agataccgga gtggtacgat 120
 gccggaatct ccgcaatat gattcctccc gcgagcaagg gtatgagcgg cggctattcg 180
 atgggctacg acccctacga ttatcttgac ctcggtgagt actaccagaa gggaacgggtg 240
 gaaacgaggt tcggctcaaa gcaggagctc ataaacatga taaacacggc ccatgcctac 300
 ggcataaagg tcatagcgga catcgtcata aaccaccgag caggcggaga cctcgagtgg 360
 aacccgttcg ttggggacta cacctggacg gacttctcaa aggtggcctc gggcaaatat 420
 actgccaact acctcgactt ccaccccaac gaggtcaagt gctgtgacga gggcacattt 480
 ggaggcttcc cagacatagc ccacgagaag agctgggacc agcactggct ctgggagcagc 540
 gatgagagct acgccgccta cctaaggagc atcggcggtg atgcctggcg cttcgactac 600
 gtcaagggct acggagcgtg ggtcgtcaag gactggctgg actggtgggg aggctggggc 660
 gtcggggagt actgggacac aaacgttgat gcactgctca actgggccta ctcgagcgat 720
 gcaaaagtct tcgacttccc gctctactac aagatggacg aggccttcga taacaacaac 780
 attcccggcc tgggtggacgc cctcagatac ggtagacag tggtagccg cgaccggtc 840
 aaggctgtga cgttttagc caaccacgat accgatataa tctggaacaa gtatccagcc 900
 tacgcgttca tcctcaccta cgagggccag ccgacaatat tctaccgca ctacgaggag 960
 tggctcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
 agcacgagca tagtttacta cgacagcgac gagatgatct tcgtgaggaa cggctatgga 1080
 agcaagcctg gccttataac ttacatcaac ctcggctcga gcaaggttgg aaggtgggtc 1140
 tacgttccga agttcgcggg agcgtgcata cagagtagca ccggcaacct cggcggctgg 1200
 gtggacaagt ggggtggactc aagcgggtgg gtgtacctcg aggccctgc ccacgacctg 1260
 gccaacggct attacggcta ctccgtctgg agctactgag tgggtgggctg a 1311

<210> 68
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Synthetically generated peptide

<400> 68
 Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala
 1 5 10 15
 Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
 20 25 30
 Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
 35 40 45
 Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp
 50 55 60
 Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val
 65 70 75 80
 Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Ile Asn Met Ile Asn Thr
 85 90 95
 Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
 100 105 110
 Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr
 115 120 125
 Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr
 130 135 140
 Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly Thr Phe
 145 150 155 160
 Gly Gly Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln His Trp

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Leu	Trp	Ala	Ser	165	Glu	Ser	Tyr	Ala	170	Tyr	Leu	Arg	Ser	175	Ile	Gly	
Val	Asp	Ala	180	Trp	Arg	Phe	Asp	Tyr	185	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val
Val	Lys	Asp	195	Trp	Leu	Asp	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	
Trp	Asp	Thr	Asn	Val	Asp	Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Asp		
Ala	Lys	Val	Phe	Asp	230	Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	
Asp	Asn	Asn	Asn	245	Ile	Pro	Ala	Leu	Val	Asp	Ala	Leu	Arg	Tyr	Gly	Gln	
Thr	Val	Val	Ser	260	Arg	Asp	Pro	Phe	265	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile		
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu		
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	Asn		
Leu	Ala	Gly	Gly	Ser	Thr	Ser	Ile	Val	Tyr	Tyr	Asp	Ser	Asp	Glu	Met		
Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	Ser	Lys	Pro	Gly	Leu	Ile	Thr	Tyr		
Ile	Asn	Leu	Gly	Ser	Ser	Lys	Val	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys		
Phe	Ala	Gly	Ala	Cys	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp		
Val	Asp	Lys	Trp	Val	Asp	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro		
Ala	His	Asp	Pro	Ala	Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr		
Cys	Val	Val	Gly														

<210> 69
 <211> 1542
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 69																	
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gagggaaata	aggcagaagc	agcaacagtg	aacaatggaa	cattaatgca	gtattttgag												120
tggtacgctc	cgaatgatgg	gaatcattgg	aatcgtttgc	gttccgatgc	tgaaagttta												180
gctcataaa	gaatcacatc	tgtatggata	ccacctgcac	ataaaggac	ttcgcaaaat												240
gatgtagggt	atggggccta	tgatttatat	gatttagggg	agttcaatca	aaaaggaacg												300
gtgcggacga	aatatgggac	aaaagcacag	ttgaaatctg	caattgacgc	tttacataag												360
caaaacatcg	acgtatacgg	tgatgtagtt	atgaatcata	aagggtggggc	tgattatact												420
gaaaccgtaa	cagctgttga	ggtagaccgt	aacaatcgaa	atattgaagt	atcaggtgat												480
tatcaaatta	gtgcatggac	ggggtttaat	tttccagggc	gcgagatgc	ttatttctaat												540
ttcaaatgga	aatggtatca	ttttgacgga	acggattggg	atgaaggaag	gaaattaaat												600
cgaatttata	aattttagggg	tgtagataaa	gcgtgggatt	gggaagtgtc	tagcgaaaat												660
ggaaattatg	attattttgat	gtatgcagat	cttgattttg	atcatcctga	tgttgcgaat												720
gagatgaaaa	attggggaac	atggtatgcg	aatgaattaa	atttagatgg	ctttcgtttg												780
gacgctgtta	aacatattga	tcatgaatat	ttacgcgatt	gggtaaatca	tgccagacag												840
caaacgggga	aagaaatggt	tacagtagct	gaatattggc	aaaatgatgt	tcaggcttta												900
aacaattatt	tagcgaaagt	caattataat	caatctgtgt	ttgatgcacc	gcttcattac												960
aatttttcatt	atgcttcaac	aggaaatggg	aattatgata	tgagaaaata	tttaaatgga												1020
acagtaatga	aaaatcacc	tgactcgc	gttactctcg	ttgagaatca	tgattctcag												1080
cctgggcagt	cattggaatc	tgtagtaagt	ccgtggttta	agccgctggc	atatgcattt												1140
attttaactc	gtgcagaggg	ctatccttca	gttttctatg	gtgattacta	tgggacaagc												1200
ggaaaatagta	gttatgaaat	tccagcgtaa	aaagataaaa	ttgatccaat	tttgacggca												1260
cgaaaaaact	ttgcataatg	tacgcagcgt	gattatttag	accatccaga	tgtgattggc												1320
tggacaagag	aaggcgatgg	tgtacatgct	aattctggtt	tagcgacatt	actctcggac												1380

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ggaccaggag gatcaaagt gatggatgtt ggaagaata acgctgggga agtatggtag 1440
gatattacgg gtaatcaaac aaatactgta acaattaata aggacggatg ggggcagttc 1500
tatgtaagtg gcggctcagt ttccatatat gttcagcggg aa 1542

<210> 70
<211> 513
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 70
Met Leu Lys Arg Ile Thr Val Val Cys Leu Leu Phe Ile Leu Leu Phe
1 5 10 15
Pro Asn Ile Tyr Glu Gly Asn Lys Ala Glu Ala Ala Thr Val Asn Asn
20 25 30
Gly Thr Leu Met Gln Tyr Phe Glu Trp Tyr Ala Pro Asn Asp Gly Asn
35 40 45
His Trp Asn Arg Leu Arg Ser Asp Ala Glu Ser Leu Ala His Lys Gly
50 55 60
Ile Thr Ser Val Trp Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Asn
65 70 75 80
Asp Val Gly Tyr Gly Ala Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn
85 90 95
Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Leu Lys
100 105 110
Ser Ala Ile Asp Ala Leu His Lys Gln Asn Ile Asp Val Tyr Gly Asp
115 120 125
Val Val Met Asn His Lys Gly Gly Ala Asp Tyr Thr Glu Thr Val Thr
130 135 140
Ala Val Glu Val Asp Arg Asn Asn Arg Asn Ile Glu Val Ser Gly Asp
145 150 155 160
Tyr Gln Ile Ser Ala Trp Thr Gly Phe Asn Phe Pro Gly Arg Gly Asp
165 170 175 180
Ala Tyr Ser Asn Phe Lys Trp Lys Trp Tyr His Phe Asp Gly Thr Asp
185 190 195
Trp Asp Glu Gly Arg Lys Leu Asn Arg Ile Tyr Lys Phe Arg Gly Val
200 205 210
Asp Lys Ala Trp Asp Trp Glu Val Ser Ser Glu Asn Gly Asn Tyr Asp
215 220 225
Tyr Leu Met Tyr Ala Asp Leu Asp Phe Asp His Pro Asp Val Ala Asn
230 235 240
Glu Met Lys Asn Trp Gly Thr Trp Tyr Ala Asn Glu Leu Asn Leu Asp
245 250 255
Gly Phe Arg Leu Asp Ala Val Lys His Ile Asp His Glu Tyr Leu Arg
260 265 270
Asp Trp Val Asn His Ala Arg Gln Gln Thr Gly Lys Glu Met Phe Thr
275 280 285
Val Ala Glu Tyr Trp Gln Asn Asp Val Gln Ala Leu Asn Asn Tyr Leu
290 295 300
Ala Lys Val Asn Tyr Asn Gln Ser Val Phe Asp Ala Pro Leu His Tyr
305 310 315 320
Asn Phe His Tyr Ala Ser Thr Gly Asn Gly Asn Tyr Asp Met Arg Asn
325 330 335
Ile Leu Asn Gly Thr Val Met Lys Asn His Pro Ala Leu Ala Val Thr
340 345 350
Leu Val Glu Asn His Asp Ser Gln Pro Gly Gln Ser Leu Glu Ser Val
355 360 365
Val Ser Pro Trp Phe Lys Pro Leu Ala Tyr Ala Phe Ile Leu Thr Arg
370 375 380
Ala Glu Gly Tyr Pro Ser Val Phe Tyr Gly Asp Tyr Tyr Gly Thr Ser
385 390 395 400
Gly Asn Ser Ser Tyr Glu Ile Pro Ala Leu Lys Asp Lys Ile Asp Pro
405 410 415
Ile Leu Thr Ala Arg Lys Asn Phe Ala Tyr Gly Thr Gln Arg Asp Tyr
420 425 430
Leu Asp His Pro Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Gly Val

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		435				440			445						
His	Ala	Asn	Ser	Gly	Leu	Ala	Thr	Leu	Leu	Ser	Asp	Gly	Pro	Gly	Gly
	450					455					460				
Ser	Lys	Trp	Met	Asp	Val	Gly	Lys	Asn	Asn	Ala	Gly	Glu	Val	Trp	Tyr
465					470					475					480
Asp	Ile	Thr	Gly	Asn	Gln	Thr	Asn	Thr	Val	Thr	Ile	Asn	Lys	Asp	Gly
				485					490					495	
Trp	Gly	Gln	Phe	Tyr	Val	Ser	Gly	Gly	Ser	Val	Ser	Ile	Tyr	Val	Gln
		500						505					510		
Arg															

<210> 71
<211> 1311
<212> DNA
<213> Artificial Sequence

<220>
<223> Synthetically generated oligonucleotide

<400> 71
atggccaagt acctggagct cgaagagggc ggggtcataa tgcaggcggt ctactgggac 60
gtgccttcag gaggaatatg gtgggacaca atacggcaga agataccgga gtggtacgat 120
gccggaatct ccgcaatatg gattcccccg gcgagcaagg gcatgggcgg cgcctattcg 180
atgggctacg acccctacga cttctttgac ctcggtgagt acgaccagaa gggaacggta 240
gagacgcgct ttggctccaa gcaggagctc gtgaacatga taaacacggc ccatgcctac 300
ggcataaagg tcatagcgga catcgtcata aaccaccgag caggcggaga cctcgagtgg 360
aaccgcgttc ttggggacta cacctggacg gactttctca aggtagcctc gggcaaatat 420
actgccaaat acctcgactt ccacccgaac gagctccatg cgggcgattc cggaacattt 480
ggaggctatc ccgacatatg ccacgacaag agctgggacc agtactggct ctggggccagc 540
caggagagct acgcggcata tctcaggagc atcggcatcg atgcctggcg cttcgactac 600
gtcaagggct atgctccctg ggtcgtcaag gactggctga actggtgggg aggcctggcg 660
gttggaaggt actgggacac caacgtcgag gctgttctca actgggcata ctcgagcggg 720
gccaaggtct ttgacttcgc cctctactac aagatggatg aggcctttga caacaaaaac 780
attccagcgc tcgtctctgc ccttcagaac ggccagactg ttgtctcccg cgacccttc 840
aaggccgtaa cttttgtagc aaaccacgac accgatataa tctggaacaa gtatccagcc 900
tacgcgttca tcttcaccta cgagggccag ccgacaatat tctaccgcca ctacgaggag 960
tggctcaaca aggataagct caagaacctc atctggatac atgacaacct cgccggagga 1020
agcactgaca tcgtttacta cgacaacgac gagctgatat tcgtgagaaa cggctacgga 1080
agcaagccgg gactgataac atacatcaac ctgcctctca gcaaagccgg aaggtggggt 1140
tatgtgccga agttcgcggg cgctgcatc cagcagatata ctggtaacct cgagggtgg 1200
gtagacaagt acgtctactc aagcggctgg gtctatctcg aagctccagc ttacgaccct 1260
gccaacgggc agtatggcta ctccgtgtgg agctactgag ggggtgggctg a 1311

<210> 72
<211> 436
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetically generated peptide

<400> 72
Met Ala Lys Tyr Leu Glu Leu Glu Gly Gly Val Ile Met Gln Ala
1 5 10 15
Phe Tyr Trp Asp Val Pro Ser Gly Gly Ile Trp Trp Asp Thr Ile Arg
20 25 30
Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
35 40 45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
50 55 60
Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Asp Gln Lys Gly Thr Val
65 70 75 80
Glu Thr Arg Phe Gly Ser Lys Gln Glu Leu Val Asn Met Ile Asn Thr
85 90 95
Ala His Ala Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
100 105 110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Gly Asp Tyr Thr

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[illegible]

<210>	73
<211>	1299
<212>	DNA
<213>	Unknown

<220>
<223> Obtained from an environmental sample.

<400>	73						
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ggaatctggt	gggacaccat	agcccagaag	ataccgcact	gggcgagcgc	cgggatttcg		120
gcaatatgga	ttcctcccgc	gagtaagggc	atgagcggcg	gctattctga	gggctacgac		180
ccctacgatt	tcttcgacct	cggtagtatc	taccagaagg	gaagcgttga	gaccgccttc		240
ggatcaaaag	aggagcttgt	gaacatgata	aacaccgccc	atgctcaca	catgaaggtc		300
atagcggaca	tagtcatcaa	ccaccgcgcc	ggcggcgacc	tggagtggaa	tcctttccacc		360
aacagctaca	cctggaccga	tttctcgaag	gtcgcgtcgg	gcaagtacac	ggccaactac		420
ctcgacttcc	acccgaacga	gcttcacgcg	ggcgattccg	gaacattttg	aggctatccc		480
gacatatgcc	acgacaagag	ctgggaccag	cactggctct	gggccagcaa	cgaaggctac		540
gccgcctacc	tccggagcat	cggcatcgac	gcctggcgct	tcgactacgt	caagggtctac		600
gctccctggg	tcgttaaagaa	ctggctgaac	cggtggggcg	gctgggcggt	tggagagtac		660
tgggacacca	acgtcgaatg	actcctgagc	tgggcctacg	acagcgggtg	taaagtcttc		720
gacttccgcg	tctactacaa	gatggacgag	gccttcgata	acaacaacat	ccccgccttc		780
gtggacgccc	tcaagaacgg	aggcacggtc	gtcagccgcg	acccgttcaa	agccgtgacc		840
ttcgtttgca	accacgatac	caacataatc	tggacaaga	atccggccta	cgccttcata		900
ctcactatga	agggacagcc	ggcaatatct	taccgcact	acgaggagt	gctcaacaag		960
gacaggctca	ggaacctcat	ctggatacac	gaccacctcg	cgggaggaag	cacagacatc		1020

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atctactacg	acagcgacga	gcttatcttc	gtgagaaacg	gctacgggga	caagccggga	1080
ctgataacct	acatcaacct	cggctcaagc	aaggccggaa	ggtgggtcta	cgttccgaag	1140
ttcgaggct	cgtgcataca	cgagtacacc	ggcaacctcg	gcggctggat	tgacaagtgg	1200
ggtgactcaa	gcggtcgggt	ctaccttgag	gccccgcgc	acgaccggc	caacggccag	1260
tacggctact	ccgtatggag	ctactgcggt	gttgggtga			1299

<210> 74
 <211> 432
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 74
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 Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro
 20 25 30
 Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser
 35 40 45
 Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe
 50 55 60
 Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Ser Val Glu Thr Arg Phe
 65 70 75 80
 Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala His Ala His
 85 90 95
 Asn Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 100 105 110
 Asp Leu Glu Trp Asn Pro Phe Thr Asn Ser Tyr Thr Trp Thr Asp Phe
 115 120 125
 Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His
 130 135 140
 Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro
 145 150 155 160
 Asp Ile Cys His Asp Lys Ser Trp Asp Gln His Trp Leu Trp Ala Ser
 165 170 175
 Asn Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp
 180 185 190
 Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Lys Asn Trp
 195 200 205
 Leu Asn Arg Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn
 210 215 220
 Val Asp Ala Leu Leu Ser Trp Ala Tyr Asp Ser Gly Ala Lys Val Phe
 225 230 235 240
 Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Asn Asn
 245 250 255
 Ile Pro Ala Leu Val Asp Ala Leu Lys Asn Gly Gly Thr Val Val Ser
 260 265 270
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asn
 275 280 285
 Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu
 290 295 300
 Gly Gln Pro Ala Ile Phe Tyr Arg Asp Tyr Glu Trp Leu Asn Lys
 305 310 315 320
 Asp Arg Leu Arg Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly
 325 330 335
 Ser Thr Asp Ile Ile Tyr Tyr Asp Ser Asp Glu Leu Ile Phe Val Arg
 340 345 350
 Asn Gly Tyr Gly Asp Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly
 355 360 365
 Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser
 370 375 380
 Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Ile Asp Lys Trp
 385 390 395 400
 Val Asp Ser Ser Gly Arg Val Tyr Leu Glu Ala Pro Ala His Asp Pro
 405 410 415
 Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly

<210> 75
 <211> 1299
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 75
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 ggaatctggg gggacacgat agcccagaag ataccgcgact gggcaagcgc cgggatttcg 120
 gcgatatgga ttccccccgc gagcaagggt atgagcggcg gctattcgat gggctacgac 180
 ccctacgatt attttgacct cggtgagtac taccagaagg gaacgggtgga aacaagattc 240
 ggctcaaagc aggagctcat aaacatgata aacaccgccc acgcctatgg catgaaggta 300
 atagccgata tagtcatcaa ccaccgcgcc ggcggcgatc tggagtggaa ccccttcgtg 360
 aacgactata cctggaccga cttctcgaag gtcgcgtcgg gtaaatacac ggccaactac 420
 ctcgacttcc acccgaacga gctccacgcg ggcgattccg gaacatttgg aggctatccc 480
 gacatatgcc acgacaagag ctgggaccag tactggctct gggccagcca ggagagctac 540
 gcggcctatc tcaggagcat cggcatcgac gcctggcgct tcgactacgt caagggctat 600
 gctccctggg tcgtcagggg ctggctgaac tgggtggggag gctgggcagt tggagagtac 660
 tgggacacca acgtcgacgc tgttctcaac tgggcatact cgagcgggtc caaggtcttt 720
 gacttcgccc tctactacaa gatggacgag gccttcgata acaacaacat tcccgccctg 780
 gtggacgccc tcagatacgg ccagacagtg gtcagccgcg acccgttcaa ggctgtgacg 840
 ttgttagcca accacgatac cgacataatc tggacaagt atccagccta cgcgttcac 900
 ctacactacg agggccagcc gacaatattc taccgcgact acgaggagtg gctcaacaag 960
 gacaagctca agaacctcat ctggatacat gacaacctcg ccggagggag cactgacatc 1020
 gtttactacg acaacgacga gctgatattc gtgagaaacg gctacggaag caagccggga 1080
 ctgataacat acatcaacct cggctcaagc aaagccggaa ggtgggttta cgttccgaag 1140
 ttcgcaggct cgtgcataca cgagtacacc ggcaacctcg gcggctgggt ggacaagtgg 1200
 gtggactcaa gcggctgggt ttacctcgag gctcctgcc acgaccggc caacggccag 1260
 tacggctact ccgtttggag ctattgcggt gttgggtga 1299

<210> 76
 <211> 432
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 76
 Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp
 1 5 10 15
 Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro
 20 25 30
 Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Ala Ser
 35 40 45
 Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr
 50 55 60
 Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe
 65 70 75 80
 Gly Ser Lys Gln Glu Ile Asn Met Ile Asn Thr Ala His Ala Tyr
 85 90 95
 Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 100 105 110
 Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe
 115 120 125
 Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His
 130 135 140
 Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro
 145 150 155 160
 Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser
 165 170 175
 Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp
 180 185 190
 Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Arg Asp Trp

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```

195      200      205
Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn
210      215      220
Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe
225      230      235
Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Asn Asn
245      250      255
Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser
260      265      270
Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp
275      280      285
Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu
290      295      300
Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
305      310      315
Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly
325      330      335
Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg
340      345      350
Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly
355      360      365
Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser
370      375      380
Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp
385      390      395
Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp Pro
405      410      415
Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly
420      425      430

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<210> 77
 <211> 1299
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 77
atggctctgg aagagggcgg gctcataatg caggccttct actgggacgt ccccatggga 60
ggaatctggg gggacacgat agcccagaag ataccgcgact gggcaagcgc cgggatttcg 120
gcgatatgga tccctccgcg gagcaagggt atgagcggcg gctattcgat gggctacgac 180
ccctacgatt attttgacct cgggtgagtac taccagaagg gaacggtgga aacgaggttc 240
ggctcaaaagc aggagctcat aaacatgata aacaccgccc acgcctatgg catgaaggta 300
atagccgata tagtcatcaa ccaccgcgcc ggcggtgacc tggagtggaa ccccttcgtg 360
aacgactata cctggaccga cttctcaaag gtcgcgctcg gtaaatacac ggccaactac 420
ctcgacttcc acccgaacga gctccatgcg ggcgattccg gaacatttgg aggctatccc 480
gacatatgcc acgacaagag ctgggaccag tactggctct gggccagcca ggagagctac 540
gcggcatatc tcaggagcat cggcatcgat gcctggcgct tcgactacgt caagggctat 600
gtccctggg tcgtcaagga ctggctgaac tgggtgggag gctgggcggt tggagagtac 660
tgggacacca acgtcgacgc tgttctcaac tgggcatact cgagcgggtgc caaggtcttt 720
gacttcgccc tctactacaa gatggacgag gccttcgata acaacaacat tcccgccctg 780
gtggacgccc tcagatacgg tcagacagtg gtcagccgcg acccgttcaa ggctgtgacg 840
tttgtagcca accacgatac cgacataatc tggacaagt atccagccta cgcgttcac 900
ctcacctacg agggccagcc gacaatattc taccgcgact acgaggagtg gctcaacaag 960
gataagctca agaacctcat ctggatacat gaccacctcg ccggaggag cactgacatc 1020
gtttactacg acaacgacga gctgatattc gtgagaaacg gctacggaag caagccggga 1080
ctgataacat acatcaacct cgcctcaagc aaagccggaa ggtgggttta cgttccgaag 1140
ttcgaggct cgtgcataca cgagtacacc ggcaatctcg gcggtgggt ggacaagtgg 1200
gtggactcaa gcggctgggt ctacctcgag gctcctgcc acgaccggc caacggccag 1260
tacggctact ccgtctggag ctactgcggt gttgggtga 1299

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<210> 78
 <211> 432
 <212> PRT
 <213> Unknown

<220>

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<223> Obtained from an environmental sample.

<400> 78

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Met Ala Leu Glu Glu Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp
 1      5      10      15
Val Pro Met Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro
 20      25      30
Asp Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Ala Ser
 35      40      45
Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr
 50      55      60
Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe
 65      70      75
Gly Ser Lys Gln Glu Ile Asn Met Ile Asn Thr Ala His Ala Tyr
 85      90      95
Gly Met Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
100      105      110
Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe
115      120      125
Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His
130      135      140
Pro Asn Glu Leu His Ala Gly Asp Ser Gly Thr Phe Gly Gly Tyr Pro
145      150      155
Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser
165      170      175
Gln Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Ile Asp Ala Trp
180      185      190
Arg Phe Asp Tyr Val Lys Gly Tyr Ala Pro Trp Val Val Lys Asp Trp
195      200      205
Leu Asn Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn
210      215      220
Val Asp Ala Val Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe
225      230      235
Asp Phe Ala Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Asn Asn
245      250      255
Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser
260      265      270
Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp
275      280      285
Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu
290      295      300
Gly Gln Pro Thr Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
305      310      315
Asp Lys Leu Lys Asn Leu Ile Trp Ile His Asp Asn Leu Ala Gly Gly
325      330      335
Ser Thr Asp Ile Val Tyr Tyr Asp Asn Asp Glu Leu Ile Phe Val Arg
340      345      350
Asn Gly Tyr Gly Ser Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Ala
355      360      365
Ser Ser Lys Ala Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser
370      375      380
Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Trp
385      390      395
Val Asp Ser Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala His Asp Pro
405      410      415
Ala Asn Gly Gln Tyr Gly Tyr Ser Val Trp Ser Tyr Cys Gly Val Gly
420      425      430

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<210> 79

<211> 1386

<212> DNA

<213> Bacterial

<400> 79

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atgaagcctg cgaaactcct cgtctttgtg ctcgtagtct ctatcctcgc ggggctctac      60
gccagccccg cgggggcggc caagtacctg gagctcgaag agggcggcgt cataatgcag      120
gcgttctact gggacgtgcc ttcaggagga atatggtggg acacaatacg gcagaagata      180
ccggagtggg acgatgccgg aatctccgca atatggattc cccggcgag caagggcag      240

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ggcggcgcct attcgatggg ctacgacccc tacgacttct ttgacctcgg tgagtacgac 300
cagaaggga cggtagagac gcgccttggc tccaagcagg agctcgtgaa catgataaac 360
accgccacg cctacggcat caaggtcatc gcagacatag taatcaacca ccgcgccgga 420
ggagaccttg agtgaaccc cttcgtcaat gactacacct ggacggactt ctcgaaggtc 480
gcttccggca agtacacggc caactacctc gacttccacc ccaacgaggt caagtgtctg 540
gacgagggca cctttggagg gttcccggac atagcccacg agaagagctg ggaccagtac 600
tggctctggg cgagcaacga gagctacgcc gcctacctca ggagcatcgg cgttgacgca 660
tggcgcttcg actacgtcaa gggctacgga gcgtgggtcg tcaaggactg gctggactgg 720
tggggaggct gggcgctcgg ggagtactgg gacacaaacg ttgatgcact gctcaactgg 780
gccctactga gcgatgcaaa agtcttcgac ttcccgtctt actacaagat ggacgcggcc 840
tttgacaaca agaacattcc cgcactcgtc gaggccctca agaacggggg cacagtcgtc 900
agccgcgacc cgtttaaggc cgtaaccttc gttgcaaacc acgacacgga cataatttgg 960
aacaagtacc cggcctacgc cttcatcctc acctacgagg gccagccgac gatattctac 1020
cgcgactacg aggagtggct caacaaggac aggtcaaga acctcatctg gatacacgac 1080
caccctcgcg gtggaagcac cgacatagtc tactacgata acgatgaact catcttcgtc 1140
aggaacggct acggggacaa gccggggcctt ataacctaca tcaacctagg ctcgagcaag 1200
gccgggaggt ggggtctacgt tccgaagttc gcgggagcgt gcattccacga gtacaccggc 1260
aaccctcgcg gctgggtgga caagtgggtg gactcaagcg ggtgggtgta cctcgaggcc 1320
cctgcccacg acccggccaa cggctattac ggctactccg tctggagcta ctgcgggggtg 1380
ggctga

```

<210> 80
 <211> 461
 <212> PRT
 <213> Bacterial

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<400> 80
Met Lys Pro Ala Lys Leu Leu Val Phe Val Leu Val Val Ser Ile Leu
1      5      10      15
Ala Gly Leu Tyr Ala Gln Pro Ala Gly Ala Ala Lys Tyr Leu Glu Leu
20
Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ser
35
Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp Tyr
50
Asp Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys Gly Met
65
Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu
85
Gly Glu Tyr Asp Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys
100
Gln Glu Leu Val Asn Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys
115
Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu
130
Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val
145
Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu
165
Val Lys Cys Cys Asp Glu Gly Thr Phe Gly Gly Phe Pro Asp Ile Ala
180
His Glu Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser Asn Glu Ser
195
Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp
210
Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Lys Asp Trp Leu Asp Trp
225
Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
245
Leu Leu Asn Trp Ala Tyr Ser Ser Asp Ala Lys Val Phe Asp Phe Pro
260
Leu Tyr Tyr Lys Met Asp Ala Ala Phe Asp Asn Lys Asn Ile Pro Ala
275
Leu Val Glu Ala Leu Lys Asn Gly Gly Thr Val Val Ser Arg Asp Pro
290
phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp
305
Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro
310
315
320

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Thr	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu	Trp	Leu	Asn	Lys	Asp	Arg	Leu
Lys	Asn	Leu	Ile	Trp	Ile	His	Asp	His	Leu	Ala	Gly	Gly	Ser	Thr	Asp
Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu	Ile	Phe	Val	Arg	Asn	Gly	Tyr
Gly	Asp	Lys	Pro	Gly	Leu	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	Ser	Ser	Lys
Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys	Phe	Ala	Gly	Ala	Cys	Ile	His
Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp	Val	Asp	Lys	Trp	Val	Asp	Ser
Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro	Ala	His	Asp	Pro	Ala	Asn	Gly
Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr	Cys	Gly	Val	Gly			

<210>	81
<211>	1386
<212>	DNA
<213>	Bacterial

<400>	81						
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gcacagccag	ctagcgccgc	aaagtattcc	gagctcgaag	aaggcggcgt	tataatgcag		120
gccttctact	gggacgtccc	aggtggagga	atctggtggg	acaccatcag	gagcaagata		180
ccggagtggg	acgagggcggg	aataatccgc	atttgatttc	cgccagccag	caaggggatg		240
agcggcgggt	actcgatggg	ctacgatccc	tacgattttct	ttgacctcgg	cgagatacaac		300
cagaagggaa	ccatcgaaac	gcgctttggc	tctaaacagg	agctcatcaa	tatgataaac		360
acggcccatg	cctacggcat	aaaggtcata	gcggacatcg	tcataaacca	ccgcgcaggc		420
ggagacctcg	agtggaaccc	gttcgttggg	gactacacct	ggacggactt	ctcaaagggtg		480
gcctcggggca	aataatactgc	caactacctc	gacttccacc	ccaacgaggt	caagtgtctgt		540
gacgagggga	catttgaggg	cttccccagac	atagcccacg	agaagagctg	ggaccagcac		600
tggctcttgg	cgagcgatga	gagctacgcc	gcctacctaa	ggagcatcgg	cgttgatgcc		660
tggcgctttg	actactgtga	gggctacgga	gcgtgggtcg	tcaaggactg	gctcaactgg		720
tggggcggct	gggccgttgg	cgagtactgg	gacaccaacg	ttgatgcact	cctcaactgg		780
gcctactcga	gcggcgccaa	ggtcttctgac	ttcccgctct	actacaagat	ggatgaggcc		840
tttgacaaca	aaaacattcc	agcgctcgtc	tctgcccttc	agaacggcca	gactgtttgtc		900
tcccgcgacc	cgttcaaggc	cgtaaccttt	gtagcaaac	acgacaccga	tataattctgg		960
aacaagtacc	ttgcttatgc	tttcatcctc	acctacaag	gccagcccgt	catattctac		1020
cgcgactacg	aggagtggct	caacaaggac	aggttgaaca	acctcatatg	gatacacgac		1080
cacctcgcag	gtggaagcac	gagcatagtc	tactacgaca	gcgacgagat	gatcttctgtg		1140
aggaacggct	atggaagcaa	gcctggcctt	ataacttaca	tcaacctcgg	ctcgagcaag		1200
gttggaaggat	gggtttatgt	gccgaagttc	gcgggcgcgt	gcattccacga	gtatactgggt		1260
aacctcggag	gctgggtaga	caagtacgtc	tactcaagcg	gctgggtcta	tctcgaagct		1320
ccagcttacg	accctgccaa	cgggcagtat	ggctactccg	tgtggagcta	ttgcggtggt		1380
qqgtqa							1386

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<210> 82
<211> 461
<212> PRT
<213> Bacterial
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<400>	82															
Met	Lys	Lys	Phe	Val	Ala	Leu	Phe	Ile	Thr	Met	Phe	Phe	Val	Val	Ser	
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Met	Ala	Val	Val	Ala	Gln	Pro	Ala	Ser	Ala	Ala	Lys	Tyr	Ser	Glu	Leu	
			20					25					30			
Glu	Glu	Gly	Gly	Val	Ile	Met	Gln	Ala	Phe	Tyr	Trp	Asp	Val	Pro	Gly	
		35					40					45				
Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg	Ser	Lys	Ile	Pro	Glu	Trp	Tyr	
	50				55						60					
Glu	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile	Pro	Pro	Ala	Ser	Lys	Gly	Met	
65					70					75					80	
Ser	Gly	Gly	Tyr	Ser	Met	Gly	Tyr	Asp	Pro	Tyr	Asp	Phe	Phe	Asp	Leu	
				85					90					95		
Gly	Glu	Tyr	Asn	Gln	Lys	Gly	Thr	Ile	Glu	Thr	Arg	Phe	Gly	Ser	Lys	

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Gln	Glu	Leu	100	Asn	Met	Ile	Asn	105	Thr	Ala	His	Ala	Tyr	110	Gly	Ile	Lys
Val	Ile	Ala	115	Asp	Ile	Val	Ile	120	Asn	His	Arg	Ala	Gly	125	Gly	Asp	Leu
Trp	Asn	Pro	130	Phe	Val	Gly	Asp	135	Tyr	Thr	Trp	Thr	Asp	140	Phe	Ser	Lys
Ala	Ser	Gly	145	Lys	Tyr	Thr	Ala	150	Asn	Tyr	Leu	Asp	Phe	155	His	Pro	Asn
Val	Lys	Cys	165	Cys	Asp	Glu	Gly	170	Thr	Phe	Gly	Gly	Phe	175	Pro	Asp	Ile
His	Glu	Lys	180	Ser	Trp	Asp	Gln	185	His	Trp	Leu	Trp	Ala	190	Ser	Asp	Glu
Tyr	Ala	Ala	195	Tyr	Leu	Arg	Ser	200	Ile	Gly	Val	Asp	Ala	205	Trp	Arg	Phe
Tyr	Val	Lys	210	Gly	Tyr	Gly	Ala	215	Trp	Val	Val	Lys	Asp	220	Trp	Leu	Asn
Trp	Gly	Gly	225	Trp	Ala	Val	Gly	230	Glu	Tyr	Trp	Asp	Thr	235	Asn	Val	Asp
Leu	Leu	Asn	245	Trp	Ala	Tyr	Ser	250	Gly	Ala	Lys	Val	Phe	255	Asp	Phe	Pro
Leu	Tyr	Tyr	260	Lys	Met	Asp	Glu	265	Ala	Phe	Asp	Asn	Lys	270	Asn	Ile	Pro
Leu	Val	Ser	275	Ala	Leu	Gln	Asn	280	Gly	Gln	Thr	Val	Val	285	Ser	Arg	Asp
Phe	Lys	Ala	290	Val	Thr	Phe	Val	295	Ala	Asn	His	Asp	Thr	300	Asp	Ile	Ile
Asn	Lys	Tyr	305	Leu	Ala	Tyr	Ala	310	Phe	Ile	Leu	Thr	Tyr	315	Glu	Gly	Gln
Val	Ile	Phe	325	Tyr	Arg	Asp	Tyr	330	Glu	Glu	Trp	Leu	Asn	335	Lys	Asp	Arg
Asn	Asn	Leu	340	Ile	Trp	Ile	His	345	Asp	His	Leu	Ala	Gly	350	Gly	Ser	Thr
Ile	Val	Tyr	355	Tyr	Asp	Ser	Asp	360	Glu	Met	Ile	Phe	Val	365	Arg	Asn	Gly
Gly	Ser	Lys	370	Pro	Gly	Leu	Ile	375	Thr	Tyr	Ile	Asn	Leu	380	Gly	Ser	Ser
Val	Gly	Arg	385	Trp	Val	Tyr	Val	390	Pro	Lys	Phe	Ala	Gly	395	Ala	Cys	Ile
Glu	Tyr	Thr	405	Gly	Asn	Leu	Gly	410	Trp	Val	Asp	Lys	Tyr	415	Val	Tyr	Ser
Ser	Gly	Trp	420	Val	Tyr	Leu	Glu	425	Pro	Ala	Tyr	Asp	Pro	430	Ala	Asn	Gly
Gln	Tyr	Gly	435	Tyr	Ser	Val	Trp	440	Ser	Tyr	Cys	Gly	Val	445	Gly		
			450					455						460			

<210> 83
 <211> 1299
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

<400> 83

atggcctctgg	aagacggcgg	gctcataatg	caggccttct	actgggatgt	tcctggagga	60
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gcgatattgga	ttccaccagc	gagtaagggc	atgagcgggtg	gttattccat	gggctacgat	180
ccctacgatt	tctttgacct	cggcgagtac	tatcagaagg	ggacagttga	gacgcgcttc	240
ggctcaaagg	aagaactggt	gaacatgata	aacaccgcac	actcctacgg	cataaagggtg	300
atagcagaca	tagtcataaa	ccaccgcgcc	ggtggagacc	ttgagtggaa	ccccttcgtg	360
aacgactata	cctggacaga	cttctcaaaa	gtcgcctccg	gtaaatatac	ggccaactac	420
cttgacttcc	acccaaacga	gcttcactgt	tgtgatgaag	gtaccttttg	aggataacct	480
gatatatgtc	acgacaaaag	ctgggaccag	tactggctct	gggagagcag	cgaaagctac	540
gctgcctacc	tcaggagcat	aggggttgac	gcctggcggt	tcgactacgt	caagggttac	600
ggagcatggg	ttgttaacga	ctggctcagc	tgggtggggag	gctggggcgt	tggagagtac	660
tgggacacga	acgttgatgc	actcctcaac	tgggcataca	gcagcggcgc	caagggtctt	720
gacttccgc	tctactacaa	gatggacgaa	gccttcgaca	acaccaacat	cccggcatta	780

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gtggatgcac	tcagatacgg	ccagacagtg	gtcagccgcg	atcccttcaa	ggcggtaact	840
ttcgttgcca	accacgatac	agatataatc	tggacaagt	atccggctta	tgcatcctac	900
cttacctatg	agggacagcc	tgttatattc	taccgcgact	acgaggagt	gctcaacaag	960
gataagctta	acaacctcat	ctggatacac	gatcaccttg	ctggagggag	tactgacatt	1020
gtttactacg	acagcgacga	gcttatcttt	gtgagaaacg	gctatggcac	caaaccagga	1080
ctgataacct	atatcaacct	cggctcaagc	aaagttggaa	ggtgggtcta	cgttccaaag	1140
ttcgccgggt	catgcatcca	cgagtacacc	ggcaacctcg	gcggttgat	agacaagtac	1200
gtctcctcca	gcggctgggt	ctatcttgag	gccccagccc	acgacccggc	gaacggctac	1260
tacggctact	ccgtatggag	ctactgcggg	gttgggtga			1299

<210> 84
 <211> 432
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 84
 Met Ala Leu Glu Asp Gly Gly Leu Ile Met Gln Ala Phe Tyr Trp Asp
 1 5 10 15
 Val Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Ala Gln Lys Ile Pro
 20 25 30
 Glu Trp Ala Ser Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser
 35 40 45
 Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe
 50 55 60
 Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe
 65 70 75 80
 Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala His Ser Tyr
 85 90 95
 Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
 100 105 110
 Asp Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe
 115 120 125
 Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His
 130 135 140
 Pro Asn Glu Leu His Cys Cys Asp Glu Gly Thr Phe Gly Gly Tyr Pro
 145 150 155 160
 Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser
 165 170 175
 Ser Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp
 180 185 190
 Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Asn Asp Trp
 195 200 205
 Leu Ser Trp Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn
 210 215 220
 Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe
 225 230 235 240
 Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Thr Asn
 245 250 255
 Ile Pro Ala Leu Val Asp Ala Leu Arg Tyr Gly Gln Thr Val Val Ser
 260 265 270
 Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp
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 Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu
 290 295 300
 Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys
 305 310 315 320
 Asp Lys Leu Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly
 325 330 335
 Ser Thr Asp Ile Val Tyr Tyr Asp Ser Asp Glu Leu Ile Phe Val Arg
 340 345 350
 Asn Gly Tyr Gly Thr Lys Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly
 355 360 365
 Ser Ser Lys Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ser
 370 375 380
 Cys Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Ile Asp Lys Tyr

30 May 2017

2013201807

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385 Val Ser Ser Ser Gly 390 Trp Val Tyr Leu Glu 395 Ala Pro Ala His Asp 400 Pro
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<211> 1299
<212> DNA
<213> Unknown

<220>
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<211> 432
<212> PRT
<213> Unknown

<220>
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35 40 45
Lys Gly Met Ser Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe
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Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe
65 70 75 80
Gly Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala His Ser Tyr
85 90 95
Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly
100 105 110
Gly Leu Glu Trp Asn Pro Phe Val Asn Asp Tyr Thr Trp Thr Asp Phe
115 120 125
Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His
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Pro Asn Glu Leu His Cys Cys Asp Glu Gly Thr Phe Gly Gly Tyr Pro
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Asp Ile Cys His Asp Lys Ser Trp Asp Gln Tyr Trp Leu Trp Ala Ser

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Ser	Glu	Ser	Tyr	165	Ala	Ala	Tyr	Leu	Arg	170	Ser	Ile	Gly	Val	Asp	175	Ala	Trp
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Leu	Ser	Trp	Trp	195	Gly	Gly	Trp	200	Ala	Val	Gly	Glu	Tyr	Trp	Asp	205	Thr	Asn
Val	Asp	Ala	Leu	210	Leu	Asn	Trp	215	Ala	Tyr	Asn	Ser	Gly	Ala	Lys	Val	Phe	
Asp	Phe	Pro	Leu	225	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	Asp	Asn	Thr	Asn		
Ile	Pro	Ala	Leu	245	Val	Tyr	Ala	Leu	Lys	250	Asn	Gly	Gly	Thr	Val	Val	Ser	
Arg	Asp	Pro	Phe	260	Lys	Ala	Val	Thr	Phe	265	Val	Ala	Asn	His	Asp	Thr	Asp	
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Gly	Gln	Pro	Val	290	Ile	Phe	Tyr	Arg	Asp	295	Tyr	Glu	Glu	Trp	Leu	Asn	Lys	
Asp	Lys	Leu	Asn	305	Asn	Leu	Ile	Trp	Ile	310	His	Asp	His	Leu	Ala	Gly	Gly	
Ser	Thr	Asp	Ile	325	Val	Tyr	Tyr	Asp	Ser	330	Asp	Glu	Leu	Ile	Phe	Val	Arg	
Asn	Gly	Tyr	Gly	340	Thr	Lys	Pro	Gly	Leu	345	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	
Ser	Ser	Lys	Ala	355	Gly	Arg	Trp	Val	Tyr	360	Val	Pro	Lys	Phe	Ala	Gly	Ser	
Cys	Ile	His	Glu	370	Tyr	Thr	Gly	Ser	Leu	375	Gly	Gly	Trp	Ile	Asp	Lys	Tyr	
Val	Ser	Ser	Ser	385	Gly	Trp	Val	Tyr	Leu	390	Glu	Ala	Pro	Ala	His	Asp	Pro	
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 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 87

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accgctcttt	ggctgccgcc	cgcttacaaa	ggaacaagcc	gcagcgacgt	aggggtacga	240
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 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 88

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 35      40      45
Ala Asn Glu Ala Asn Asn Leu Ser Ser Leu Gly Ile Thr Ala Leu Trp
 50      55      60
Leu Pro Pro Ala Tyr Lys Gly Thr Ser Arg Ser Asp Val Gly Tyr Gly
 65      70      75      80
Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val
 85      90      95
Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Leu Gln Ala Ile Gln Ala
100      105      110
Ala His Ala Ala Gly Met Gln Val Tyr Ala Asp Val Val Phe Asp His
115      120      125
Lys Gly Gly Ala Asp Gly Thr Glu Trp Val Asp Ala Val Glu Val Asn
130      135      140
Pro Ser Asp Arg Asn Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala
145      150      155      160
Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn Thr Tyr Ser Ser Phe
165      170      175
Lys Trp Arg Trp Tyr His Phe Asp Gly Val Asp Trp Asp Glu Ser Arg
180      185      190
Lys Leu Ser Arg Ile Tyr Lys Phe Arg Gly Ile Gly Lys Ala Trp Asp
195      200      205
Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala
210      215      220
Asp Leu Asp Met Asp His Pro Glu Val Val Thr Glu Leu Lys Asn Trp
225      230      235      240
Gly Lys Trp Tyr Val Asn Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp
245      250      255
Ala Val Lys His Ile Lys Phe Ser Phe Phe Pro Asp Trp Leu Ser Tyr
260      265      270
Val Arg Ser Gln Thr Gly Lys Pro Leu Phe Thr Val Gly Glu Tyr Trp
275      280      285
Ser Tyr Asp Ile Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asp Gly
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Ser Lys Ser Gly Gly Ala Phe Asp Met Arg Thr Leu Met Thr Asn Thr
325      330      335
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340      345      350
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355      360      365
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370      375      380
Cys Val Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro Gln Tyr Asn Ile Pro
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465

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 <212> DNA
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<213> Bacterial

<400> 90

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Asp	Gln	755	Ile	Ala	Pro	Ala	760	Ala	Ile	Glu	Leu	Arg	Gln	Pro	Ala
Glu	Ser	770	Gly	Gln	Val	Asn	775	Leu	Ser	Trp	Thr	Phe	Val	Gly	Lys
Gly	Asp	785	Ala	Tyr	Leu	Ala	790	Ile	Glu	Arg	Asn	Gly	Asp	Ile	Val
Thr	Thr	805	Thr	Ser	Ile	Gly	810	Thr	Asp	Tyr	Asp	Val	Val	Glu	Asn
Gly	Thr	820	Glu	Tyr	Thr	Tyr	825	Val	Val	Lys	Leu	Tyr	Asp	Arg	Ala
Val	Val	835	Ala	Ser	Asn	Thr	840	Lys	Val	Thr	Pro	Asp	Ile	Val	Met
Lys	Val	850	Ile	Phe	Lys	Val	855	Arg	Ala	Pro	Asp	Tyr	Thr	Pro	Leu
Arg	Ile	865	Thr	Ile	Pro	Asn	870	Ser	Leu	Asn	Gly	Trp	Asn	Thr	Gly
Glu	Met	885	Ser	Arg	Asn	Gly	890	Ala	Val	Thr	Pro	Asp	Trp	Gln	Phe
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Ser	Trp	915	Asp	Gln	Glu	Gly	920	Ala	Asp	His	Thr	Arg	Glu	Asp	Asp
Asp	Asp	930	Val	Ser	Tyr	Thr	935	Gly	Tyr	Gly	Thr	Ile	Gly	Thr	Asp
Lys	Val	945	Thr	Val	His	Asn	950	Glu	Gly	Asn	Asn	Thr	Met	Ile	Val
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Lys	Gln	980	Gly	Ser	Gln	Val	985	Ile	Lys	Gly	Asn	Ala	Ile	Lys	Asn
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phe	Ser	1010	Tyr	Thr	Phe	Ala	1015	Pro	Ala	Ser	His	Gln	Lys	Glu	Val
His	Ile	1025	Glu	Pro	Ser	Ala	1030	Glu	Ser	Lys	Thr	Ala	Ile	Phe	Asn
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<211> 1650
<212> DNA
<213> Bacterial
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<211> 549
<212> PRT
<213> Bacterial
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Lys	Ala	Ala	Ala	Pro	Phe	Asn	Gly	Thr	Met	Met	Gln	Tyr	Phe	Glu	Trp	
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Tyr	Asp	Leu	Gly	Glu	Phe	Asn	Gln	Lys	Gly	Thr	Val	Arg	Thr	Lys	Tyr	
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Ile	Tyr	Lys	195	Phe	Arg	Gly	Ile	200	Gly	Lys	Ala	Trp	205	Asp	Trp	Glu	Val	Asp
Thr	210	Glu	Asn	Gly	Asn	Tyr	215	Asp	Tyr	Leu	Met	Tyr	220	Ala	Asp	Leu	Asp	Met
225	Asp	His	Pro	Glu	Val	Val	Thr	Glu	Leu	Lys	235	Asn	Trp	Gly	Lys	Trp	Tyr	240
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Ile	Lys	Phe	260	Ser	Phe	Phe	Pro	265	Asp	Trp	Leu	Ser	Tyr	Val	270	Arg	Ser	Gln
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Gln	Pro	Thr	Leu	Ala	Val	Thr	Phe	Val	Asp	Asn	His	Asp	Thr	Glu	Pro	355	360	375
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Tyr	Ala	Phe	Ile	Leu	Thr	Arg	Gln	Glu	Gly	Tyr	Pro	Cys	Val	Phe	Tyr	385	400	415
Gly	Asp	Tyr	Tyr	Gly	Ile	Pro	Gln	Tyr	Asn	Ile	Pro	Ser	Leu	Lys	Ser	405	415	430
Lys	Ile	Asp	Pro	Leu	Leu	Ile	Ala	Arg	Asp	Tyr	Ala	Tyr	Gly	Thr	Arg	Glu	420	435
Gln	His	Asp	Tyr	Leu	Asp	His	Ser	Asp	Ile	Ile	Gly	Trp	Thr	Arg	Glu	435	445	460
Gly	Val	Thr	Glu	Lys	Pro	Gly	Ser	Gly	Leu	Ala	Ala	Leu	Ile	Thr	Asp	445	455	470
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Lys	Val	Phe	Tyr	Asp	Leu	Thr	Gly	Asn	Arg	Ser	Asp	Thr	Val	Thr	Ile	475	485	500
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 <213> Bacterial

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 <213> Bacterial

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 <212> DNA
 <213> Bacterial

<400> 95

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 <212> PRT
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<400> 96

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Leu	Gly	Val	Asp	Thr	Val	Trp	Phe	Leu	Pro	Phe	Asn	Lys	Ser	Lys	Ser
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Tyr	His	Gly	Tyr	Asp	Val	Glu	Asp	Tyr	Tyr	Asp	Val	Glu	Pro	Asp	Tyr
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Gly	Thr	Leu	Gln	Asp	Leu	Asp	Asn	Met	Ile	Lys	Val	Leu	Asn	Glu	Asn

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	Trp	Asn	Tyr	Tyr	Ile	100	Met	Ser	Leu	Asp	Glu	105	Pro	Gln	Asn	Lys	Asn	110	His
	Trp	His	Tyr	Lys	Val	115	Asn	Ser	Lys	Gly	Gln	120	Thr	Val	125	Trp	Tyr	125	Phe
	Leu	Phe	Asp	Ser	Ser	130	Met	Pro	Asp	Leu	Asn	135	Tyr	Asp	Asn	Pro	Lys	140	Val
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Arg Trp Ile Ser Ile Leu Ile Asp Gln Lys Tyr Ser Gly Asn Thr Gln
      290      295      300
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      420      425      430
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 gatgtaccct atagcagtaa tgactttaac tcctgtacag gagatattga ctataataac 540
 cgttggcaaa cacagcattg tgatttagtc ggtcttaatg atctaaaaac aggatctgac 600
 tacgtccgcc aaaaaatagc ggattatatg aacgacgcaa tcagtatggg ttagctggtt 660
 ttccgtattg atgcagccaa acatatacca gcaggtgata tagctgccat taaaggtaaa 720
 ttaaatggta atccatacat cttccaagag gtaattgggt catccggcga acctgttcga 780
 ccgactgaat acacctttat cgggtggtgtc acggaatttc aatttgctcg aaaattgggt 840
 ccagccttcc gcaatagtaa tattgcttgg ttaaaagaca ttggcagtc aatggaatta 900
 tccagtgtct atgccgtaac atttgtaacg aatcatgatg aagagcgtca taaccggaat 960
 ggtcctattt ggcacggcgt tcaaggtaat gggtatgcat tagcaaatat tttcacctta 1020
 gcttaccctt acggctatcc aaaaatcatg tcaggatact tcttcacag tgactttaac 1080
 gcagctccac caagcagttg tatacacaca ggaaatgcgt gtgggttttga tggcggagac 1140
 tgggtatgca aacacaaatg gcgcggtatt gctaacatgg ttgccttccg caactataca 1200
 gcaagcgaat ggcgtatcag taattgggtg caaaacagta acgaccaa atgttttgggt 1260
 cgcggtgggt taggttttgt tgttattaat aaacgtgcta atggtagcat taatcaaagt 1320
 tttgatcagg gaatgcctga tggccaatac tgtaacataa tagaagctaa ctttgatgaa 1380
 agcacgggcc aatgtagtgc agctacagat tccaacgggt aagccgttat taccgtcagt 1440
 ggtgggcaag ctaactttta tgtagcaggc gatcatgctg ctgcaattca tgttggcgca 1500
 aaaattgggt atcaatgtag tgggtgatgat tgcccattga caggatccga ttgtaataat 1560
 gatcctaaac ctgattttgc agtaccagca acatcaattt gtacatcaga aaatttacct 1620
 acgctatatt actggggagc acagcctaca gatagcttag cgaatgcagc ttggccaggt 1680
 gtcgcaatgc aaacaaatgg cgactttaag tgtcatgatt taggtgtcga actaaccaaa 1740
 attaacgcca tctttagtga caatgggtgca aataaaacag ctgatctaac tgttactgggt 1800
 gcaggttggt ataaagacgg gacttgagac accttaca aaatttggtt tgaaattacc 1860
 ggtgcacaaa ccaatccagt cgggtggcgac gaagtcgtgt acttccgagg tactgtctaat 1920
 gactggggta aagcacaatt agattatgac gcaactagcg gtttgtatta cacaatacaa 1980
 agctttaatg gtgaagaagc acctgcgcgt tttaaaattg ataattgtag ttggactgaa 2040
 gcttatccaa cagctgatta ccaagttaca gataacaatt cataccgcat taactttaat 2100
 agcgatagca aagcgattac agtaaacgca caataa 2136

<210> 102
 <211> 711
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 102
 Met Arg Phe Phe Pro Lys Leu Ile Ser Pro Phe Pro Gln Asn Thr Arg
 1 5 10 15
 Glu Trp Gln Arg Ser Ala Val Ser Arg Asp Thr Glu Gln Leu Gln Arg
 20 25 30
 Lys Val Ile Met Ile Asn Leu Lys Asn Thr Ile Ser Ala Leu Val
 35 40 45
 Ala Gly Met Val Leu Gly Phe Ala Ser Asn Ala Met Ala Val Pro Arg
 50 55 60

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Thr 65	Ala	Phe	Val	His	Leu 70	Phe	Glu	Trp	Lys	Trp 75	Glu	Asp	Val	Ala	Gln 80
Glu	Cys	Glu	Thr	Phe 85	Leu	Gly	Pro	Lys	Gly 90	Phe	Ala	Ala	Val	Gln 95	Val
Ser	Pro	Pro	Thr 100	Lys	Ser	His	Asn	Thr 105	Asp	Ala	Trp	Trp	Gly 110	Arg	Tyr
Gln	Pro	Val 115	Ser	Tyr	Ala	Phe	Glu 120	Gly	Arg	Ser	Gly	Asn 125	Arg	Ser	Gln
Phe	Lys 130	Asn	Met	Val	Gln	Arg 135	Cys	Lys	Ala	Val	Gly 140	Val	Asp	Ile	Tyr
Val 145	Asp	Ala	Val	Ile	Asn 150	His	Met	Ala	Ala	Tyr 155	Asp	Arg	Asn	Phe	Pro 160
Asp	Val	Pro	Tyr	Ser 165	Ser	Asn	Asp	Phe	Asn 170	Ser	Cys	Thr	Gly	Asp 175	Ile
Asp	Tyr	Asn	Asn 180	Arg	Trp	Gln	Thr	Gln 185	His	Cys	Asp	Leu	Val 190	Gly	Leu
Asn	Asp	Leu 195	Lys	Thr	Gly	Ser	Asp 200	Tyr	Val	Arg	Gln	Lys 205	Ile	Ala	Asp
Tyr	Met 210	Asn	Asp	Ala	Ile	Ser 215	Met	Gly	Val	Ala	Gly 220	Phe	Arg	Ile	Asp
Ala 225	Ala	Lys	His	Ile	Pro 230	Ala	Gly	Asp	Ile	Ala 235	Ala	Ile	Lys	Gly	Lys 240
Leu	Asn	Gly	Asn	Pro 245	Tyr	Ile	Phe	Gln	Glu 250	Val	Ile	Gly	Ala	Ser 255	Gly
Glu	Pro	Val	Arg 260	Pro	Thr	Glu	Tyr	Thr 265	Phe	Ile	Gly	Gly	Val 270	Thr	Glu
Phe	Gln	Phe 275	Ala	Arg	Lys	Leu	Gly 280	Pro	Ala	Phe	Arg	Asn 285	Ser	Asn	Ile
Ala	Trp 290	Leu	Lys	Asp	Ile	Gly 295	Ser	Gln	Met	Glu	Leu 300	Ser	Ser	Ala	Asp
Ala 305	Val	Thr	Phe	Val	Thr 310	Asn	His	Asp	Glu	Glu 315	Arg	His	Asn	Pro	Asn 320
Gly	Pro	Ile	Trp	His 325	Gly	Val	Gln	Gly	Asn 330	Gly	Tyr	Ala	Leu	Ala 335	Asn
Ile	Phe	Thr	Leu 340	Ala	Tyr	Pro	Tyr	Gly 345	Tyr	Pro	Lys	Ile	Met 350	Ser	Gly
Tyr	Phe	Phe 355	His	Gly	Asp	Phe	Asn 360	Ala	Ala	Pro	Pro	Ser 365	Ser	Gly	Ile
His	Thr 370	Gly	Asn	Ala	Cys	Gly 375	Phe	Asp	Gly	Gly	Asp 380	Trp	Val	Cys	Glu
His 385	Lys	Trp	Arg	Gly	Ile 390	Ala	Asn	Met	Val	Ala 395	Phe	Arg	Asn	Tyr	Thr 400
Ala	Ser	Glu	Trp	Arg 405	Ile	Ser	Asn	Trp	Trp 410	Gln	Asn	Ser	Asn	Asp 415	Gln
Ile	Ala	Phe	Gly	Arg	Gly	Gly	Leu	Gly 425	Phe	Val	Val	Ile	Asn 430	Lys	Arg
Ala	Asn	Gly 435	Ser	Ile	Asn	Gln	Ser 440	Phe	Asp	Thr	Gly	Met 445	Pro	Asp	Gly
Gln	Tyr 450	Cys	Asn	Ile	Ile	Glu 455	Ala	Asn	Phe	Asp	Glu 460	Ser	Thr	Gly	Gln
Cys 465	Ser	Ala	Ala	Thr	Asp 470	Ser	Asn	Gly	Gln	Ala 475	Val	Ile	Thr	Val	Ser 480
Gly	Gly	Gln	Ala	Asn 485	Phe	Asn	Val	Ala	Gly 490	Asp	His	Ala	Ala	Ala 495	Ile
His	Val	Gly	Ala 500	Lys	Ile	Gly	Asp	Gln 505	Cys	Ser	Gly	Asp	Asp 510	Cys	Pro
Cys	Thr	Gly 515	Ser	Asp	Cys	Asn	Asn 520	Asp	Pro	Lys	Pro	Asp 525	Phe	Ala	Val
Pro	Ala 530	Thr	Ser	Ile	Cys	Thr 535	Ser	Glu	Asn	Leu	Pro 540	Thr	Leu	Tyr	Tyr
Trp 545	Gly	Ala	Gln	Pro	Thr 550	Asp	Ser	Leu	Ala	Asn 555	Ala	Ala	Trp	Pro	Gly 560
Val	Ala	Met	Gln	Thr 565	Asn	Gly	Asp	Phe	Lys 570	Cys	His	Asp	Leu	Gly 575	Val
Glu	Leu	Thr	Lys 580	Ile	Asn	Ala	Ile	Phe 585	Ser	Asp	Asn	Gly	Ala 590	Asn	Lys
Thr	Ala	Asp 595	Leu	Thr	Val	Thr	Gly 600	Ala	Gly	Cys	Tyr	Lys 605	Asp	Gly	Thr

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Trp 610	Ser 615	Thr	Leu	Gln	Asn	Cys 615	Gly	Phe	Glu	Ile	Thr 620	Gly	Ala	Gln	Thr
Asn 625	Pro	Val	Gly	Gly	Asp 630	Glu	Val	Trp	Tyr	Phe 635	Arg	Gly	Thr	Ala	Asn 640
Asp	Trp	Gly	Lys	Ala 645	Gln	Leu	Asp	Tyr	Asp 650	Ala	Thr	Ser	Gly	Leu 655	Tyr
Tyr	Thr	Ile	Gln 660	Ser	Phe	Asn	Gly	Glu 665	Glu	Ala	Pro	Ala	Arg 670	Phe	Lys
Ile	Asp	Asn 675	Gly	Ser	Trp	Thr	Glu 680	Ala	Tyr	Pro	Thr	Ala 685	Asp	Tyr	Gln
Val	Thr 690	Asp	Asn	Asn	Ser	Tyr 695	Arg	Ile	Asn	Phe	Asn 700	Ser	Asp	Ser	Lys
Ala 705	Ile	Thr	Val	Asn	Ala 710	Gln									

<210>	103
<211>	1650
<212>	DNA
<213>	Unknown

<220>
<223> Obtained from an environmental sample.

<400>	103						
gtgctaacgt	ttcaccgcat	cattcgaaaa	ggatggatgt	tcctgctcgc	gtttttgctc		60
actgcctcgc	tgttctgccc	aacaggacag	cccgccaaag	ctgccgcacc	gtttaacggc		120
accatgatgc	agtattttga	atggtaactg	ccggatgatg	gcacgttatg	gaccaaagtg		180
gccaatgaag	ccaacaactt	atccagcctt	ggcatcacgc	ctctttggct	gcgcgccgct		240
tacaaaaggc	caagccgcag	cgacgtatgg	tacggagtat	acgacttgta	tgacctcggc		300
gaattcaatc	aaaaagggac	cgtcgcgaca	aaatacggaa	caaaagctca	atatcttcaa		360
gccattcaag	ccgcccacgc	cgctggaatg	caagtgtacg	ccgatgtcgt	gttcgaccat		420
aaaggcggcg	ccgacggcac	ggaatgggtg	gacgccgtcg	aagtcaatcc	gtccgaccgc		480
aaccaagaaa	tcctggggcac	ctatcaaatc	caagcatgga	cgaatttga	ttttccggg		540
cggggcaaca	ccctaccagc	ctttaagtgg	cgctggtacc	attttgacgg	cgttgatctg		600
gacgaaaagc	gaaaattgag	ccgcattttac	aaattccgcg	gcatacggca	agcgtgggat		660
tggaagtag	acacggaaaa	cggaaactat	gactacttaa	tgtatgccga	ccttgatatg		720
gatcatcccg	aagtcgtgac	cgagctgaaa	aactgggggg	aatggtatgt	caacacaacg		780
aacattgatg	ggttcgggct	tgatgccgtc	aagcatatta	agttcagttt	ttttcctgat		840
tggttgctgt	atgtgcgttc	tcagatggc	aagccgctat	ttaccgtcgg	ggaatatctg		900
agctatgaca	tcacaagatt	gcacaattac	attacgaaaa	caaacggaac	gatgtctttg		960
tttgatgccc	cgttacacaa	caaattttat	accgcttcca	aatcaggggg	cgcatttgat		1020
atgcgcacgt	taatgaccaa	tactctcatg	aaagatcaac	cgacattggc	cgtcaccttc		1080
gttgataatc	atgacaccga	acccggccaa	gcgctgcagt	catgggtcga	cccattggtc		1140
aaaccgttgg	cttacgcctt	tattctaaat	cggcaggaag	gatacccggt	cgtcttttat		1200
ggtgactatt	atggcatctc	acaatataac	attctcttcg	tgaaaaagca	aatcgatccg		1260
ctctcatatg	cgcgaggga	ttatgcttac	ggaacgcaac	atgattatct	tgatcactcc		1320
gacatcatcg	ggtggacaag	ggaaggggtc	actgaaaaac	caggatccgg	gctggccgca		1380
ctgatcaccg	atgggcccgg	aggaagcaaa	tggatgtacg	ttggcaaaac	acacgctgga		1440
aaagtgttct	atgaccttac	cggcaaccgg	agtgaaccgc	tcaccatcaa	cagtgatgga		1500
tggggggaa	tcaaagtcaa	tggcggttcg	gtttcggttt	gggttcctag	aaaaacgacc		1560
gtttctacca	tcgctcggcc	gatcacaacc	cgaccgtgga	ctggtgaatt	cgtccgttgg		1620
accgaaccac	ggttgggtgc	atggccttga					1650

<210>	104
<211>	549
<212>	PRT
<213>	Unknown

<220>
<223> obtained from an environmental sample.

<400> 104
Val Leu Thr Phe His Arg Ile Ile Arg Lys Gly Trp Met Phe Leu Leu
1 5 10 15
Ala Phe Leu Leu Thr Ala Ser Leu Phe Cys Pro Thr Gly Gln Pro Ala
20 25 30
Lys Ala Ala Pro Phe Asn Gly Thr Met Met Gln Tyr Phe Glu Trp
35 40 45

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Tyr	Leu	Pro	Asp	Asp	Gly	Thr	Leu	Trp	Thr	Lys	Val	Ala	Asn	Glu	Ala
50	50					55				60					
Asn	Asn	Leu	Ser	Ser	Leu	Gly	Ile	Thr	Ala	Leu	Trp	Leu	Pro	Pro	Ala
65					70					75					80
Tyr	Lys	Gly	Thr	Ser	Arg	Ser	Asp	Val	Gly	Tyr	Gly	Val	Tyr	Asp	Leu
				85					90					95	
Tyr	Asp	Leu	Gly	Glu	Phe	Asn	Gln	Lys	Gly	Thr	Val	Arg	Thr	Lys	Tyr
			100					105					110		
Gly	Thr	Lys	Ala	Gln	Tyr	Leu	Gln	Ala	Ile	Gln	Ala	Ala	His	Ala	Ala
		115					120				125				
Gly	Met	Gln	Val	Tyr	Ala	Asp	Val	Val	Phe	Asp	His	Lys	Gly	Gly	Ala
	130					135					140				
Asp	Gly	Thr	Glu	Trp	Val	Asp	Ala	Val	Glu	Val	Asn	Pro	Ser	Asp	Arg
145					150					155					160
Asn	Gln	Glu	Ile	Ser	Gly	Thr	Tyr	Gln	Ile	Gln	Ala	Trp	Thr	Lys	Phe
				165					170					175	
Asp	Phe	Pro	Gly	Arg	Gly	Asn	Thr	Tyr	Ser	Ser	Phe	Lys	Trp	Arg	Trp
			180					185					190		
Tyr	His	Phe	Asp	Gly	Val	Asp	Trp	Asp	Glu	Ser	Arg	Lys	Leu	Ser	Arg
		195					200					205			
Ile	Tyr	Lys	Phe	Arg	Gly	Ile	Gly	Lys	Ala	Trp	Asp	Trp	Glu	Val	Asp
	210					215					220				
Thr	Glu	Asn	Gly	Asn	Tyr	Asp	Tyr	Leu	Met	Tyr	Ala	Asp	Leu	Asp	Met
225					230					235					240
Asp	His	Pro	Glu	Val	Val	Thr	Glu	Leu	Lys	Asn	Trp	Gly	Glu	Trp	Tyr
				245					250					255	
Val	Asn	Thr	Thr	Asn	Ile	Asp	Gly	Phe	Arg	Leu	Asp	Ala	Val	Lys	His
			260					265					270		
Ile	Lys	Phe	Ser	Phe	Phe	Pro	Asp	Trp	Leu	Ser	Tyr	Val	Arg	Ser	Gln
		275					280					285			
Thr	Gly	Lys	Pro	Leu	Phe	Thr	Val	Gly	Glu	Tyr	Trp	Ser	Tyr	Asp	Ile
	290					295					300				
Asn	Lys	Leu	His	Asn	Tyr	Ile	Thr	Lys	Thr	Asn	Gly	Thr	Met	Ser	Leu
305					310					315					320
Phe	Asp	Ala	Pro	Leu	His	Asn	Lys	Phe	Tyr	Thr	Ala	Ser	Lys	Ser	Gly
				325					330					335	
Gly	Ala	Phe	Asp	Met	Arg	Thr	Leu	Met	Thr	Asn	Thr	Leu	Met	Lys	Asp
			340					345					350		
Gln	Pro	Thr	Leu	Ala	Val	Thr	Phe	Val	Asp	Asn	His	Asp	Thr	Glu	Pro
		355					360					365			
Gly	Gln	Ala	Leu	Gln	Ser	Trp	Val	Asp	Pro	Trp	Phe	Lys	Pro	Leu	Ala
	370					375					380				
Tyr	Ala	Phe	Ile	Leu	Thr	Arg	Gln	Glu	Gly	Tyr	Pro	Cys	Val	Phe	Tyr
385					390					395					400
Gly	Asp	Tyr	Tyr	Gly	Ile	Pro	Gln	Tyr	Asn	Ile	Pro	Ser	Leu	Lys	Ser
				405					410					415	
Lys	Ile	Asp	Pro	Leu	Leu	Ile	Ala	Arg	Arg	Asp	Tyr	Ala	Tyr	Gly	Thr
			420					425					430		
Gln	His	Asp	Tyr	Leu	Asp	His	Ser	Asp	Ile	Ile	Gly	Trp	Thr	Arg	Glu
		435					440					445			
Gly	Val	Thr	Glu	Lys	Pro	Gly	Ser	Gly	Leu	Ala	Ala	Leu	Ile	Thr	Asp
	450					455					460				
Gly	Pro	Gly	Gly	Ser	Lys	Trp	Met	Tyr	Val	Gly	Lys	Gln	His	Ala	Gly
465					470					475					480
Lys	Val	Phe	Tyr	Asp	Leu	Thr	Gly	Asn	Arg	Ser	Asp	Thr	Val	Thr	Ile
				485					490					495	
Asn	Ser	Asp	Gly	Trp	Gly	Glu	Phe	Lys	Val	Asn	Gly	Gly	Ser	Val	Ser
			500					505					510		
Val	Trp	Val	Pro	Arg	Lys	Thr	Thr	Val	Ser	Thr	Ile	Ala	Arg	Pro	Ile
	515						520					525			
Thr	Thr	Arg	Pro	Trp	Thr	Gly	Glu	Phe	Val	Arg	Trp	Thr	Glu	Pro	Arg
	530					535					540				
Leu	Val	Ala	Trp	Pro											
545															

<210> 105
 <211> 1650
 <212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 105

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attgtcctt	tttccattca	aacagaaaaa	gtccgcgctg	gaagtgttcc	agtgaatgga	120
acgatgatgc	aatatcttcga	atgggtacctt	ccagacgatg	gaacactatg	gacgaaagta	180
gcaaaataacg	cccaatcttt	agcgaatctt	ggcattactg	ccctttggct	ttccccctgcc	240
tataaaaggaa	caagcagcag	tgacgtttgga	tatggcgttt	atgattttata	tgacctagga	300
gagttttaatc	aaaaaggaac	tgtccgaaca	aaatacggaa	caaaaacaca	atatatccaa	360
gcaatccaag	cggcgcatat	agcaggaatg	caagtatatg	cagatgtcgt	ctttaacat	420
aaagccgggtg	cagatgggac	agaactagt	gatgcagtag	aagtaaacc	ttctgaccgc	480
aatcaagaaa	tatcaggaac	atatcaaatc	caagcggtga	caaaatttga	ttttcctggt	540
cgtggaaaca	cctattctag	ttttaaatgg	cgttggtatc	atttcgatgg	aacggactgg	600
gatgagagta	gaaaactaaa	tcgtattttac	aaattccgcg	gcacgggaaa	agcatgggat	660
tgggaagtag	atacagaaaa	tgggaattat	gactatctca	tgtatgcaga	tttggaatatg	720
gatcatccag	aggttgatc	tgaactaaaa	aatgggggaa	agtgggtatgt	aaccacaacc	780
aatatcgacg	gattccgtct	ggatgcagtg	aagcatatta	aatatagctt	tttcccagac	840
tggctatcgt	atgtacgaac	ccaaacacaa	aagcctcttt	ttgccgttgg	cgaatttttg	900
agctatgaca	ttaacaagct	acacaactat	attacaaaga	cgaacggctc	tatgtcccta	960
ttcgatgccc	cgctgcataa	caattttttat	atagcatcga	aatcagggtg	ctattttgat	1020
atgcgcacat	tactcaacaa	cacattgatg	aaagatcaac	caacactatc	ggtcacatta	1080
gtagacaatc	acgatactga	gccagggcaa	tctttgcagt	cgtgggtcga	gccgtgggtt	1140
aaaccggttag	cttacgcatt	tatcttgacc	cgccaagaag	gttatccgtg	catcttttat	1200
ggagattact	atgggtatttc	aaaatacaac	attcctgcgc	tgaaaagcaa	acttgatccg	1260
ctgttaattg	ctcgaagaga	ttatgcctac	ggaacacagc	acgactatat	tgacaatgca	1320
gatattatcg	gctggacgcg	ggaaggagta	gctgaaaaag	caaattcggg	acttgctgca	1380
ctcattaccg	acggacctgg	cgggaagcaa	tggatgtatg	ttggcaaaca	acacgctggc	1440
aaaacgtttt	atgatctaac	cggcaatcga	agtgtatacag	tgacaatcaa	cgctgatgga	1500
tggggagaat	ttaaagtcaa	tggagggtct	gtatccatat	gggttccaaa	aacatcaacc	1560
acttcccaaa	tcacattttac	tgtaaataat	gccacaaccg	tttgggggaca	aaatgtatac	1620
gttgtcggga	atatttcgca	gctggggcaac				1650

<210> 106

<211> 550

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 106

Met	Ser	Leu	Phe	Lys	Lys	Ile	Phe	Pro	Trp	Ile	Val	Ser	Leu	Leu	Leu
1				5					10					15	
Leu	Phe	Ser	Phe	Ile	Ala	Pro	Phe	Ser	Ile	Gln	Thr	Glu	Lys	Val	Arg
			20					25					30		
Ala	Gly	Ser	Val	Pro	Val	Asn	Gly	Thr	Met	Met	Gln	Tyr	Phe	Glu	Trp
		35					40					45			
Tyr	Leu	Pro	Asp	Asp	Gly	Thr	Leu	Trp	Thr	Lys	Val	Ala	Asn	Asn	Ala
	50				55					60					
Gln	Ser	Leu	Ala	Asn	Leu	Gly	Ile	Thr	Ala	Leu	Trp	Leu	Pro	Pro	Ala
65				70					75					80	
Tyr	Lys	Gly	Thr	Ser	Ser	Asp	Val	Gly	Tyr	Gly	Val	Tyr	Asp	Leu	
			85					90					95		
Tyr	Asp	Leu	Gly	Glu	Phe	Asn	Gln	Lys	Gly	Thr	Val	Arg	Thr	Lys	Tyr
		100						105					110		
Gly	Thr	Lys	Thr	Gln	Tyr	Ile	Gln	Ala	Ile	Gln	Ala	Ala	His	Thr	Ala
		115					120					125			
Gly	Met	Gln	Val	Tyr	Ala	Asp	Val	Val	Phe	Asn	His	Lys	Ala	Gly	Ala
	130					135					140				
Asp	Gly	Thr	Glu	Leu	Val	Asp	Ala	Val	Glu	Val	Asn	Pro	Ser	Asp	Arg
145				150					155					160	
Asn	Gln	Glu	Ile	Ser	Gly	Thr	Tyr	Gln	Ile	Gln	Ala	Trp	Thr	Lys	Phe
			165					170						175	
Asp	Phe	Pro	Gly	Arg	Gly	Asn	Thr	Tyr	Ser	Ser	Phe	Lys	Trp	Arg	Trp
			180					185					190		

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Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg
 195 200 205
 Ile Tyr Lys Phe Arg Gly Thr Lys Ala Trp Asp Trp Glu Val Asp
 210 215 220
 Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met
 225 230 235 240
 Asp His Pro Glu Val Val Ser Glu Leu Lys Asn Trp Gly Lys Trp Tyr
 245 250 255
 Val Thr Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His
 260 265 270
 Ile Lys Tyr Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Thr Gln
 275 280 285
 Thr Gln Lys Pro Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Ile
 290 295 300
 Asn Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu
 305 310 315 320
 Phe Asp Ala Pro Leu His Asn Asn Phe Tyr Ile Ala Ser Lys Ser Gly
 325 330 335
 Gly Tyr Phe Asp Met Arg Thr Leu Leu Asn Asn Thr Leu Met Lys Asp
 340 345 350
 Gln Pro Thr Leu Ser Val Thr Leu Val Asp Asn His Asp Thr Glu Pro
 355 360 365
 Gly Gln Ser Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Pro Leu Ala
 370 375 380
 Tyr Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Ile Phe Tyr
 385 390 395 400
 Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Ala Leu Lys Ser
 405 410 415
 Lys Leu Asp Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr
 420 425 430
 Gln His Asp Tyr Ile Asp Asn Ala Asp Ile Ile Gly Trp Thr Arg Glu
 435 440 445
 Gly Val Ala Glu Lys Ala Asn Ser Gly Leu Ala Ala Leu Ile Thr Asp
 450 455 460
 Gly Pro Gly Gly Ser Lys Trp Met Tyr Val Gly Lys Gln His Ala Gly
 465 470 475 480
 Lys Thr Phe Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile
 485 490 495
 Asn Ala Asp Gly Trp Gly Glu Phe Lys Val Asn Gly Gly Ser Val Ser
 500 505 510
 Ile Trp Val Pro Lys Thr Ser Thr Thr Ser Gln Ile Thr Phe Thr Val
 515 520 525
 Asn Asn Ala Thr Thr Val Trp Gly Gln Asn Val Tyr Val Val Gly Asn
 530 535 540
 Ile Ser Gln Leu Gly Asn
 545 550

<210> 107

<211> 1509

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 107

atggacagcc	tcgacgcgcc	ggagcagaag	ccctgggtga	aggatggcag	gctctccgcg	60
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ccggccgagg	aagtccggcc	cgtggacaag	tggaataaac	atatcatcta	tttcgtcctc	180
accgaccgtt	tccaggatgg	cgacaagacc	aacaacatgg	acgtgggtccc	gacggacatg	240
aaaaaatatc	atggcggcga	catccagggg	ctcatcgaca	agctcgacta	tatcaaggag	300
accggttcga	cggccatctg	gctcacgccc	cctatgaagg	ggcagacca	cttcttcgag	360
accgacaatt	accatgggta	ctggcccat	gacttctatg	acacggaccc	ccatgtgggc	420
accatgcaga	aatttgagga	gcttatcgag	aaagcccatg	agaaagggct	gaagatcgtg	480
ctcgatattc	ccctgaacca	cacggcctgg	gagcatccct	tctacaagga	cgacagcaag	540
aaggactggg	tccaccatat	aggagatgtg	aaggactggg	aagatcccta	ctgggctgaa	600
aacggctcca	tattcggctc	tcctgacctg	gcgcaggaaa	accctgccgt	ggaaaagtac	660
ctcatcgacg	tggccaagtt	ctgggtagac	aagggtattg	acggcttcag	gcttgacgcc	720

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gtgaagaacg	tgccccctcaa	cttctggg	cg	aagtttgacc	ggg	cgattatgcg	780
ggcaaggact	tcctcctcgt	cg	gggaatac	tttgacggaa	acccggcgaa	agtcgcgaac	840
taccagagag	aggacatgag	ctcactcttc		gattaccgc	tctactggac	cctgaaggac	900
accttcgcca	aggacgggag	catgcgcaac	ctggcggcga	agcttgatga	gtgcgacagg		960
aattatccccg	acccgggcct	catgtcgggt	ttccttgata	accacgacac	gccgaggttc		1020
ctcaccgagg	ccaacggcaa	caaggataag	ctcaaactgg	ccctcgctt	cgcgatgacc		1080
atcaaccgca	tgcttaccat	ttattatggc	accgaggttg	ccatggaagg	caactgcat		1140
atcatggg	cgtagataa	ccggaggac	atgcagtggg	acaaggatcc	tgacatgttc		1200
aaatacttca	agactctcac	cactgcccgc	aatgagcatg	aatccctcag	ggaaggaaag		1260
aagctcgaga	tgtggcagga	tgacaaagtc	tacgcgtacg	ggaggcagac	cccgaaggac		1320
gagtcctatcg	tggtgcttaa	caacggctat	gatacgacag	aacgggacat	accgctccgc		1380
cccgaagagcg	gcatacaagaa	cggcacgggtg	ctgaaggatg	tcatacccg	cgaaaccgtg		1440
acggtacaga	acggaaaaat	ccatgcgaaa	tgcggcggca	aacaggcgcg	gatctacgtg		1500
cccgcgtag							1509

<210> 108
 <211> 502
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 108

Met	Asp	Ser	Leu	Asp	Ala	Pro	Glu	Gln	Lys	Pro	Trp	Val	Lys	Asp	Gly
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Arg	Leu	Ser	Ala	Tyr	Leu	Asp	Thr	Gly	Thr	Gly	Thr	Val	Val	Ala	Pro
			20					25					30		
Glu	Ala	Pro	Ala	Pro	Pro	Pro	Pro	Pro	Ala	Glu	Glu	Val	Arg	Pro	Val
			35				40					45			
Asp	Lys	Trp	Lys	Asn	Asp	Ile	Ile	Tyr	Phe	Val	Leu	Thr	Asp	Arg	Phe
	50					55					60				
Gln	Asp	Gly	Asp	Lys	Thr	Asn	Asn	Met	Asp	Val	Val	Pro	Thr	Asp	Met
65					70				75						80
Lys	Lys	Tyr	His	Gly	Gly	Asp	Ile	Gln	Gly	Leu	Ile	Asp	Lys	Leu	Asp
				85					90					95	
Tyr	Ile	Lys	Glu	Thr	Gly	Ser	Thr	Ala	Ile	Trp	Leu	Thr	Pro	Pro	Met
			100					105					110		
Lys	Gly	Gln	Thr	His	Phe	Phe	Glu	Thr	Asp	Asn	Tyr	His	Gly	Tyr	Trp
			115				120					125			
Pro	Ile	Asp	Phe	Tyr	Asp	Thr	Asp	Pro	His	Val	Gly	Thr	Met	Gln	Lys
	130					135					140				
Phe	Glu	Glu	Leu	Ile	Glu	Lys	Ala	His	Glu	Lys	Gly	Leu	Lys	Ile	Val
145					150				155						160
Leu	Asp	Ile	Pro	Leu	Asn	His	Thr	Ala	Trp	Glu	His	Pro	Phe	Tyr	Lys
				165					170					175	
Asp	Asp	Ser	Lys	Lys	Asp	Trp	Phe	His	His	Ile	Gly	Asp	Val	Lys	Asp
			180					185					190		
Trp	Glu	Asp	Pro	Tyr	Trp	Ala	Glu	Asn	Gly	Ser	Ile	Phe	Gly	Leu	Pro
		195					200					205			
Asp	Leu	Ala	Gln	Glu	Asn	Pro	Ala	Val	Glu	Lys	Tyr	Leu	Ile	Asp	Val
	210					215					220				
Ala	Lys	Phe	Trp	Val	Asp	Lys	Gly	Ile	Asp	Gly	Phe	Arg	Leu	Asp	Ala
225					230					235					240
Val	Lys	Asn	Val	Pro	Leu	Asn	Phe	Trp	Ala	Lys	Phe	Asp	Arg	Ala	Ile
			245						250					255	
His	Asp	Tyr	Ala	Gly	Lys	Asp	Phe	Leu	Leu	Val	Gly	Glu	Tyr	Phe	Asp
			260					265					270		
Gly	Asn	Pro	Ala	Lys	Val	Ala	Asn	Tyr	Gln	Arg	Glu	Asp	Met	Ser	Ser
		275					280					285			
Leu	phe	Asp	Tyr	Pro	Leu	Tyr	Trp	Thr	Leu	Lys	Asp	Thr	Phe	Ala	Lys
	290					295					300				
Asp	Gly	Ser	Met	Arg	Asn	Leu	Ala	Ala	Lys	Leu	Asp	Glu	Cys	Asp	Arg
305					310					315					320
Asn	Tyr	Pro	Asp	Pro	Gly	Leu	Met	Ser	Val	Phe	Leu	Asp	Asn	His	Asp
				325					330					335	
Thr	Pro	Arg	Phe	Leu	Thr	Glu	Ala	Asn	Gly	Asn	Lys	Asp	Lys	Leu	Lys
			340					345					350		

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Leu Ala Leu Ala Phe Ala Met Thr Ile Asn Arg Met Pro Thr Ile Tyr
 355 360 365
 Tyr Gly Thr Glu Val Ala Met Gly Asn Cys Asp Ile Met Gly Ala
 370 375 380
 Val Asp Asn Arg Arg Asp Met Gln Trp Asp Lys Asp Pro Asp Met Phe
 385 390 395 400
 Lys Tyr Phe Lys Thr Leu Thr Thr Ala Arg Asn Glu His Glu Ser Leu
 405 410 415
 Arg Glu Gly Lys Lys Leu Glu Met Trp Gln Asp Asp Lys Val Tyr Ala
 420 425 430
 Tyr Gly Arg Gln Thr Pro Lys Asp Glu Ser Ile Val Val Leu Asn Asn
 435 440 445
 Gly Tyr Asp Thr Gln Glu Arg Asp Ile Pro Leu Arg Pro Glu Ser Gly
 450 455 460
 Ile Lys Asn Gly Thr Val Leu Lys Asp Val Ile Thr Gly Glu Thr Val
 465 470 475 480
 Thr Val Gln Asn Gly Lys Ile His Ala Lys Cys Gly Gly Lys Gln Ala
 485 490 495
 Arg Ile Tyr Val Pro Ala
 500

<210> 109
 <211> 1374
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 109
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 gttccggcaa aggcagaaac tctagagaat ggtggagtta taatgcaggc tttctattgg 120
 gatgttcttg gaggaggaat ctggtgggac acaatagctc aaaagatacc cgaatgggca 180
 agtgcaggaa tctcagcgat atggattcca ccagcgagta agggcatgag cgggtggtat 240
 tccatgggct acgatcccta cgatttcttt gacctcggcg agtactatca gaaggggaca 300
 gttgagacgc gcttcggctc aaaggaagaa ctggtgaaca tgataaacac cgcacactcc 360
 tacggcataa aggtgatagc ggacatagtc ataaaccacc gcgccggtgg agaccttgag 420
 tggaaacctt tcgtgaacga ctataacctgg acagacttct caaaagtcgc ctccggtaaa 480
 tatakggcca actaccttga cttccaccca aacgagcttc actgttgtga tgaagggtacc 540
 tttggaggat accctgatat atgtcacgac aaaagctggg accagtactg gctctgggcg 600
 agcagcgaag gctacgctgc ctacctcagg agcatagggg ttgacgcctg gcgtttcgac 660
 tacgtcaagg gctacggagc atgggttggt aacgactggc tcagctggtg gggaggctgg 720
 gccgttggag agtactggga cacgaacggt gatgcactcc tcaactgggc atacagcagc 780
 ggcgccaagg tctttgactt cccgctctac tacaagatgg acgaagcctt cgacaacacc 840
 aacatcccgg cattagtggg tgcactcaga tacggccaga cagtgggtcag ccgcgatccc 900
 ttcaaggcgg taactttcgt tgccaaccac gatacagata taatctggaa caagtatccg 960
 gcttatgcat tcatccttac ctatgagggg cagcctgtta tattctaccg cgactacgag 1020
 gagtggctca acaaggataa gcttaacaac ctcatctgga tacacgatca ccttgctgga 1080
 gggagtactg acattgttta ctacgacagc gacgagctta tctttgtgag aaacggctat 1140
 ggcaccaaac caggactgat aaccttatatc aacctcggct caagcaaagt tgggaagggtg 1200
 gtctacgttc caaagttcgc cggttcatgc atccacgagt acaccggcaa cctcggcggt 1260
 tggatagaca agtacgtctc ctccagcggc tgggtctatc ttgaggcccc agcccacgac 1320
 ccggcgaacg gctactacgg ctactctgtc tggagctact gcggtgtggg ttga 1374

<210> 110
 <211> 457
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 110
 Met Ala Arg Lys Thr Leu Ala Ile Phe Phe Val Leu Leu Val Leu Leu
 1 5 10 15
 Ser Leu Ser Ala Val Pro Ala Lys Ala Glu Thr Leu Glu Asn Gly Gly
 20 25 30
 Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Gly Ile Trp

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Trp	Asp	35	Thr	Ile	Ala	Gln	Lys	40	Ile	Pro	Glu	Trp	Ala	45	Ser	Ala	Gly	Ile
Ser	Ala	50	Ile	Trp	Ile	Pro	Pro	Ala	Ser	Lys	Gly	Met	Ser	60	Gly	Gly	Tyr	
Ser	Met	65	Gly	Tyr	Asp	Pro	Tyr	Asp	Phe	Phe	Asp	Leu	Gly	75	Glu	Tyr	Tyr	
Gln	Lys	85	Thr	Val	Glu	Thr	Arg	Phe	Gly	Ser	Lys	Glu	Glu	90	Leu	Val		
Asn	Met	100	Ile	Asn	Thr	Ala	His	Ser	Tyr	Gly	Ile	Lys	Val	105	Ile	Ala	Asp	
Ile	Val	115	Ile	Asn	His	Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	120	Asn	Pro	Phe	
Val	Asn	130	Asp	Tyr	Thr	Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	135	Ser	Gly	Lys	
Tyr	Thr	145	Ala	Asn	Tyr	Leu	Asp	Phe	His	Pro	Asn	Glu	Leu	140	His	Cys	Cys	
Asp	Glu	165	Gly	Thr	Phe	Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	170	Asp	Lys	Ser	
Trp	Asp	180	Gln	Tyr	Trp	Leu	Trp	Ala	Ser	Ser	Glu	Ser	Tyr	185	Ala	Ala	Tyr	
Leu	Arg	195	Ser	Ile	Gly	Val	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	200	Val	Lys	Gly	
Tyr	Gly	210	Ala	Trp	Val	Val	Asn	Asp	Trp	Leu	Ser	Trp	Trp	215	Gly	Gly	Trp	
Ala	Val	225	Gly	Glu	Tyr	Trp	Asp	Thr	Asn	Val	Asp	Ala	Leu	230	Leu	Asn	Trp	
Ala	Tyr	245	Ser	Ser	Gly	Ala	Lys	Val	Phe	Asp	Phe	Pro	Leu	250	Tyr	Tyr	Lys	
Met	Asp	260	Glu	Ala	Phe	Asp	Asn	Thr	Asn	Ile	Pro	Ala	Leu	265	Val	Asp	Ala	
Leu	Arg	275	Tyr	Gly	Gln	Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	280	Lys	Ala	Val	
Thr	Phe	290	Val	Ala	Asn	His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	295	Lys	Tyr	Pro	
Ala	Tyr	305	Ala	Phe	Ile	Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Val	310	Ile	Phe	Tyr	
Arg	Asp	325	Tyr	Glu	Trp	Leu	Asn	Lys	Asp	Lys	Leu	Asn	Asn	330	Leu	Ile		
Trp	Ile	340	His	Asp	His	Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	345	Val	Tyr	Tyr	
Asp	Ser	355	Asp	Glu	Leu	Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	360	Thr	Lys	Pro	
Gly	Leu	370	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	Ser	Ser	Lys	Val	375	Gly	Arg	Trp	
Val	Tyr	385	Val	Pro	Lys	Phe	Ala	Gly	Ser	Cys	Ile	His	Glu	390	Tyr	Thr	Gly	
Asn	Leu	405	Gly	Trp	Ile	Asp	Lys	Tyr	Val	Ser	Ser	Ser	Gly	410	Trp	Val		
Tyr	Leu	420	Glu	Ala	Pro	Ala	His	Asp	Pro	Ala	Asn	Gly	Tyr	425	Gly	Tyr		
Ser	Val	435	Trp	Ser	Tyr	Cys	Gly	Val	Gly					440				
		450												455				

<210> 111
 <211> 1416
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 111	
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ttggctgtgg cttcgatagg cctcctctcg actccagtgg gtgctgccaa gtactccgaa	120
ctcgaagagg gcggtgttat aatgcaggcc ttctactggg acgtccctac cgggtgggatc	180
tggtgggaca ccataagaca gaaaatcccg gagtggtagc acgctggaat ctcggcgata	240
tggtattctc cagctagcaa aggtatgggt ggtgcatact ccatgggtta tgaccctac	300

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```

gattttctttg acctcggcga gtactatcag aagggaaacag ttgagacgcg cttcgggtca 360
aaggaggaac tggatgaacat gataaacacc gcacactcct atggcataaa ggtgatagcg 420
gacatagtca taaaccaccg cgccggcggc gacctggagt ggaacccctt tgtaacaac 480
tatacttggga cagactttctc caaggctcgcc tccggtaaat acacggccaa ctaccttgac 540
ttccacccaa acgaggtcaa gtgctgcatg gagggtagat ttggtgactt tccggacatc 600
gcccacgaga agagctggga tcagtactgg ctctgggcaa gcaatgagag ctacgccgcc 660
tatctccgga gcatagggat cgatgcatgg cgtttcgact acgtcaaagg ttacggagcg 720
tgggttggtta acgactggct cagctgggtg ggaggttggg ccgttggaga gtactgggac 780
accaacgttg atgcactcct taactgggca tacaacagcg gtgccaagggt ctttgacttc 840
ccgctctact acaagatgga cgaagccttt gacaacacca acatccccgc tttggtttac 900
gccctccaga acggaggaac agtcgtttcc cgcgatccct tcaaggcagt aactttcgtt 960
gccaaccacg ataccgatat aatctggaac aagtatccgg cttatgctgt catccttacc 1020
tatgaggggac agcctgttat attctaccgc gactacgagg agtggctcaa caaggataag 1080
cttaacaacc ttatctggat acacgagcac cttgccggag gaagtaccaa gatcctctac 1140
tacgataacg atgagctaata attcatgagg gagggctacg ggagcaagcc gggcctcata 1200
acctacataa acctcggaaa cgactggggc gagcgctggg tgaacgtcgg ctcaaagttt 1260
gccggctaca caatccatga atacacaggc aatctcgggt gctgggttga caggtgggtt 1320
cagtacgacg gatgggttaa actgacggca cctcctcacg atccagccaa cggatattac 1380
ggctactcag tctggagcta cgcaggcgctc ggatga 1416

```

<210> 112
 <211> 471
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 112
 Met Pro Ala Phe Lys Ser Lys Val Met His Met Lys Leu Lys Tyr Leu
 1 5 10 15
 Ala Leu Val Leu Ala Val Ala Ser Ile Gly Leu Leu Ser Thr Pro
 20 25 30
 Val Gly Ala Ala Lys Tyr Ser Glu Leu Glu Glu Gly Gly Val Ile Met
 35 40 45
 Gln Ala Phe Tyr Trp Asp Val Pro Thr Gly Gly Ile Trp Trp Asp Thr
 50 55 60
 Ile Arg Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile
 65 70 75 80
 Trp Ile Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly
 85 90 95
 Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Tyr Gln Lys Gly
 100 105 110
 Thr Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile
 115 120 125
 Asn Thr Ala His Ser Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile
 130 135 140
 Asn His Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Val Asn Asn
 145 150 155 160
 Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala
 165 170 175
 Asn Tyr Leu Asp Phe His Pro Asn Glu Val Lys Cys Cys Asp Glu Gly
 180 185 190
 Thr Phe Gly Asp Phe Pro Asp Ile Ala His Glu Lys Ser Trp Asp Gln
 195 200 205
 Tyr Trp Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala Tyr Leu Arg Ser
 210 215 220
 Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala
 225 230 235 240
 Trp Val Val Asn Asp Trp Leu Ser Trp Trp Gly Gly Trp Ala Val Gly
 245 250 255
 Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Asn
 260 265 270
 Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu
 275 280 285
 Ala Phe Asp Asn Thr Asn Ile Pro Ala Leu Val Tyr Ala Leu Gln Asn
 290 295 300
 Gly Gly Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val

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```

305      310      315      320
Ala Asn His Asp Thr Asp Ile Ile Trp Asn Lys Tyr Pro Ala Tyr Ala
      325      330      335
Phe Ile Leu Thr Tyr Glu Gly Gln Pro Val Ile Phe Tyr Arg Asp Tyr
      340      345      350
Glu Glu Trp Leu Asn Lys Asp Lys Leu Asn Asn Leu Ile Trp Ile His
      355      360      365
Glu His Leu Ala Gly Gly Ser Thr Lys Ile Leu Tyr Tyr Asp Asn Asp
      370      375      380
Glu Leu Ile Phe Met Arg Glu Gly Tyr Gly Ser Lys Pro Gly Leu Ile
385      390      395      400
Thr Tyr Ile Asn Leu Gly Asn Asp Trp Ala Glu Arg Trp Val Asn Val
      405      410      415
Gly Ser Lys Phe Ala Gly Tyr Thr Ile His Glu Tyr Thr Gly Asn Leu
      420      425      430
Gly Gly Trp Val Asp Arg Trp Val Gln Tyr Asp Gly Trp Val Lys Leu
      435      440      445
Thr Ala Pro Pro His Asp Pro Ala Asn Gly Tyr Tyr Gly Tyr Ser Val
450      455      460
Trp Ser Tyr Ala Gly Val Gly
465      470

```

<210> 113
 <211> 1539
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 113
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ttgctgcctc attctgcagc agcggcggca aatcttaatg ggacgctgat gcagtatttt 120
gaatggtaca tgcccaatga cggccaacat tggaagcgct tgcaaaacga ctcggcatat 180
ttggctgaac acggtattac tgccgtctgg attcccccg catataaggg aacgagccaa 240
gcggatgtgg gctacggtgc ttacgacctt tatgatttag gggagtttca tcaaaaaggg 300
acggttcgga caaagtacgg cacaaaagga gagctgcaat ctgcgatcaa aagtcttcat 360
tcccgcgaca ttaacgttta cggggatgtg gtcatcaacc acaaaggcgg cgctgatgcg 420
accgaagatg taaccgcggt tgaagtcgat cccgctgacc gcaaccgcgt aatttcagga 480
gaacaccgaa ttaaagcctg gacacatttt cattttccgg ggcgcggcag cacatacagc 540
gattttaaat ggcattggta ccattttgac ggaaccgatt gggacgagtc ccgaaagctg 600
aaccgcatct ataagtttca aggaaaggct tgggattggg aagttttcaa tgaaaacggc 660
aactatgatt atttgatgta tgccgacatc gattatgacc atcctgatgt cgcagcagaa 720
attaagagat ggggcacttg gtatgccaat gaactgcaat tggacggttt ccgtcttgat 780
gctgtcaaac acattaaatt ttcttttttg cgggattggg ttaatcatgt cagggaaaaa 840
acggggaagg aaatgtttac ggtagctgaa tattggcaga atgacttggg cgcgctggaa 900
aactatttga acaaaacaaa ttttaatcat tcagtgtttg acgtgccgct tcattatcag 960
ttccatgctg catcgacaca gggaggcggc tatgatatga ggaaattgct gaacggtagc 1020
gtcgtttcca agcatccgtt gaaagcgggt acatttgcgc ataaccatga tacacagccg 1080
gggcaatcgc ttgagtcgac tgtccaacaa tggtttaagc cgcttgctta cgctttcatt 1140
ctcacaaggg aatctggata ccctcagggt ttctacgggg atatgtacgg gacgaaagga 1200
gactcccagc gcgaaattcc tgccttgaaa cacaaaattg aaccgatctt aaaagcgaga 1260
aaacagtatg cgtacggagc acagcatgat tatttcgacc accatgacat tgtcggctgg 1320
acaagggaag gcgacagctc ggttgcaaat tcaggtttgg cggcattaat aacagacgga 1380
cccgggtggg caaagcgaat gtatgtcggc cggcaaaacg ccggtgagac atggcatgac 1440
attaccggaa accgttcgga gccggttgtc atcaattcgg aaggctgggg agagtttcac 1500
gtaaacggcg ggtcggtttc aatttatgtt caaagatag 1539

```

<210> 114
 <211> 512
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

```

<400> 114
Met Lys Gln Gln Lys Arg Leu Tyr Ala Arg Leu Leu Thr Leu Leu Phe

```

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1	Ala	Leu	Ile	20	5	Leu	Leu	Pro	His	25	10	Ala	Ala	Ala	Ala	30	15	Asn	Leu
Asn	Gly	Thr	35	Leu	Met	Gln	Tyr	Phe	40	Glu	Trp	Tyr	Met	Pro	45	Asn	Asp	Gly	
Gln	His	Trp	50	Lys	Arg	Leu	Gln	Asn	55	Asp	Ser	Ala	Tyr	60	Leu	Ala	Glu	His	
Gly	Ile	Thr	65	Ala	Val	Trp	Ile	Pro	70	Pro	Ala	Tyr	Lys	75	Gly	Thr	Ser	Gln	
Ala	Asp	Val	85	Gly	Tyr	Gly	Ala	Tyr	90	Asp	Leu	Tyr	Asp	95	Leu	Gly	Glu	Phe	
His	Gln	Lys	100	Gly	Thr	Val	Arg	Thr	105	Lys	Tyr	Gly	Thr	110	Lys	Gly	Glu	Leu	
Gln	Ser	Ala	115	Ile	Lys	Ser	Leu	His	120	Ser	Arg	Asp	Ile	125	Asn	Val	Tyr	Gly	
Asp	Val	Val	130	Ile	Asn	His	Lys	Gly	135	Gly	Ala	Asp	Ala	140	Thr	Glu	Asp	Val	
Thr	Ala	Val	145	Glu	Val	Asp	Pro	Ala	150	Asp	Arg	Asn	Arg	155	Val	Ile	Ser	Gly	
Glu	His	Arg	165	Ile	Lys	Ala	Trp	Thr	170	His	Phe	His	Phe	175	Pro	Gly	Arg	Gly	
Ser	Thr	Tyr	180	Ser	Asp	Phe	Lys	Trp	185	His	Trp	Tyr	His	190	Phe	Asp	Gly	Thr	
Asp	Trp	Asp	195	Glu	Ser	Arg	Lys	Leu	200	Asn	Arg	Ile	Tyr	205	Lys	Phe	Gln	Gly	
Lys	Ala	Trp	210	Asp	Trp	Glu	Val	Ser	215	Asn	Glu	Asn	Gly	220	Asn	Tyr	Asp	Tyr	
Leu	Met	Tyr	225	Ala	Asp	Ile	Asp	Tyr	230	Asp	His	Pro	Asp	235	Val	Ala	Ala	Glu	
Ile	Lys	Arg	245	Trp	Gly	Thr	Trp	Tyr	250	Ala	Asn	Glu	Leu	255	Gln	Leu	Asp	Gly	
Phe	Arg	Leu	260	Asp	Ala	Val	Lys	His	265	Ile	Lys	Phe	Ser	270	Phe	Leu	Arg	Asp	
Trp	Val	Asn	275	His	Val	Arg	Glu	Lys	280	Thr	Gly	Lys	Glu	285	Met	Phe	Thr	Val	
Ala	Glu	Tyr	290	Trp	Gln	Asn	Asp	Leu	295	Gly	Ala	Leu	Glu	300	Asn	Tyr	Leu	Asn	
Lys	Thr	Asn	305	Phe	Asn	His	Ser	Val	310	Phe	Asp	Val	Pro	315	Leu	His	Tyr	Gln	
Phe	His	Ala	325	Ala	Ser	Thr	Gln	Gly	330	Gly	Gly	Tyr	Asp	335	Met	Arg	Lys	Leu	
Leu	Asn	Gly	340	Thr	Val	Val	Ser	Lys	345	His	Pro	Leu	Lys	350	Ala	Val	Thr	Phe	
Val	Asp	Asn	355	His	Asp	Thr	Gln	Pro	360	Gly	Gln	Ser	Leu	365	Glu	Ser	Thr	Val	
Gln	Thr	Trp	370	Phe	Lys	Pro	Leu	Ala	375	Tyr	Ala	Phe	Ile	380	Leu	Thr	Arg	Glu	
Ser	Gly	Tyr	385	Pro	Gln	Val	Phe	Tyr	390	Gly	Asp	Met	Tyr	395	Gly	Thr	Lys	Gly	
Asp	Ser	Gln	405	Arg	Glu	Ile	Pro	Ala	410	Leu	Lys	His	Lys	415	Ile	Glu	Pro	Ile	
Leu	Lys	Ala	420	Arg	Lys	Gln	Tyr	Ala	425	Tyr	Gly	Ala	Gln	430	His	Asp	Tyr	Phe	
Asp	His	His	435	Asp	Ile	Val	Gly	Trp	440	Thr	Arg	Glu	Gly	445	Asp	Ser	Ser	Val	
Ala	Asn	Ser	450	Gly	Leu	Ala	Ala	Leu	455	Ile	Thr	Asp	Gly	460	Pro	Gly	Gly	Ala	
Lys	Arg	Met	465	Tyr	Val	Gly	Arg	Gln	470	Asn	Ala	Gly	Glu	475	Thr	Trp	His	Asp	
Ile	Thr	Gly	485	Asn	Arg	Ser	Glu	Pro	490	Val	Val	Ile	Asn	495	Ser	Glu	Gly	Trp	
Gly	Glu	Phe	500	His	Val	Asn	Gly	Gly	505	Ser	Val	Ser	Ile	510	Tyr	Val	Gln	Arg	

<210> 115
 <211> 1338
 <212> DNA
 <213> Eukaryote

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<400> 115
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gttccggagg gaggaatctg gtgggacaca atacggcaga agatccctga atggtacgat      120
gcaggcatat ccgccatctg gatacccccg gcgagcaagg gcatgggcgg ggcctactcg      180
atgggctacg acccctacga ttacttcgat ctgggcgagt tttaccagaa gggaaccgtt      240
gagacccgct tcggctccaa ggaagagctc gtcaacatga tctccacggc ccaccagtac      300
ggcatcaagg ttatagcgga catagtgata aaccaccgcg caggtggaga cctcgaatgg      360
aacccatacg tcggcgacta tacctggacg gacttttcta aggtcgctc cgggaaatac      420
aaggcccaact acatggactt ccatccaaac aactacagca cctcagacga gggaaccttc      480
ggtggcttcc cagacattga tcacctcgtg cccttcaacc agtactggct gtgggcgagc      540
aacgagagct acgccgccta cctcaggagc atagggatcg atgctggcg ctttgactac      600
gttaagggct acggcgcggt ggtcgtcaag gactggctga gtcagtgggg cggctgggcc      660
gtcggcgagt actgggacac caacgtcgat gcgctcctca actgggccta cagcagcggc      720
gccaagggtct tcgacttccc gctctactac aagatggacg aggcctttga caacaagaac      780
attcccgccc tcgtttacgc catccagaac ggtgaaaccg tcgtcagcag ggatcccttc      840
aaggccgtta ccttcgtggc taaccacgat acgaacataa tctggaacaa gtaccctgcc      900
tatgccttca tcctgaccta cgaaggtcag cccgtcatct tctaccgcga ctacgaggag      960
tggctcaaca aggacaaact caacaacctc atatggattc acgagcacct ggcaggggga      1020
agcaccaaga tccttacta cgacgacgat gagctcatct tcatgagggg aggtctacggc      1080
gcagggcccc ggcttataac ctacatcaac ctcggtagcg actgggcgga gagatgggtg      1140
aacgttggct caaagttcgc gggctataca atccacgaat acaccggaaa cctcggcggc      1200
tgggtcgaca ggtacgtcca gtacgacggc tgggtcaagc ttaccgctcc gccacacgat      1260
ccggcaaacg gctattacgg ctactcggtc tggagctacg cggagttgg aagatctcat      1320
caccatcacc atcactaa
1338

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<210> 116
<211> 445
<212> PRT
<213> Eukaryote

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<400> 116
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Phe Tyr Trp Asp Val Pro Glu Gly Gly Ile Trp Trp Asp Thr Ile Arg
          20          25          30
Gln Lys Ile Pro Glu Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile
          35          40          45
Pro Pro Ala Ser Lys Gly Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp
          50          55          60
Pro Tyr Asp Tyr Phe Asp Leu Gly Glu Phe Tyr Gln Lys Gly Thr Val
          65          70          75          80
Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val Asn Met Ile Ser Thr
          85          90          95
Ala His Gln Tyr Gly Ile Lys Val Ile Ala Asp Ile Val Ile Asn His
          100          105          110
Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Tyr Val Gly Asp Tyr Thr
          115          120          125
Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Lys Ala His Tyr
          130          135          140
Met Asp Phe His Pro Asn Asn Tyr Ser Thr Ser Asp Glu Gly Thr Phe
          145          150          155          160
Gly Gly Phe Pro Asp Ile Asp His Leu Val Pro Phe Asn Gln Tyr Trp
          165          170          175
Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala Tyr Leu Arg Ser Ile Gly
          180          185          190
Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly Tyr Gly Ala Trp Val
          195          200          205
Val Lys Asp Trp Leu Ser Gln Trp Gly Gly Trp Ala Val Gly Glu Tyr
          210          215          220
Trp Asp Thr Asn Val Asp Ala Leu Leu Asn Trp Ala Tyr Ser Ser Gly
          225          230          235          240
Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys Met Asp Glu Ala Phe
          245          250          255
Asp Asn Lys Asn Ile Pro Ala Leu Val Tyr Ala Ile Gln Asn Gly Glu
          260          265          270
Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn
          275          280          285

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His	Asp	Thr	Asn	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
	290					295					300				
Leu	Thr	Tyr	Glu	Gly	Gln	Pro	Val	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu
305				310						315					320
Trp	Leu	Asn	Lys	Asp	Lys	Leu	Asn	Asn	Leu	Ile	Trp	Ile	His	Glu	His
				325					330					335	
Leu	Ala	Gly	Gly	Ser	Thr	Lys	Ile	Leu	Tyr	Tyr	Asp	Asp	Asp	Glu	Leu
			340					345					350		
Ile	Phe	Met	Arg	Glu	Gly	Tyr	Gly	Asp	Arg	Pro	Gly	Leu	Ile	Thr	Tyr
		355					360					365			
Ile	Asn	Leu	Gly	Ser	Asp	Trp	Ala	Glu	Arg	Trp	Val	Asn	Val	Gly	Ser
	370					375					380				
Lys	Phe	Ala	Gly	Tyr	Thr	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly
385					390					395					400
Trp	Val	Asp	Arg	Tyr	Val	Gln	Tyr	Asp	Gly	Trp	Val	Lys	Leu	Thr	Ala
				405					410					415	
Pro	Pro	His	Asp	Pro	Ala	Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser
			420					425					430		
Tyr	Ala	Gly	Val	Gly	Arg	Ser	His	His	His	His	His	His			
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<210> 117
 <211> 1476
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 117

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caag	acag	ggg	acata	aca	aat	gaa	acac	aca	gcg	gga	atgc	tgg	cga	tcgc	agg	tat	gctg	120
atc	gcccc	ct	tgg	cgc	atgc	cgat	gtc	ata	ctgc	acgc	ct	tca	act	ggaa	ata	cagt	gaa	180
gtc	acc	gcca	agg	ccg	atct	cat	caagg	ct	gcc	ggt	taca	agc	agg	tgt	cat	ctc	accg	240
cct	ctg	aagt	cct	cgg	gcaa	cgag	tgg	tgg	gct	cgt	tacc	agc	ccc	cagg	tct	gcgc	ctg	300
gtc	gac	accc	ccct	tgg	caa	caag	cagg	at	ctg	gag	cagc	tga	tcg	ccgc	gat	gcag	acc	360
cg	ggg	cattg	ccgt	ctac	gc	ggac	gtg	gtg	ctc	aacc	aca	tgg	cca	acga	aag	ctg	gaag	420
cg	cag	cga	cc	tca	ctacc	cgg	cag	cga	ctg	ctg	caaa	gct	acg	ccg	caat	ccg	gcc	480
tact	ttg	aac	gcc	aga	aag	ct	ttt	ggc	gat	ctg	ggg	caga	act	tcct	cgc	cgg	cagg	540
ttt	cat	ccg	aggg	gtg	cat	cacc	gact	gg	aaca	atcc	cg	gcc	atg	tcca	gtact	ggc	ga	600
ctg	tgc	ggc	gg	ctg	ga	caag	ggg	ctg	ccg	gat	ctg	accc	ca	aaca	ctg	gg	tgg	660
aacc	agca	ac	agg	ctt	ac	gc	agg	cgtc	aagg	ggg	atgg	ggat	ca	aagg	ttt	tcg	gg	720
gat	gcg	tca	agca	cat	gag	cgatt	tacc	ag	atca	acgc	cg	tgt	tca	cccc	cgag	atca	aa	780
cagg	ggat	gc	acgt	cttt	gg	cgag	gtgat	c	acc	acg	ggg	gcg	ccg	gcaa	cag	cga	ctat	840
gaga	act	tcc	tcaa	acc	cta	cct	cga	cgc	agc	ggc	cagg	ggg	cct	acga	ctt	ccc	gctc	900
ttc	gc	ctcc	tgc	gtg	gag	gct	ggg	ctac	ggc	ggc	agca	tga	acct	gct	ggc	gat	ccc	960
ggt	gc	ctat	gtc	agg	cg	gccc	gtag	c	cg	cgc	gtca	cct	tcg	ccat	cacc	cac	gac	1020
at	cccc	acca	acg	acg	gttt	ccg	ctacc	ag	atc	ctca	aacc	agacc	gacga	gag	act	ggcc		1080
tat	gc	ctacc	tgc	tcg	gtcg	cgat	ggc	gg	tcg	ctct	tgg	tct	act	ccga	tcac	ggt	gaa	1140
acc	agg	gaca	agg	acg	gatt	gcg	ctgg	cag	gact	act	atc	tg	cga	caccga	tct	caa	agg	1200
at	gat	ccg	ct	tca	taac	agt	gc	agg	ca	acc	gat	gc	ag	ctcat	cgg	cag	taac	1260
tg	ctt	ctg	ct	caag	cg	tgg	caag	cag	ggc	gtg	gtcg	gc	atca	acaa	gtg	cga	ctac	1320
gag	cagg	gag	act	gg	ctcga	tacc	gcc	caga	ttc	gag	atga	act	gg	tatcg	caact	tacc	cg	1380
gat	gtg	ctcg	acc	aga	atgc	cgt	ggt	caac	gtg	cag	agcc	agt	ggg	taag	gct	ga	ccatc	1440
ccg	gccc	cg	gcg	ccaga	at	gtg	gct	gcag	gag	tga								1476

<210> 118
 <211> 491
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 118

Met	Arg	Val	Phe	Leu	Val	Val	Pro	Lys	Leu	Ser	Arg	Pro	Phe	Gln	Ala
1				5					10					15	
Glu	Ser	Gln	Gln	Gln	Asp	Arg	Asp	Ile	Thr	Met	Lys	His	Thr	Ala	Gly

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Met	Leu	Ala	20	Ile	Ala	Gly	Met	Leu	25	Ile	Ala	Pro	Leu	30	Ala	Asp
Val	Ile	Leu	35	His	Ala	Phe	Asn	Trp	40	Lys	Tyr	Ser	Glu	45	Val	Thr
Ala	Asp	Leu	50	Ile	Lys	Ala	Ala	Gly	55	Tyr	Lys	Gln	Val	60	Leu	Ile
Pro	Leu	Lys	65	Ser	Ser	Gly	70	Asn	75	Glu	Trp	Trp	Ala	80	Arg	Tyr
Asp	Leu	Arg	85	Val	Val	Asp	Thr	Pro	90	Leu	Gly	Asn	Lys	95	Gln	Pro
Gln	Leu	Ile	100	Ala	Ala	Met	Gln	Thr	105	Arg	Gly	Ile	Ala	110	Val	Tyr
Val	Val	Leu	115	Asn	His	Met	Ala	Asn	120	Glu	Ser	Trp	Lys	125	Arg	Ser
Asn	Tyr	Pro	130	Gly	Ser	Glu	Leu	Leu	135	Gln	Ser	Tyr	Ala	140	Gly	Asn
Tyr	Phe	Glu	145	Arg	Gln	Lys	Leu	Phe	150	Gly	Asp	Leu	Gly	155	Gln	Asn
Ala	Gly	Gln	165	Asp	Phe	His	Pro	Glu	170	Gly	Cys	Ile	Thr	175	Asp	Trp
Pro	Gly	His	180	Val	Gln	Tyr	Trp	Arg	185	Leu	Cys	Gly	Gly	190	Ala	Gly
Gly	Leu	Pro	195	Asp	Leu	Asp	Pro	Asn	200	Asn	Trp	Val	Val	205	Asn	Gln
Ala	Tyr	Leu	210	Gln	Ala	Leu	Lys	Gly	215	Met	Gly	Ile	Lys	220	Gly	Phe
Asp	Ala	Val	225	Lys	His	Met	Ser	Asp	230	Tyr	Gln	Ile	Asn	235	Ala	Val
Pro	Glu	Ile	245	Lys	Gln	Gly	Met	His	250	Val	Phe	Gly	Glu	255	Val	Ile
Gly	Gly	Ala	260	Gly	Asn	Ser	Asp	Tyr	265	Glu	Asn	Phe	Leu	270	Lys	Pro
Asp	Ser	Ser	275	Gly	Gln	Gly	Ala	Tyr	280	Asp	Phe	Pro	Leu	285	Phe	Ala
Arg	Gly	Ala	290	Leu	Gly	Tyr	Gly	Gly	295	Ser	Met	Asn	Leu	300	Leu	Ala
Gly	Ala	Tyr	305	Gly	Gln	Ala	Leu	Pro	310	Gly	Ser	Arg	Ala	315	Val	Thr
Ile	Thr	His	325	Asp	Ile	Pro	Thr	Asn	330	Asp	Gly	Phe	Arg	335	Tyr	Gln
Asn	Gln	Thr	340	Asp	Glu	Arg	Leu	Ala	345	Tyr	Ala	Tyr	Leu	350	Leu	Gly
Gly	Gly	Ser	355	Pro	Leu	Val	Tyr	Ser	360	Asp	His	Gly	Glu	365	Thr	Arg
Asp	Gly	Leu	370	Arg	Trp	Gln	Asp	Tyr	375	Tyr	Leu	Arg	Thr	380	Asp	Leu
Met	Ile	Arg	385	Phe	His	Asn	Thr	Val	390	Gln	Gly	Gln	Pro	395	Met	Lys
Gly	Ser	Asn	400	Asp	Cys	Phe	Val	Leu	405	Phe	Lys	Arg	Gly	410	Lys	Gln
Val	Gly	Ile	415	Asn	Lys	Cys	Asp	Tyr	420	Glu	Gln	Glu	Tyr	425	Trp	Leu
Ala	Arg	Phe	430	Glu	Met	Asn	Trp	Tyr	435	Arg	Asn	Tyr	Arg	440	Asp	Val
Gln	Asn	Ala	445	Val	Val	Asn	Val	Gln	450	Ser	Gln	Trp	Val	455	Arg	Leu
Pro	Ala	Arg	460	Gly	Ala	Arg	Met	Trp	465	Leu	Gln	Glu		470		Thr
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									490							

<210> 119
 <211> 1695
 <212> DNA
 <213> Bacterial

<400> 119
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60
 120

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tcttcactag	gtatcacagc	actatggctc	cctccagcat	ataaaggaa	gagccaaagc	300
gatgtcggat	acgggtgttta	cgatttatat	gaccttgggg	aattttaatca	aaaagggacg	360
atccgaacga	aatacggaa	aaaaacacaa	tatattcaag	ccattcaaac	tgcccaagcc	420
gcagggatgc	aagtatatgc	ggatgttgta	tttaatcata	aggcaggggc	tgacagtaca	480
gaatttgtcg	atgcagttga	ggtaaaccct	tctaatcgaa	atcaagaaac	atctggcaca	540
tatcaaatc	aagcatggac	aaaatttgat	tttcctggtc	gtggaacac	atactccagc	600
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gaatttaaaaa	actggggaac	gtgggtacgtc	aatactacaa	atatcgatgg	attccgctta	840
gatgccgtaa	aacatatttaa	atacagcttt	ttccctgact	ggctaacata	tgtacgtaat	900
caaacaggaa	aaaattttatt	tgccgtttggg	gaatttttga	gctatgacgt	caataagctg	960
cataattaca	ttacaaaaaac	aaatgggtcg	atgtcattat	ttgatgcacc	cttgcataac	1020
aactttttata	ccgctttccaa	atcgagtggg	tatttttgaca	tgctgttattt	attgaataat	1080
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aaatacaata	tcccgggggt	aaaaagtaaa	atcgaccgcg	ttttaattgc	tcgtcgtgat	1320
tacgcttatg	gaacacacag	tgattacatt	gatcatcaag	acattatcgg	atggacacga	1380
gaaggcattg	atgcaaaacc	gaactctgga	ctggcggcctt	taattaccga	cggctcctggt	1440
ggaagtaaat	ggatgtatgt	cggtaaaaaag	catgccggga	aagtatttta	tgattttaact	1500
ggaaatcgaa	gtgacacagt	aacgattaat	gcggatgggt	ggggagaatt	taaagtaaac	1560
ggaggatccg	tctcaatttg	ggtggctaaa	acgtcaaacg	tcacattttac	agtcaataac	1620
gccacaacaa	caagcggaca	aaacgtatat	gttgtcggca	acattccaga	gctaggcaat	1680
tgctgcacgg	gttaa					1695

<210> 120
 <211> 564
 <212> PRT
 <213> Bacterial

<400> 120
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 Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val Ala Thr
 20 25 30
 Leu Ile Ile Ser Phe Leu Thr Pro Phe Ser Thr Ala Gln Ala Asn Thr
 35 40 45
 Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp Leu Pro
 50 55 60
 Asn Asp Gly Thr Leu Trp Thr Lys Val Lys Asn Glu Ala Thr Asn Leu
 65 70 75 80
 Ser Ser Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Gly
 85 90 95
 Thr Ser Gln Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu Tyr Asp Leu
 100 105 110
 Gly Glu Phe Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly Thr Lys
 115 120 125
 Thr Gln Tyr Ile Gln Ala Ile Gln Thr Ala Gln Ala Ala Gly Met Gln
 130 135 140
 Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala Asp Ser Thr
 145 150 155 160
 Glu Phe Val Asp Ala Val Glu Val Asn Pro Ser Asn Arg Asn Gln Glu
 165 170 175
 Thr Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp Phe Pro
 180 185 190
 Gly Arg Gly Asn Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe
 195 200 205
 Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile Tyr Lys
 210 215 220
 Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp Thr Glu Asn
 225 230 235 240
 Gly Asn Tyr Asp Tyr Leu Met Phe Ala Asp Leu Asp Met Asp His Pro
 245 250 255
 Glu Val Val Thr Glu Leu Lys Asn Trp Gly Thr Trp Tyr Val Asn Thr
 260 265 270

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Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile Lys Tyr
 275 280 285
 Ser Phe Phe Pro Asp Trp Leu Thr Tyr Val Arg Asn Gln Thr Gly Lys
 290 295 300
 Asn Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Val Asn Lys Leu
 305 310 315 320
 His Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu Phe Asp Ala
 325 330 335
 Pro Leu His Asn Asn Phe Tyr Thr Ala Ser Lys Ser Ser Gly Tyr Phe
 340 345 350
 Asp Met Arg Tyr Leu Leu Asn Asn Thr Leu Met Lys Asp Gln Pro Ser
 355 360 365
 Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln Pro Gly Gln Ser
 370 375 380
 Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Gln Leu Ala Tyr Ala Phe
 385 390 395 400
 Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly Asp Tyr
 405 410 415
 Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Gly Leu Lys Ser Lys Ile Asp
 420 425 430
 Pro Leu Leu Ile Ala Arg Arg Asp Tyr Ala Tyr Gly Thr Gln Arg Asp
 435 440 445
 Tyr Ile Asp His Gln Asp Ile Ile Gly Trp Thr Arg Glu Gly Ile Asp
 450 455 460
 Ala Lys Pro Asn Ser Gly Leu Ala Ala Leu Ile Thr Asp Gly Pro Gly
 465 470 475 480
 Gly Ser Lys Trp Met Tyr Val Gly Lys Lys His Ala Gly Lys Val Phe
 485 490 495
 Tyr Asp Leu Thr Gly Asn Arg Ser Asp Thr Val Thr Ile Asn Ala Asp
 500 505 510
 Gly Trp Gly Glu Phe Lys Val Asn Gly Gly Ser Val Ser Ile Trp Val
 515 520 525
 Ala Lys Thr Ser Asn Val Thr Phe Thr Val Asn Asn Ala Thr Thr Thr
 530 535 540
 Ser Gly Gln Asn Val Tyr Val Val Gly Asn Ile Pro Glu Leu Gly Asn
 545 550 555 560
 Cys Arg Thr Gly

<210> 121
 <211> 1556
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 121
 atgctcgccc tgtcgctcgg cggtcgcgcc atcgacgcgg gcccgacagg ccctcgcgctc 60
 gtggagccgc tgccgcagcg cccacgcgtt ccgcaggagt accgcgccag cggccacgcg 120
 gccgcccggc acgtgttcgt gcacctgttc gaggcgaagt ggccggacat cgcggaggaa 180
 tgcgagaacg tgctggggcc ggccggctac gaggcgggtgc aggtgtcgcc gccgcaggag 240
 cacttggtgc agcagggggc gccgtggttg cagcgggtacc agccgggtgag ctactcgggtg 300
 gcgctgagcc gcagcggcac gggcgtggag ttacagcaaca tgatcagccg gtgcaaggcc 360
 gccggcgctg acatctacgt ggacgccgtc atcaaccaca tgacggccgg tgcggggacg 420
 gggagcaacg gcaccgccta caccaagtac aactaccccg gcctgtacgc gcaggcggac 480
 ttacaccgc agtgcgcggt gggcgactac accagcgccg ccaacgtgca ggactgcaaa 540
 ctgctggggc tggctgacct gaacaccggc gcggccggcg tgcagcagaa gatcgcgga 600
 tacctgggtc cgctggcgcg gctgggcgtg gcgggttttc gcatcgacgc cgccaagcac 660
 atccagccgg tggaaactga cgccatcggt gaccgcgtga accagacgct ggccggcggag 720
 gggcgccgcg ttccctactg gttcgccgag gtgatcgaca acggcggcga gggggtgcgg 780
 cgcgagcact actacggcct gggatacggc accggcggcg ccgcggacat cacggagttc 840
 cgctacaagg gcgtgggcga caagtctctg ggcagcggcg gccagcggct ggtggacctg 900
 aagaacttct cggcgggtgac gtggaacctg atgccgtcgg acaaggccgt cgtctttctg 960
 gagaaccacg atacgcagcg cggcggcgcc atcggctacc gcgatggcac ggcgttcagg 1020
 ctggccaacg tgtggtgctt ggcgcagccg tacggctatc cgtcgggtgat gtccagctac 1080
 gcctttgacc gcacctccc ctttgccgcg gacgcggccc cgccctccga ggacggcgcg 1140
 acgaaggacg tgacgtgcgc gccacgcgtg gagacggcgg tgctgggcac ctgggtgtgc 1200

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gagcaccg	cgccccgt	cat	tcagcggat	gtgggctt	tcgccg	cgat	ggcggg	cacg	1260	
gacctga	aacc	gctgg	ggga	caacgg	cggc	aacgcc	attg	ccttttc	cg	1320
ggcttc	g	cg	ccatcag	ccg	cgagcc	gaag	gtgacc	atgg	cg	1380
tccccgg	ca	cctact	gcga	cgtgct	gacc	ggcgg	caagg	tgggca	acgc	1440
accagc	gtga	cggtc	gactc	tcaggg	cgtg	gtgcag	ctga	gc	atcgtcga	1500
ctggtg	atcc	acctc	ggggc	caagct	gtaa	cggcgc	gctg	gcggat	gtgc	1556
								ggaggg		

<210> 122
 <211> 517
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 122

Met	Leu	Ala	Leu	Ser	Leu	Gly	Gly	Cys	Gly	Ile	Asp	Ala	Gly	Pro	Thr
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Gly	Pro	Arg	Val	Val	Glu	Pro	Leu	Pro	Gln	Arg	Pro	Thr	Leu	Pro	Gln
			20					25					30		
Glu	Tyr	Arg	Ala	Ser	Gly	His	Ala	Ala	Ala	Gly	Asp	Val	Phe	Val	His
		35					40					45			
Leu	Phe	Glu	Trp	Lys	Trp	Pro	Asp	Ile	Ala	Glu	Glu	Cys	Glu	Asn	Val
	50					55					60				
Leu	Gly	Pro	Ala	Gly	Tyr	Glu	Ala	Val	Gln	Val	Ser	Pro	Pro	Gln	Glu
65					70					75					80
His	Leu	Val	Gln	Gln	Gly	Ala	Pro	Trp	Trp	Gln	Arg	Tyr	Gln	Pro	Val
			85						90					95	
Ser	Tyr	Ser	Val	Ala	Leu	Ser	Arg	Ser	Gly	Thr	Gly	Val	Glu	Phe	Ser
			100					105					110		
Asn	Met	Ile	Ser	Arg	Cys	Lys	Ala	Ala	Gly	Val	Asp	Ile	Tyr	Val	Asp
		115					120					125			
Ala	Val	Ile	Asn	His	Met	Thr	Ala	Gly	Ala	Gly	Thr	Gly	Ser	Asn	Gly
	130					135					140				
Thr	Ala	Tyr	Thr	Lys	Tyr	Asn	Tyr	Pro	Gly	Leu	Tyr	Ala	Gln	Ala	Asp
145					150					155					160
Phe	His	Pro	Gln	Cys	Ala	Val	Gly	Asp	Tyr	Thr	Ser	Ala	Ala	Asn	Val
			165						170					175	
Gln	Asp	Cys	Glu	Leu	Leu	Gly	Leu	Ala	Asp	Leu	Asn	Thr	Gly	Ala	Ala
			180					185					190		
Gly	Val	Gln	Gln	Lys	Ile	Ala	Asp	Tyr	Leu	Val	Ser	Leu	Ala	Arg	Leu
		195					200					205			
Gly	Val	Ala	Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His	Ile	Gln	Pro	Val
	210					215					220				
Glu	Leu	Asp	Ala	Ile	Val	Asp	Arg	Val	Asn	Gln	Thr	Leu	Ala	Ala	Glu
225					230					235					240
Gly	Arg	Pro	Leu	Pro	Tyr	Trp	Phe	Ala	Glu	Val	Ile	Asp	Asn	Gly	Gly
			245						250					255	
Glu	Gly	Val	Arg	Arg	Glu	His	Tyr	Tyr	Gly	Leu	Gly	Tyr	Gly	Thr	Gly
			260					265					270		
Gly	Ala	Ala	Asp	Ile	Thr	Glu	Phe	Arg	Tyr	Lys	Gly	Val	Gly	Asp	Lys
		275					280					285			
Phe	Leu	Gly	Ser	Gly	Gly	Gln	Arg	Leu	Val	Asp	Leu	Lys	Asn	Phe	Ser
	290					295					300				
Ala	Val	Thr	Trp	Asn	Leu	Met	Pro	Ser	Asp	Lys	Ala	Val	Val	Phe	Leu
305					310					315					320
Glu	Asn	His	Asp	Thr	Gln	Arg	Gly	Gly	Gly	Ile	Gly	Tyr	Arg	Asp	Gly
				325					330					335	
Thr	Ala	Phe	Arg	Leu	Ala	Asn	Val	Trp	Met	Leu	Ala	Gln	Pro	Tyr	Gly
			340					345					350		
Tyr	Pro	Ser	Val	Met	Ser	Ser	Tyr	Ala	Phe	Asp	Arg	Thr	Ser	Pro	Phe
		355					360					365			
Gly	Arg	Asp	Ala	Gly	Pro	Pro	Ser	Glu	Asp	Gly	Ala	Thr	Lys	Asp	Val
	370					375					380				
Thr	Cys	Ala	Pro	Thr	Leu	Glu	Thr	Ala	Val	Leu	Gly	Thr	Trp	Val	Cys
385					390					395					400
Glu	His	Arg	Asp	Pro	Val	Ile	Gln	Arg	Met	Val	Gly	Phe	Arg	Arg	Ala
				405					410					415	

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Met Ala Gly Thr Asp Leu Asn Arg Trp Trp Asp Asn Gly Gly Asn Ala
 420 425 430
 Ile Ala Phe Ser Arg Gly Asp Arg Gly Phe Val Ala Ile Ser Arg Glu
 435 440 445
 Pro Lys Val Thr Met Ala Ala Val Pro Ser Gly Leu Ser Pro Gly Thr
 450 455 460
 Tyr Cys Asp Val Leu Thr Gly Gly Lys Val Gly Asn Ala Cys Ala Gly
 465 470 475
 Thr Ser Val Thr Val Asp Ser Gln Gly Val Val Gln Leu Ser Ile Val
 485 490 495
 Glu Asn Ser Ala Leu Val Ile His Leu Gly Ala Lys Leu Arg Arg Ala
 500 505 510
 Gly Gly Cys Ala Glu
 515

<210> 123
 <211> 1770
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 123
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 ctgcttggtc tcgtcttttc tgtcccaccc cgggcaatcc aggccagac aaccccggcc 120
 cgtaccggtta tggttcacct cttcgagtgg aaatggaccg acatcgctaa agaatgcbag 180
 aatttcctcg gaccgaaagg ctttgccgca atccagggtat cgccgcccc aaggatgctc 240
 caggggtcgc aatggtggac ccgctatcag ccggtcagct acaagatcga gagccgctcc 300
 ggcacccggg ccgagttcgc caatatggtc tcgctgtgca aagccgtcgg ggtcgatata 360
 tatgtcgatg ccgtgatcaa ccatatgacg actgtcggct ccggcactgg tatggctgga 420
 tcgacctaca ccagctacac ctatccgggg ctgtatcaga ccaggactt ccaccactgc 480
 gggcgcaatg gcaacgatga tatcagcagc tacggcgatc gctgggaagt acaaaactgc 540
 gaactgctca acctagccga cctcaacacc ggcgctgagt atgtccgggg taaactcgc 600
 gcctatatga acgatctgcg cggcctgggc gtcgcccgat ttcggatcga tgccgccaag 660
 cacatggata ccaacgacat caacaatatc gttggccgcc tgcccaacgc gccctacatc 720
 taccaggaag tgatcgacca gggcggcgag ccaattaccg ccggcgaata cttccagaat 780
 ggcgatgtga ccgagttcaa gtacagccgc gagatctcgc gcatgttcaa aaccggccag 840
 ctgacccata tgagccagtt cggcactgcc tggggcttca tgtccagcga cctggcagta 900
 gttttcaccg ataaccacga caaccagcgc ggtcacggcg gcgcccggcg tgtcttgacc 960
 taaaaagatg gccagctgta caccctgggc aatatcttcg agctagcctg gccgataggc 1020
 taccacaggg tcatgtcgag ctacacgttc agcaacggcg accagggggc gccatcgacc 1080
 aatgtgtacg caaccacaac gcctgattgt ggcaacggcc gctgggtctg tgagcaccgc 1140
 tggcgaggaa tcgccaacat ggtcgcgttc cgcaactaca ccgccccgac cttcagcacc 1200
 agcaactggg ggagcaacgg caacaaccag atcgctttca gccgcgggac cctgggcttt 1260
 gtggcgatca atcggaagg tggcagcctg aaccgcacct tccaaaccgg cctgcccgtc 1320
 ggacactact gcgatgtcat tcacggcgat ttcaatgcc a gcgcccggc ctgttcggc 1380
 ccaactatcg ctgtcaacgg ctccggacag gcaaccatca cgggtcaacgc gatggacgcg 1440
 gtggcgatct acggcggagc caggctcgcc actccggcca gtgtcaacgt gacattcaac 1500
 gaaaacgcca cgaccacctg ggggcagaat gtgtatatcg tcggcaacgt cgccgcccctg 1560
 ggcagctgga acgcaggcag cgcggtctta ctctcctccg ctaactacc aatctggagc 1620
 aagaccatcg ccctgccagc caacaccgcc attgagtaca agtacatcaa aaaggatggc 1680
 gcgggcaatg tgggtgagg aagcggcgcc aaccgcgtct ttaccacccc cggcagcggc 1740
 agtgccacgc gcaacgatac ctggaatatg 1770

<210> 124
 <211> 589
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 124
 Met Pro Gln Ala Ile Arg Thr Phe Ser Arg Trp Thr Leu Phe Gly Leu
 1 5 10 15
 Ile Gly Val Phe Leu Leu Gly Leu Val Phe Ser Val Pro Pro Arg Ala
 20 25 30

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Ile	Gln	Ala	Gln	Thr	Thr	Pro	Ala	Arg	Thr	Val	Met	Val	His	Leu	Phe
Glu	Trp	Lys	Trp	Thr	Asp	Ile	Ala	Lys	Glu	Cys	Glu	Asn	Phe	Leu	Gly
Pro	Lys	Gly	Phe	Ala	Ala	Ile	Gln	Val	Ser	Pro	Pro	Gln	Glu	His	Val
Gln	Gly	Ser	Gln	Trp	Trp	Thr	Arg	Tyr	Gln	Pro	Val	Ser	Tyr	Lys	Ile
Glu	Ser	Arg	Ser	Gly	Thr	Arg	Ala	Glu	Phe	Ala	Asn	Met	Val	Ser	Arg
Cys	Lys	Ala	Val	Gly	Val	Asp	Ile	Tyr	Val	Asp	Ala	Val	Ile	Asn	His
Met	Thr	Thr	Val	Gly	Ser	Gly	Thr	Gly	Met	Ala	Gly	Ser	Thr	Tyr	Thr
Ser	Tyr	Thr	Tyr	Pro	Gly	Leu	Tyr	Gln	Thr	Gln	Asp	Phe	His	His	Cys
Gly	Arg	Asn	Gly	Asn	Asp	Asp	Ile	Ser	Ser	Tyr	Gly	Asp	Arg	Trp	Glu
Val	Gln	Asn	Cys	Glu	Leu	Leu	Asn	Leu	Ala	Asp	Leu	Asn	Thr	Gly	Ala
Glu	Tyr	Val	Arg	Gly	Lys	Leu	Ala	Ala	Tyr	Met	Asn	Asp	Leu	Arg	Gly
Leu	Gly	Val	Ala	Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His	Met	Asp	Thr
Asn	Asp	Ile	Asn	Asn	Ile	Val	Gly	Arg	Leu	Pro	Asn	Ala	Pro	Tyr	Ile
Tyr	Gln	Glu	Val	Ile	Asp	Gln	Gly	Gly	Glu	Pro	Ile	Thr	Ala	Gly	Glu
Tyr	Phe	Gln	Asn	Gly	Asp	Val	Thr	Glu	Phe	Lys	Tyr	Ser	Arg	Glu	Ile
Ser	Arg	Met	Phe	Lys	Thr	Gly	Gln	Leu	Thr	His	Met	Ser	Gln	Phe	Gly
Thr	Ala	Trp	Gly	Phe	Met	Ser	Ser	Asp	Leu	Ala	Val	Val	Phe	Thr	Asp
Asn	His	Asp	Asn	Gln	Arg	Gly	His	Gly	Gly	Ala	Gly	Asp	Val	Leu	Thr
Tyr	Lys	Asp	Gly	Gln	Leu	Tyr	Thr	Leu	Gly	Asn	Ile	Phe	Glu	Leu	Ala
Trp	Pro	Tyr	Gly	Tyr	Pro	Gln	Val	Met	Ser	Ser	Tyr	Thr	Phe	Ser	Asn
Gly	Asp	Gln	Gly	Pro	Pro	Ser	Thr	Asn	Val	Tyr	Ala	Thr	Thr	Thr	Pro
Asp	Cys	Gly	Asn	Gly	Arg	Trp	Val	Cys	Glu	His	Arg	Trp	Arg	Gly	Ile
Ala	Asn	Met	Val	Ala	Phe	Arg	Asn	Tyr	Thr	Ala	Pro	Thr	Phe	Ser	Thr
Ser	Asn	Trp	Trp	Ser	Asn	Gly	Asn	Asn	Gln	Ile	Ala	Phe	Ser	Arg	Gly
Thr	Leu	Gly	Phe	Val	Ala	Ile	Asn	Arg	Glu	Gly	Gly	Ser	Leu	Asn	Arg
Thr	Phe	Gln	Thr	Gly	Leu	Pro	Val	Gly	Thr	Tyr	Cys	Asp	Val	Ile	His
Gly	Asp	Phe	Asn	Ala	Ser	Ala	Gly	Thr	Cys	Ser	Gly	Pro	Thr	Ile	Ala
Val	Asn	Gly	Ser	Gly	Gln	Ala	Thr	Ile	Thr	Val	Asn	Ala	Met	Asp	Ala
Val	Ala	Ile	Tyr	Gly	Gly	Ala	Arg	Leu	Ala	Thr	Pro	Ala	Ser	Val	Asn
Val	Thr	Phe	Asn	Glu	Asn	Ala	Thr	Thr	Thr	Trp	Gly	Gln	Asn	Val	Tyr
Ile	Val	Gly	Asn	Val	Ala	Ala	Leu	Gly	Ser	Trp	Asn	Ala	Gly	Ser	Ala
Val	Leu	Leu	Ser	Ser	Ala	Asn	Tyr	Pro	Ile	Trp	Ser	Lys	Thr	Ile	Ala
Leu	Pro	Ala	Asn	Thr	Ala	Ile	Glu	Tyr	Lys	Tyr	Ile	Lys	Lys	Asp	Gly
Ala	Gly	Asn	Val	Val	Trp	Glu	Ser	Gly	Ala	Asn	Arg	Val	Phe	Thr	Thr

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Pro Gly Ser Gly Ser Ala Thr Arg Asn Asp Thr Trp Lys
580 585

<210> 125
<211> 1395
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 125
gtggtgcaca tgaagttgaa gtaccttgcc ttagttttgt tggctgtggc ttcgataggc 60
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atgcaggcct tctactggga tgttcccggg gggggaatct ggtgggacac cataagacag 180
aaaatcccgg agtggtacga cgctggaatc tcggcgatat ggattcctcc agctagcaaa 240
gggatgggag gtggttattc catgggctac gatccctacg atttctttga cctcggcgag 300
tactatcaga agggaacagt tgagacgcgc ttcggctcaa aggaggaact ggtgaacatg 360
ataaacaccg cacactccta tggcataaag gtgatagcgg acatagtcac aaaccaccgc 420
gccggtggag accttgagtg gaaccccttt gtaaacaact atacttggac agacttctcc 480
aaggtcgcct ccggtaaata cacggccaac taccttgact tccacccaaa cgagggtcaag 540
tgctgcgatg aggggtacatt tgggtgacttt ccggacatcg cccacgagaa gagctgggat 600
cagtactggc tctgggcaag caatgagagc tacgccgcac atctccggag catagggatc 660
gatgcatggc gtttcgacta cgtcaaaggt tacggagcgt ggggttgtaa tgactggctc 720
agctgggtggg gaggctgggc cgttgagagag tactgggaca cgaacgttga tgcactcctt 780
aacggggcat acgacagcgg tgccaaggtc tttgacttcc cgctctacta caagatggac 840
gaagcccttg acaacaccaa catccccgct ttggtttacg ccctccagaa cggaggaaca 900
gtcgtttccc gcgatccctt caaggcagta actttcgttg ccaaccacga tacagatata 960
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cacgagcacc ttgccggagg aagtaccaag atcctctact acgataacga tgagctaata 1140
ttcatgaggg agggctacgg gagcaagccg ggcctcataa cctacataaa cctcggaac 1200
gactgggccc agcgtgggtg gaacgtcggc tcaaagtttg ccggctacac aatccatgaa 1260
tacacaggca atctcgttgg ctgggttgac aggtgggttc agtacgatgg atgggttaaa 1320
ctgacggcac ctctcatga tccagccaac ggatattacg gctactcagt ctggagctac 1380
gcaggcgctcg gatga 1395

<210> 126
<211> 464
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 126
Val Val His Met Lys Leu Lys Tyr Leu Ala Leu Val Leu Leu Ala Val
1 5 10 15
Ala Ser Ile Gly Leu Leu Ser Thr Pro Val Gly Ala Ala Lys Tyr Ser
20 25 30
Glu Leu Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val
35 40 45
Pro Gly Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu
50 55 60
Trp Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys
65 70 75 80
Gly Met Gly Gly Gly Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe
85 90 95
Asp Leu Gly Glu Tyr Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly
100 105 110
Ser Lys Glu Glu Leu Val Asn Met Ile Asn Thr Ala His Ser Tyr Gly
115 120 125
Ile Lys Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Gly Asp
130 135 140
Leu Glu Trp Asn Pro Phe Val Asn Asn Tyr Thr Trp Thr Asp Phe Ser
145 150 155 160
Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro
165 170 175

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Asn	Glu	Val	Lys	Cys	Cys	Asp	Glu	Gly	Thr	Phe	Gly	Asp	Phe	Pro	Asp
Ile	Ala	His	Glu	Lys	Ser	Trp	Asp	Gln	Tyr	Trp	Leu	Trp	Ala	Ser	Asn
Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly	Ile	Asp	Ala	Trp	Arg
Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val	Val	Asn	Asp	Trp	Leu
Ser	Trp	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	Trp	Asp	Thr	Asn	Val
Asp	Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Asp	Ser	Gly	Ala	Lys	Val	Phe	Asp
Phe	Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	Asp	Asn	Thr	Asn	Ile
Pro	Ala	Leu	Val	Tyr	Ala	Leu	Gln	Asn	Gly	Gly	Thr	Val	Val	Ser	Arg
Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn	His	Asp	Thr	Asp	Ile
Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile	Leu	Thr	Tyr	Glu	Gly
Gln	Pro	Val	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu	Trp	Leu	Asn	Lys	Asp
Lys	Leu	Asn	Asn	Leu	Ile	Trp	Ile	His	Glu	His	Leu	Ala	Gly	Gly	Ser
Thr	Lys	Ile	Leu	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu	Ile	Phe	Met	Arg	Glu
Gly	Tyr	Gly	Ser	Lys	Pro	Gly	Leu	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	Asn
Asp	Trp	Ala	Glu	Arg	Trp	Val	Asn	Val	Gly	Ser	Lys	Phe	Ala	Gly	Tyr
Thr	Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp	Val	Asp	Arg	Trp
Val	Gln	Tyr	Asp	Gly	Trp	Val	Lys	Leu	Thr	Ala	Pro	Pro	His	Asp	Pro
Ala	Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr	Ala	Gly	Val	Gly

<210> 127
 <211> 1848
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

<400> 127

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atgatgcaat	atttcgaatg	ggattttaccg	aatgatggca	cacttttgac	gaaagtaaaa	180
aacgaagcaa	gcagtccttc	ttcttttaggt	attactgcgt	tatgggttacc	acctgcatac	240
aaaggaacga	gcccaagggga	tgtcgggtat	ggcgtgtacg	atttgtatga	cttaggagaa	300
tttaaatcaa	aagggacgat	tcgaacgaaa	tacggaacaa	aaacgcaata	tttacaagcc	360
attcaagcgg	caaaaagcgc	tggcatgcaa	gtatacgctg	atgtcgtatt	taatcacaag	420
gcggggggcag	atagtacaga	atgggttgac	gcagtcgaag	tgaatccttc	taatcgaaac	480
caagaacat	ctggcacata	tcaaattcaa	gcatggacaa	aatttgattt	ccctggccgt	540
gggaacacat	actcaagcct	taaatggcga	tggtatcatt	ttgacggtag	ggattgggat	600
gaaagccgaa	aactaaatcg	tattttacaaa	tttcgtggca	caggaaaagc	atgggattgg	660
gaagtagaca	cagagaacgg	aaactatgac	tacttaatgt	ttgctgattt	agatatggat	720
caccctgaag	tcgtgacaga	gctaaaaaac	tggggaacat	ggtacgtcaa	tacgacaaat	780
gtcgtatgggt	ttcgcttaga	tgcagtaaaag	catattaaat	atagcttctt	cccagattgg	840
ttaacacatg	tgcgttcaca	aacacgaaaa	aatctttttg	cagtaggaga	atgttgagc	900
tacgatgtca	ataaactgca	taactacatt	acaaaaacaa	gtggaaccat	gtcgtttatt	960
gatgcgccac	ttcataacaa	cttttacact	gcttcaaaat	ctagcgggta	ttttgacatg	1020
cgctatttgt	taaataatac	gttgatgaaa	gaccagcctt	ctcttgcggt	cacactcggt	1080
gataatcatg	acacgcaacc	gggacaatct	ttacaatcat	gggtagagcc	ttggtttaag	1140
ccgcttgctt	atgcctttat	tttgacaaga	caagaaggat	atccttgctg	atgttaccgc	1200
gactattacg	gcattccctaa	atacaacatt	ccgggattga	aaagtaaaat	cgatccgctt	1260
ctcattgccc	gtagagacta	cgcatacgga	acacaacgtg	attatattga	ccatcaagac	1320

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```

attattggat ggacacggga aggaattgac tcaaaaccga actctggact tgcggccttta 1380
attactgacg gccctgggtgg aagtaaatgg atgtatgtag gtaaaaagca tgctggaaaa 1440
gtgttttacg atctcactgg aaatcgaagc gatacggtaa cgattaatgc agacggctgg 1500
ggagagttta aagtaaacgg tggctccgtt tccatttggg ttgccaaaac atcacaagtc 1560
acgtttaccg tcaacaatgc gacaacgata agcggacaaa atgtgtatgt cgttggtaac 1620
attccagagc tcggaaattg gaacacagca aacgcaatca aaatgacccc atcttcttat 1680
ccaacgtgga aagcaaccat tgctcttcca caaggaaaag ccattgaatt taaattttatt 1740
aaaaaagacc aatcgggaaa tgttggttgg gaaagcattc caaacccaac atacaccgtt 1800
ccatttttat caacaggctc atatacagct agttggaatg taccttaa 1848

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<210> 128

<211> 615

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 128

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Val Cys Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val
1      5      10      15
Ala Thr Leu Ile Ile Tyr Phe Leu Thr Pro Phe Ser Thr Ala Gln Ala
20      25      30
Asn Thr Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp
35      40      45
Leu Pro Asn Asp Gly Thr Leu Trp Thr Lys Val Lys Asn Glu Ala Ser
50      55      60
Ser Leu Ser Ser Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala Tyr
65      70      75      80
Lys Gly Thr Ser Gln Gly Asp Val Gly Tyr Gly Val Tyr Asp Leu Tyr
85      90      95
Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Ile Arg Thr Lys Tyr Gly
100      105      110
Thr Lys Thr Gln Tyr Leu Gln Ala Ile Gln Ala Ala Lys Ser Ala Gly
115      120      125
Met Gln Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala Asp
130      135      140
Ser Thr Glu Trp Val Asp Ala Val Glu Val Asn Pro Ser Asn Arg Asn
145      150      155      160
Gln Glu Thr Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe Asp
165      170      175
Phe Pro Gly Arg Gly Asn Thr Tyr Ser Phe Lys Trp Arg Trp Tyr
180      185      190
His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg Ile
195      200      205
Tyr Lys Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp Thr
210      215      220
Glu Asn Gly Asn Tyr Asp Tyr Leu Met Phe Ala Asp Leu Asp Met Asp
225      230      235      240
His Pro Glu Val Val Thr Glu Leu Lys Asn Trp Gly Thr Trp Tyr Val
245      250      255
Asn Thr Thr Asn Val Asp Gly Phe Arg Leu Asp Ala Val Lys His Ile
260      265      270
Lys Tyr Ser Phe Phe Pro Asp Trp Leu Thr His Val Arg Ser Gln Thr
275      280      285
Arg Lys Asn Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Val Asn
290      295      300
Lys Leu His Asn Tyr Ile Thr Lys Thr Ser Gly Thr Met Ser Leu Phe
305      310      315      320
Asp Ala Pro Leu His Asn Asn Phe Tyr Thr Ala Ser Lys Ser Ser Gly
325      330      335
Tyr Phe Asp Met Arg Tyr Leu Leu Asn Asn Thr Leu Met Lys Asp Gln
340      345      350
Pro Ser Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Gln Pro Gly
355      360      365
Gln Ser Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Pro Leu Ala Tyr
370      375      380
Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr Gly

```


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385	Asp	Tyr	Tyr	Gly	Ile	390	Pro	Lys	Tyr	Asn	Ile	395	Pro	Gly	Leu	Lys	Ser	400	Lys
				405	Leu	Ile	Ala	Arg	Arg	Asp	Tyr	Ala	Tyr	Gly	Thr	Gln			
			420	Leu	His	Gln	Asp	440	Ile	Ile	Gly	Trp	Thr	Arg	Glu	Gly			
		435	Asp	Pro	Asn	Ser	455	Gly	Leu	Ala	Ala	Leu	Ile	Thr	Asp	Gly			
		450	Ser	Lys	Pro	Trp	Met	Tyr	Val	Gly	Lys	Lys	His	Ala	Gly	Lys			
		465	Gly	Ser	Lys	Trp	Met	Tyr	Val	Gly	Lys	Lys	His	Ala	Gly	Lys			
		485	Leu	Thr	Gly	Asn	Arg	Ser	490	Asp	Thr	Val	Thr	Ile	Asn				
		500	Gly	Glu	Phe	Lys	Val	Asn	505	Gly	Gly	Ser	Val	Ser	Ile				
		515	Ala	Lys	Thr	Ser	Gln	Val	520	Thr	Phe	Thr	Val	Asn	Asn	Ala	Thr		
		530	Ile	Ser	Gly	Gln	Asn	Val	535	Tyr	Val	Val	Gly	Asn	Ile	Pro	Glu	Leu	
		545	Gly	Asn	Trp	Asn	Thr	Ala	550	Asn	Ala	Ile	Lys	Met	Thr	Pro	Ser	Ser	Tyr
		565	Pro	Thr	Trp	Lys	Ala	Thr	570	Ile	Ala	Leu	Pro	Gln	Gly	Lys	Ala	Ile	Glu
		580	Phe	Lys	Phe	Ile	Lys	Lys	585	Asp	Gln	Ser	Gly	Asn	Val	Val	Trp	Glu	Ser
		595	Ile	Pro	Asn	Arg	Thr	Tyr	600	Thr	Val	Pro	Phe	Leu	Ser	Thr	Gly	Ser	Tyr
		610	Thr	Ala	Ser	Trp	Asn	Val	615	Pro									

<210> 129

<211> 1854

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 129

atgctgtgcc	gccgtggcag	ggacgggtgt	tgggtgcgggc	ggcgtaatgc	gctgccgcga	60
cacccgcgtg	aacaaaataa	tatgaattat	ttgaatagga	tgggggtgtc	aagaatgaca	120
aaatctcgag	agttgcgggtg	ttcatggaaa	gtatttgttg	ttgggtgcct	gttgtggatg	180
gcttggggat	cttccgcgtc	cgccggcgta	ttgatgcaag	gcttctactg	ggacgccagt	240
accgggacca	gtgattcgtg	gtggacgcat	ttggccaagc	aagccaacgg	tctaaaacgg	300
gcgggggttca	ccgccgtatg	gattcctccg	gtgcttaaag	gggcttcagg	gggctatttc	360
aacgggtacg	atccctttga	cgactatgat	atcggaagca	aggaccagaa	aggtaccgtg	420
gcgacgcgat	gggggacgcg	agaagaactg	caacgtgccg	tggccgtgat	gcgcgcgaac	480
ggtctggatg	tgtatgtgga	tctggtgctg	aaccaccgca	acggggacga	cgggaattgg	540
aattttcatt	acaaagatgc	gtacggcaaa	gtgggttacg	ggcggtttca	aaaggggttt	600
tacgattttc	accccaacta	caacattcag	gatgccaatg	ttcccaacga	ggattccagc	660
ttcgggcgcg	atttagccca	tgacaatccg	tatgtggccg	atggactgaa	ggctgcaggc	720
gattggctga	ccaaagccct	cgatgttcag	ggatatcgtc	tggattacgt	gaaaggcatc	780
agctacacct	tcctgaaaag	ttatctgtcc	tatggggcca	tgaacggaaa	atttgccgtc	840
ggtgagtact	gggatgccaa	ccgggatacg	ttgaactggt	gggcgaacac	ggcgaatgaa	900
gggcgggtccc	atgtgtttga	ttttgcgttg	cgcgaggagc	tgaaaaacat	gtgcaatgcg	960
gacgggtact	acgacatgcg	tcgattggac	cacgcgggtc	tggtcggaat	cgaccctggg	1020
aaggcgggtga	cgtttgtcga	aaaccatgat	acggatcggc	acgaccccat	ctacaataac	1080
aagcatttgg	cgatgccta	catcttgacg	tcggaagggt	atccgacggt	gttctggaag	1140
gattactacc	aatacggaa	gaagccgatc	atcgacaacc	tcattttgat	ccacgaacac	1200
attgcgtacg	gaacgaccca	agagcgttgg	aaagacgaag	atgtctttgt	gtatgagcgg	1260
accggaggga	agcgcctatt	ggtggggcct	aacgacaatc	gcgccaccag	caaaacggtc	1320
accgtacaga	ccggcttttg	tgccaacgtg	gccttgcacg	actacaccgg	caacggcccc	1380
gatctccgta	ccgacgccta	cggtcgggta	accttgacca	ttcctgcaaa	cgggtacgtg	1440
gcctattccg	ttccgggcat	ctccggatcc	tttgtgccgg	tcgagaaaac	cgtgacgcag	1500
gagtttgccg	gggcgtccga	cttgatatt	cgtccggccg	ataacacgca	atttgtgcag	1560
gtcgggcgga	tatacgccaa	ggcaaacaag	ccggttacag	cgggaattga	ttgggatgcc	1620
aaagactgga	cgacctcac	gtcgattctc	ctagaagtgc	gttcggcttc	gggaacgctc	1680
atcacgacaa	agaccgtgac	ccaattgtcg	tcccagggtg	cccgcgtttc	cttcacgcct	1740

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tcggctaccg gatggtacgt cttttccatt cgaagctata acacgccttc gacgaaccca 1800
aagccggcct actggttaaa ggtaacgtat acggcgccgc aattgcttca gtaa 1854

<210> 130
<211> 617
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 130
Met Arg Cys Arg Arg Gly Arg Asp Gly Cys Trp Cys Gly Arg Arg Asn
1 5 10 15
Ala Leu Pro Arg His Pro Arg Glu Gln Asn Asn Met Asn Tyr Leu Asn
20 25 30
Arg Met Gly Val Ser Arg Met Thr Lys Ser Arg Glu Leu Arg Cys Ser
35 40 45
Trp Lys Val Phe Val Val Gly Cys Leu Leu Trp Met Ala Trp Gly Ser
50 55 60
Ser Ala Ser Ala Gly Val Leu Met Gln Gly Phe Tyr Trp Asp Ala Ser
65 70 75 80
Thr Gly Thr Ser Asp Ser Trp Trp Thr His Leu Ala Lys Gln Ala Asn
85 90 95
Gly Leu Lys Arg Ala Gly Phe Thr Ala Val Trp Ile Pro Pro Val Leu
100 105 110
Lys Gly Ala Ser Gly Gly Tyr Ser Asn Gly Tyr Asp Pro Phe Asp Asp
115 120 125
Tyr Asp Ile Gly Ser Lys Asp Gln Lys Gly Thr Val Ala Thr Arg Trp
130 135 140
Gly Thr Arg Glu Glu Leu Gln Arg Ala Val Ala Val Met Arg Ala Asn
145 150 155 160
Gly Leu Asp Val Tyr Val Asp Leu Val Leu Asn His Arg Asn Gly Asp
165 170 175
Asp Gly Asn Trp Asn Phe His Tyr Lys Asp Ala Tyr Gly Lys Val Gly
180 185 190
Tyr Gly Arg Phe Gln Lys Gly Phe Tyr Asp Phe His Pro Asn Tyr Asn
195 200 205
Ile Gln Asp Ala Asn Val Pro Asn Glu Asp Ser Ser Phe Gly Arg Asp
210 215 220
Leu Ala His Asp Asn Pro Tyr Val Ala Asp Gly Leu Lys Ala Ala Gly
225 230 235 240
Asp Trp Leu Thr Lys Ala Leu Asp Val Gln Gly Tyr Arg Leu Asp Tyr
245 250 255
Val Lys Gly Ile Ser Tyr Thr Phe Leu Lys Ser Tyr Leu Ser Tyr Gly
260 265 270
Ala Met Asn Gly Lys Phe Ala Val Gly Glu Tyr Trp Asp Ala Asn Arg
275 280 285
Asp Thr Leu Asn Trp Trp Ala Asn Thr Ala Met Glu Gly Arg Ala His
290 295 300
Val Phe Asp Phe Ala Leu Arg Glu Glu Leu Lys Asn Met Cys Asn Ala
305 310 315 320
Asp Gly Tyr Tyr Asp Met Arg Arg Leu Asp His Ala Gly Leu Val Gly
325 330 335
Ile Asp Pro Trp Lys Ala Val Thr Phe Val Glu Asn His Asp Thr Asp
340 345 350
Arg His Asp Pro Ile Tyr Asn Asn Lys His Leu Ala Tyr Ala Tyr Ile
355 360 365
Leu Thr Ser Glu Gly Tyr Pro Thr Val Phe Trp Lys Asp Tyr Tyr Gln
370 375 380
Tyr Gly Met Lys Pro Ile Asp Asn Leu Ile Trp Ile His Glu His
385 390 395 400
Ile Ala Tyr Gly Thr Thr Gln Glu Arg Trp Lys Asp Glu Asp Val Phe
405 410 415
Val Tyr Glu Arg Thr Gly Gly Lys Arg Leu Leu Val Gly Leu Asn Asp
420 425 430
Asn Arg Ala Thr Ser Lys Thr Val Thr Val Gln Thr Gly Phe Gly Ala
435 440 445

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```

Asn Val Ala Leu His Asp Tyr Thr Gly Asn Gly Pro Asp Leu Arg Thr
 450 455
Asp Ala Tyr Gly Arg Val Thr Leu Thr Ile Pro Ala Asn Gly Tyr Val
 465 470 475 480
Ala Tyr Ser Val Pro Gly Ile Ser Gly Ser Phe Val Pro Val Glu Lys
 485 490 495
Thr Val Thr Gln Glu Phe Ala Gly Ala Ser Asp Leu Asp Ile Arg Pro
 500 505 510
Ala Asp Asn Thr Gln Phe Val Gln Val Gly Arg Ile Tyr Ala Lys Ala
 515 520 525
Asn Lys Pro Val Thr Ala Glu Leu Tyr Trp Asp Ala Lys Asp Trp Thr
 530 535 540
Thr Ser Thr Ser Ile Leu Leu Glu Val Arg Ser Ala Ser Gly Thr Leu
 545 550 555 560
Ile Thr Thr Lys Thr Val Thr Gln Leu Ser Ser Gln Gly Thr Arg Val
 565 570 575
Ser Phe Thr Pro Ser Ala Thr Gly Trp Tyr Val Phe Ser Ile Arg Ser
 580 585 590
Tyr Asn Thr Pro Ser Thr Asn Pro Lys Pro Ala Tyr Trp Leu Lys Val
 595 600 605
Thr Tyr Thr Ala Pro Gln Leu Leu Gln
 610 615

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<210> 131
 <211> 1881
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

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<400> 131
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gccttgacgc tggccaccac ggccctgggc atctcgacgg cccaggccca gaggtcaccg 120
cgacggcct tcgtgcatct gttcgaatgg aagtggaccg acatcgcgcg cgagtgcgag 180
accttcctcg ggcccaaggg cttcgcgcg gtgcaggtgt cgcgccgaa cgagcacaac 240
tgggtgacca gcggtgatgg tgcaccttat ccgtggtgga tgcgtacca gccggtgagc 300
tacagcctgg accgcagccg cagcggcacg cgcgccgagt tccaggacat ggtcaaccga 360
tgcaatgccg tgggctgagg catctacgtg gacgccgtga tcaatcacat gtccggcggc 420
acggggcgga cctcgagcgc tgggcgcagc tggagctatc acaactacc tgggctctat 480
ggccccaacg acttcacca gccggtgtgc agcatcacca actacgggga tgcgaacaat 540
gtgcagcgtt gcgagctctc gggcttgacg gacctggaca ctgggagcgc ttatgtgcgc 600
ggcaagatcg ccgactatct ggtgatctg gtcaaatagg gggtaagggt cttccgggtg 660
gatcgggcca cccgaccag cccgaccgac ctgggcgcca tcacgatgc ggtcaacagc 720
cgacccggcg cgaaccgccc tttctggttt ctggaggtga ttggcgcggc cggcgaggca 780
gtgcagccga accagtaact ctcgctcggc ggcggccagg tcaccgtgac cgagttcaac 840
tatgggaagc aaatcttcgg caagttcgcc ggtggcggcc gtctggccga gctgcgcagc 900
ttcgtgaaa cctggggcct gatgccagc agcaaagcga ttgctttcat cgacaaccac 960
gacaagcagc gcggtcatgg cggcggtggc aactatctga cctaccacca tggctcgacc 1020
tacgatctgg ccaacatctt catgctggct tggccttatg gctaccgggc gctgatgtcc 1080
agctatgcct tcaaccgcag cacggcctac gacacgagct ttggcccgcc acacgacagt 1140
ggtggcgcca cccgtggccc ctgggatggt ggcggcagcc agccggcctg cttcaaccag 1200
agcatcggtg gctgggtgtg tgagcaccgc tggcggggca tcgccaatat ggtggccttc 1260
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ttcaacaatg gtcagtgcac gggccatgtg gtgacggtcg atgccggcg ctacgtgacg 1500
ctgacggccg ggcccaatgg tgcggcggcc atccacgtgg gcgcccgtct ggacggcgcc 1560
tctagccgc cgacgaccgc ctcggtgacg ttcaacgcgt cggccgatac cttttgggga 1620
cagaacctgt tcgtcgtggg caaccacagc gcactgggca actggtcgcc ggcggccgcc 1680
aggccgatga cttggatttc cgttcgggc acgcgcggga actggcgcg ggtgctcaat 1740
ttgccggcca ataccaccta ccaatacaag ttcacgaaga aggacggggc tggaaacgtg 1800
gtttgggagg gcggtggcaa tcgctcgtg accacgccgt ctggggcgcg atcggtgagc 1860
acgggcggca attggcagta g

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<210> 132
 <211> 626
 <212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 132

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Met 1 Pro Gln Leu Tyr 5 Pro Leu Pro Pro Arg 10 Trp Arg Arg Ala 15 Ala Arg
Gln 1 Gly Leu Ala 20 Leu Thr Leu Ala 25 Thr Thr Ala Leu Gly 30 Ile Ser
Thr Ala Gln 35 Ala Gln Ser Ala 40 Pro Arg Thr Ala Phe Val 45 His Leu Phe
Glu Trp 50 Lys Trp Thr Asp 55 Ala Arg Glu Cys 60 Thr Phe Leu Gly
Pro 65 Lys Gly Phe Ala 70 Val Gln Val Ser 75 Pro Asn Glu His Asn 80
Trp Val Thr Ser Gly 85 Asp Gly Ala Pro Tyr 90 Pro Trp Trp Met Arg 95 Tyr
Gln Pro Val Ser 100 Tyr Ser Leu Asp Arg 105 Ser Arg Ser Gly Thr Arg Ala
Glu Phe Gln 115 Asp Met Val Asn Arg 120 Cys Asn Ala Val Gly Val Gly Ile
Tyr Val Asp Ala Val Ile Asn 135 His Met Ser Gly Gly Thr Gly Gly Thr
Ser 145 Ser Ala Gly Arg Ser 150 Trp Ser Tyr His Asn 155 Tyr Pro Gly Leu Tyr 160
Gly Pro Asn Asp Phe 165 His Gln Pro Val Cys 170 Ser Ile Thr Asn Tyr Gly 175
Asp Ala Asn Asn Val Gln Arg Cys Glu 185 Leu Ser Gly Leu Gln Asp Leu 190
Asp Thr Gly 195 Ser Ala Tyr Val Arg 200 Gly Lys Ile Ala Asp Tyr Leu Val
Asp Leu Val Asn Met Gly Val 215 Lys Gly Phe Arg Val Asp Ala Ala Lys
His 225 Ile Ser Pro Thr Asp 230 Leu Gly Ala Ile Ile Asp Ala Val Asn Ser 240
Arg Thr Gly Ala Asn 245 Arg Pro Phe Trp Phe 250 Leu Glu Val Ile Gly Ala 255
Ala Gly Glu Ala Val Gln Pro Asn Gln Tyr Phe Ser Leu Gly Gly Gly 270
Gln Val Thr Val Thr Glu Phe Asn 280 Tyr Gly Lys Gln Ile Phe Gly Lys
Phe Ala Gly Gly Gly Arg Leu 295 Ala Glu Leu Arg Ser 300 Phe Gly Glu Thr
Trp 305 Gly Leu Met Pro Ser 310 Ser Lys Ala Ile Ala Phe Ile Asp Asn His 320
Asp Lys Gln Arg Gly 325 His Gly Gly Gly Gly Asn Tyr Leu Thr Tyr His 335
His Gly Ser Thr 340 Tyr Asp Leu Ala Asn 345 Ile Phe Met Leu Ala Trp Pro 350
Tyr Gly Tyr 355 Pro Ala Leu Met Ser 360 Ser Tyr Ala Phe Asn Arg Ser Thr 365
Ala Tyr Asp Thr Ser Phe Gly 375 Pro Pro His Asp Ser Gly Gly Ala Thr 380
Arg Gly Pro Trp Asp Gly 390 Gly Gly Ser Gln Pro Ala Cys Phe Asn Gln 400
Ser Ile Gly Gly Trp Val Cys Glu His Arg 410 Trp Arg Gly Ile Ala Asn 415
Met Val Ala Phe 420 Arg Asn Ala Thr Leu 425 Pro Asn Trp Thr Val Thr Asp 430
Trp Trp Asp 435 Asn Gly Asn Asn Gln 440 Ile Ala Phe Gly Arg Gly Asp Lys 445
Gly Phe Val Val Ile Asn Arg 455 Glu Asp Ala Ala Leu Thr Arg Asn Phe 460
Lys 465 Thr Ser Leu Pro Ala Gly Gln Tyr Cys Asp Val Ile Ser Gly Asp 480
Phe Asn Asn Gly Gln 485 Cys Thr Gly His Val 490 Val Thr Val Asp Ala Gly 495

```

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Gly	Tyr	Val	Thr	Leu	Thr	Ala	Gly	Pro	Asn	Gly	Ala	Ala	Ala	Ile	His
			500					505					510		
Val	Gly	Ala	Arg	Leu	Asp	Gly	Ala	Ser	Gln	Pro	Pro	Thr	Thr	Ala	Ser
		515					520					525			
Val	Thr	Phe	Asn	Ala	Ser	Ala	Asp	Thr	Phe	Trp	Gly	Gln	Asn	Leu	Phe
		530				535					540				
Val	Val	Gly	Asn	His	Ser	Ala	Leu	Gly	Asn	Trp	Ser	Pro	Ala	Ala	Ala
545					550					555					560
Arg	Pro	Met	Thr	Trp	Ile	Ser	Gly	Ser	Gly	Thr	Arg	Gly	Asn	Trp	Arg
				565					570					575	
Ala	Val	Leu	Asn	Leu	Pro	Ala	Asn	Thr	Thr	Tyr	Gln	Tyr	Lys	Phe	Ile
			580					585					590		
Lys	Lys	Asp	Gly	Ala	Gly	Asn	Val	Val	Trp	Glu	Gly	Gly	Gly	Asn	Arg
		595					600					605			
Val	Val	Thr	Thr	Pro	Ser	Gly	Gly	Gly	Ser	Val	Ser	Thr	Gly	Gly	Asn
	610					615					620				
Trp	Gln														
625															

<210> 133
 <211> 1638
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 133

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caatatttcg	aatgggattt	accgaatgat	gggacgcttt	ggacgaaagt	aaaaaatgaa	180
gctaccaatc	tttcttcgct	aggtattaca	gcgttatggc	tccctccagc	atataaagga	240
acgagccaaa	gcgatgtcgg	atatggcgtg	tacgatttat	atgaccttgg	ggaattttaat	300
caaaaaggga	cgatccgaac	gaaatacggg	acaaaagcac	aatatattca	agccatccaa	360
gctgccaaag	ccgcagggat	gcaagtatat	gcagatgttg	tatttaatac	taaggcgggg	420
gctgacggca	cagaatttgt	cgatgcagtt	gaggtaaacc	cttctaatac	aaatcaagaa	480
acatctggca	catatcaaat	tcaagcatgg	acaaaatttg	attttctctg	tcgtggaaac	540
acatactcca	gcttcaaatg	gcgctgggat	cattttgacg	gtaccgattg	ggatgaaagt	600
cgtaaattaa	atcgtattta	caaattccgc	ggtacaggaa	aagcgtggga	ctgggaagtc	660
gatacagaaa	acggaaacta	tgattattta	atgttcgctg	atttagatat	ggatcacctt	720
gaagtgttga	cagagttaaa	aaactgggga	aaatgggatg	taaatacgac	aaatgtagac	780
ggatttcggt	tggaatgccg	aaaacatatt	aaatacagct	ttttccctga	ctggctaaca	840
tatgtacgta	atcaaacagg	aaaaaattta	tttgctgttg	gggaattttg	gagctatgac	900
gtcaataaag	tgcataacta	cattacaata	acaaatggat	cgatgtcgtt	atttgatgca	960
ccittgcata	acaactttta	tatcgcttcc	aaatcgagtg	gatattttga	catgcgttat	1020
ttattgaata	atacattaat	gaaagatcaa	ccttcactcg	ctgtaacact	tgatcgataac	1080
catgatacac	aaccagggtca	atctttacaa	tcatgggtag	aagcttggtt	taaaccgctt	1140
gcttacgcct	ttattttaac	aagacaagag	gggtatcctt	gcgtatttta	cgggtgactat	1200
tacggaatcc	cgaaatacaa	tattccggga	ttaaaaagta	aaattgatcc	gcttttaatt	1260
gctcgtcgtg	attatgcctt	tggaacacaa	cgatgattaca	ttgatcatca	agacattatc	1320
ggatggacac	gagaaggcat	tgatgcaaaa	ccgaactctg	gacttgccgg	tttaattacc	1380
gacggccctg	gcggaagtaa	atggatgtat	gtcggtaaaa	aacatgctgg	gaaagtgttt	1440
tatgatttaa	ctggaaatcg	aagtgcacac	gtaacgatta	atgcggacgg	ttggggagaa	1500
tttaaagtaa	acggcggctc	cgtttcgatt	tgggtggcta	aaacatcaaa	cgtcacattt	1560
acagtcaata	acgccacaac	aacaagtgga	caaacgatat	atgttgtttg	caacattcca	1620
gagctaggca	attctttt					1638

<210> 134
 <211> 546
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 134

Met	Asn	Asn	Val	Lys	Lys	Val	Trp	Leu	Tyr	Tyr	Ser	Ile	Ile	Ala	Thr
1				5					10					15	

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Leu	Val	Ile	Ser	Phe	Phe	Thr	Pro	Phe	Ser	Thr	Ala	Gln	Ala	Asn	Thr
Ala	Pro	Val	Asn	Gly	Thr	Met	Met	25	Gln	Tyr	Phe	Glu	Trp	Asp	Leu
Asn	Asp	Gly	Thr	Leu	Trp	Thr	Lys	Val	Lys	Asn	Glu	Ala	Thr	Asn	Leu
Ser	Ser	Leu	Gly	Ile	Thr	Ala	Leu	Trp	Leu	Pro	Ala	Tyr	Lys	Gly	80
Thr	Ser	Gln	Ser	Asp	Val	Gly	Tyr	Gly	Val	Tyr	Asp	Leu	Tyr	Asp	Leu
Gly	Glu	Phe	Asn	Gln	Lys	Gly	Thr	Ile	Arg	Thr	Lys	Tyr	Gly	Thr	Lys
Ala	Gln	Tyr	Ile	Gln	Ala	Ile	Gln	Ala	Ala	Lys	Ala	Ala	Gly	Met	Gln
Val	Tyr	Ala	Asp	Val	Val	Phe	Asn	His	Lys	Ala	Gly	Ala	Asp	Gly	Thr
Glu	Phe	Val	Asp	Ala	Val	Glu	Val	Asn	Pro	Ser	Asn	Arg	Asn	Gln	Glu
Thr	Ser	Gly	Thr	Tyr	Gln	Ile	Gln	Ala	Trp	Thr	Lys	Phe	Asp	Phe	Pro
Gly	Arg	Gly	Asn	Thr	Tyr	Ser	Ser	Phe	Lys	Trp	Arg	Trp	Tyr	His	Phe
Asp	Gly	Thr	Asp	Trp	Asp	Glu	Ser	Arg	Lys	Leu	Asn	Arg	Ile	Tyr	Lys
Phe	Arg	Gly	Thr	Gly	Lys	Ala	Trp	Asp	Trp	Glu	Val	Asp	Thr	Glu	Asn
Gly	Asn	Tyr	Asp	Tyr	Leu	Met	Phe	Ala	Asp	Leu	Asp	Met	Asp	His	Pro
Glu	Val	Val	Thr	Glu	Leu	Lys	Asn	Trp	Gly	Lys	Trp	Tyr	Val	Asn	Thr
Thr	Asn	Val	Asp	Gly	Phe	Arg	Leu	Asp	Ala	Val	Lys	His	Ile	Lys	Tyr
Ser	Phe	Phe	Pro	Asp	Trp	Leu	Thr	Tyr	Val	Arg	Asn	Gln	Thr	Gly	Lys
Asn	Leu	Phe	Ala	Val	Gly	Glu	Phe	Trp	Ser	Tyr	Asp	Val	Asn	Lys	Leu
His	Asn	Tyr	Ile	Thr	Lys	Thr	Asn	Gly	Ser	Met	Ser	Leu	Phe	Asp	Ala
Pro	Leu	His	Asn	Asn	Phe	Tyr	Ile	Ala	Ser	Lys	Ser	Ser	Gly	Tyr	Phe
Asp	Met	Arg	Tyr	Leu	Leu	Asn	Asn	Thr	Leu	Met	Lys	Asp	Gln	Pro	Ser
Leu	Ala	Val	Thr	Leu	Val	Asp	Asn	His	Asp	Thr	Gln	Pro	Gly	Gln	Ser
Leu	Gln	Ser	Trp	Val	Glu	Ala	Trp	Phe	Lys	Pro	Leu	Ala	Tyr	Ala	Phe
Ile	Leu	Thr	Arg	Gln	Glu	Gly	Tyr	Pro	Cys	Val	Phe	Tyr	Gly	Asp	Tyr
Tyr	Gly	Ile	Pro	Lys	Tyr	Asn	Ile	Pro	Gly	Leu	Lys	Ser	Lys	Ile	Asp
Pro	Leu	Leu	Ile	Ala	Arg	Arg	Asp	Tyr	Ala	Tyr	Gly	Thr	Gln	Arg	Asp
Tyr	Ile	Asp	His	Gln	Asp	Ile	Ile	Gly	Trp	Thr	Arg	Glu	Gly	Ile	Asp
Ala	Lys	Pro	Asn	Ser	Gly	Leu	Ala	Ala	Leu	Ile	Thr	Asp	Gly	Pro	Gly
Gly	Ser	Lys	Trp	Met	Tyr	Val	Gly	Lys	Lys	His	Ala	Gly	Lys	Val	Phe
Tyr	Asp	Leu	Thr	Gly	Asn	Arg	Ser	Asp	Thr	Val	Thr	Ile	Asn	Ala	Asp
Gly	Trp	Gly	Glu	Phe	Lys	Val	Asn	Gly	Ser	Val	Ser	Ile	Trp	Val	
Ala	Lys	Thr	Ser	Asn	Val	Thr	Phe	Thr	Val	Asn	Asn	Ala	Thr	Thr	Thr
Ser	Gly	Gln	Asn	Val	Tyr	Val	Val	Gly	Asn	Ile	Pro	Glu	Leu	Gly	Asn
Ser	Leu														

<210> 135
 <211> 1935
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 135
 gtgacaggca ccccgctctt atacattcct ccacataaaa taaccataca gctttcaaat 60
 ttgttgaaat gtataaaaat aaaaaatagt attgtaagcg ttaacatccg tcattataat 120
 aacttcaaac gcgtttatgt tttaatgcaa acgtttgcat cctcatttta tttaaagaaa 180
 ggatgtgtgt gcatgaatta tttgaaaaaa gtgtgggtgt attacgctat cgtcgtctacc 240
 ttaatcattt cctttcttac gcccttttca actgcacaag ccaacactgc accagtcaac 300
 ggaacgatga tgcaatattt cgaatgggat ttaccgaatg atggcacact ttggacgaaa 360
 gtaaaaaacg aagcaagcag cctttcttct ttaggtatta ctgcttatg gttaccacct 420
 gcatacaaaag gaacgagcca aggggatgtc ggggatggcg tgtacgattt gtatgactta 480
 ggagaattta atcaaaaagg gacgattcga acgaaatagc gaacaaaaac gcaatattta 540
 caagccattc aagcggcaaa aagcgtggc atgcaagtat acgctgatgt cgtatttaat 600
 cacaaggcgg gggcagatag tacagaatgg gttgacgcag tcgaagtga tcttcttaat 660
 cgaaaccaag aaacatctgg cacatatcaa attcaagcat ggacaaaatt tgatttcctt 720
 gaccgtggga acacatactc aagctttaaa tggcgtggg atcattttga cggtagcgat 780
 tgggatgaaa gtcgaaaact aaatcgcat tacaattttc gtggcacagg aaaagcatgg 840
 gattgggaag tagacacaga gaacggaac tatgactact taatgtttgc tgatttagat 900
 atggatcacc ctgaagtcgt gacagagcta aaaaactggg gaacatggta cgtcaatacg 960
 acaaatgtcg atgggtttcg cttagatgca gtaaaagcata ttaaatatag ctttttccca 1020
 gattgggttaa catatgtgcg ctcaaaaaca caaaaaaatc tgtttgcagt aggagaattt 1080
 tggagctacg atgtcaataa actgcataac tacattacaa aaacaagtgg aaccatgtcg 1140
 ttatttgatg cgccacttca taacaacttt tacttgctt caaaatctag cgggtatttt 1200
 gacatgcgct atttgttaaa taatacgttg atgaaagacc agccttctct tgcggtcaca 1260
 ctggttgata atcatgacac gcaaccggga caatctttac aatcatgggt agagccttgg 1320
 ttaagccgc ttgcttatgc ctttattttg acaagacaag aaggatatcc ttgcgtattt 1380
 tacggcgact attacggcat ccctaaatac aatattccgg gattgaaaag taaaatcgat 1440
 ccgcttctca ttgcccgtag agactacgca tacggaacac aacgtgatta tattgaccat 1500
 caagacatta ttggatggac acgggaagga attgactcaa aaccgaactc tggacttgcg 1560
 gctttaatta ctgacggctc tgggtggaagt aaatggatgt atgtaggtaa aaagcatgct 1620
 ggaaaagtgt tttacgatct cactggaaat cgaagcgata cggtaacgat taatgcagac 1680
 ggctggggag agtttaaagt aaacgggtggc tccgtttcca tttgggttgc caaaacatca 1740
 caagtcacgt ttaccgtcaa caatgcgaca acgacaagcg gacaaaatgt gtatgtcggt 1800
 ggcaacattc cagagctcgg aaattggaac acagcaaacg caatcaaaat gaccccatct 1860
 tcttatccaa cgtggaaaac aaccattgct cttccacaag gaaaagcaat tggcggcgta 1920
 cgccatggcc ctga 1935

<210> 136
 <211> 644
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 136
 Val Thr Gly Thr Pro Ser Leu Tyr Ile Pro Pro His Lys Ile Thr Ile
 1 5 10 15
 Gln Leu Ser Asn Leu Leu Lys Cys Ile Lys Ile Lys Asn Ser Ile Val
 20 25 30
 Ser Val Asn Ile Arg His Tyr Asn Asn Phe Lys Arg Val Tyr Val Leu
 35 40 45
 Met Gln Thr Phe Ala Ser Ser Phe Tyr Leu Lys Lys Gly Cys Val Cys
 50 55 60
 Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val Ala Thr
 65 70 75 80
 Leu Ile Ile Ser Phe Leu Thr Pro Phe Ser Thr Ala Gln Ala Asn Thr
 85 90 95
 Ala Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp Asp Leu Pro
 100 105 110
 Asn Asp Gly Thr Leu Trp Thr Lys Val Lys Asn Glu Ala Ser Ser Leu

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Ser	Ser	115	Gly	Ile	Thr	Ala	120	Leu	Trp	Leu	Pro	Pro	125	Ala	Tyr	Lys	Gly
Thr	Ser	130	Gln	Gly	Asp	Val	135	Gly	Tyr	Gly	Val	Tyr	140	Asp	Leu	Tyr	Asp
145	Gly	Glu	Phe	Asn	Gln	150	Lys	Gly	Thr	Ile	Arg	155	Thr	Lys	Tyr	Gly	Thr
Thr	Gln	Tyr	Leu	165	Gln	Ala	Ile	Gln	Ala	170	Ala	Lys	Ser	Ala	Gly	175	Met
Val	Tyr	Ala	180	Asp	Val	Val	Phe	Asn	185	His	Lys	Ala	Gly	Ala	190	Asp	Ser
Glu	Trp	195	Val	Asp	Ala	Val	Glu	200	Val	Asn	Pro	Ser	Asn	205	Arg	Asn	Gln
Thr	210	Ser	Gly	Thr	Tyr	Gln	215	Ile	Gln	Ala	Trp	Thr	220	Lys	Phe	Asp	Phe
225	Asp	Arg	Gly	Asn	Thr	Tyr	230	Ser	Ser	Phe	Lys	Trp	235	Arg	Trp	Tyr	His
Asp	Gly	Thr	245	Asp	Trp	Asp	Glu	Ser	Arg	250	Lys	Leu	Asn	Arg	Ile	255	Tyr
Phe	Arg	Gly	260	Thr	Gly	Lys	Ala	Trp	265	Asp	Trp	Glu	Val	270	Asp	Thr	Glu
Gly	Asn	Tyr	275	Asp	Tyr	Leu	Met	280	Phe	Ala	Asp	Leu	Asp	285	Met	Asp	His
Glu	305	Val	Val	Thr	Glu	Leu	295	Lys	Asn	Trp	Gly	Thr	300	Trp	Tyr	Val	Asn
Thr	Asn	Val	Asp	Gly	310	Phe	Arg	Leu	Asp	Ala	315	Val	Lys	His	Ile	Lys	320
Ser	Phe	Phe	Pro	Asp	Trp	Leu	Thr	Tyr	325	Val	Arg	Ser	Gln	330	Thr	Gln	Lys
Asn	Leu	Phe	340	Ala	Val	Gly	Glu	Phe	345	Trp	Ser	Tyr	Asp	350	Val	Asn	Lys
His	355	Asn	Tyr	Ile	Thr	Lys	Thr	360	Ser	Gly	Thr	Met	Ser	365	Leu	Phe	Asp
Pro	370	Leu	His	Asn	Asn	Phe	Tyr	375	Thr	Ala	Ser	Lys	Ser	380	Ser	Gly	Tyr
385	Asp	Met	Arg	Tyr	Leu	Leu	Asn	390	Asn	Thr	Leu	Met	Lys	395	Asp	Gln	Pro
Leu	Ala	Val	Thr	405	Leu	Val	Asp	Asn	His	410	Asp	Thr	Gln	415	Pro	Gly	Gln
Leu	Gln	Ser	Trp	Val	Glu	Pro	Trp	425	Phe	Lys	Pro	Leu	Ala	430	Tyr	Ala	Phe
Ile	Leu	Thr	Arg	Gln	Glu	Gly	440	Tyr	Pro	Cys	Val	Phe	445	Tyr	Gly	Asp	Tyr
Tyr	450	Gly	Ile	Pro	Lys	Tyr	455	Asn	Ile	Pro	Gly	Leu	460	Lys	Ser	Lys	Ile
465	Pro	Leu	Leu	Ile	Ala	Arg	Arg	470	Asp	Tyr	Ala	Tyr	475	Gly	Thr	Gln	Arg
Tyr	Ile	Asp	His	Gln	Asp	Ile	Ile	485	Gly	Trp	Thr	Arg	490	Glu	Gly	Ile	Asp
Ser	Lys	Pro	500	Asn	Ser	Gly	Leu	505	Ala	Ala	Leu	Ile	510	Asp	Gly	Pro	Gly
Gly	Ser	Lys	Trp	Met	Tyr	Val	515	Gly	Lys	Lys	His	Ala	520	Gly	Lys	Val	Phe
Tyr	530	Asp	Leu	Thr	Gly	Asn	Arg	535	Ser	Asp	Thr	Val	540	Thr	Ile	Asn	Ala
545	Gly	Trp	Gly	Glu	Phe	Lys	Val	550	Asn	Gly	Gly	Ser	555	Val	Ser	Ile	Trp
Ala	Lys	Thr	Ser	Gln	Val	Thr	Phe	565	Thr	Val	Asn	Asn	570	Ala	Thr	Thr	Thr
Ser	Gly	Gln	Asn	Val	Tyr	Val	Val	580	Gly	Asn	Ile	Pro	585	Glu	Leu	Gly	Asn
Trp	595	Asn	Thr	Ala	Asn	Ala	Ile	600	Lys	Met	Thr	Pro	605	Ser	Ser	Tyr	Pro
Trp	610	Lys	Thr	Thr	Ile	Ala	615	Leu	Pro	Gln	Gly	Lys	620	Ala	Ile	Gly	Gly
625	Arg	His	Gly	Pro		630							635				640

<210> 137
 <211> 1320
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 137
 gtggggacggg caggcttggc gcatacactcg aacacttccg ccaaggggac atacgggtca 60
 cctctcgaac tgcgtccgga tcgccccgcc gtggccgggg cggtcgagct tgaagatgtc 120
 cagcggggag ccgccgccga ggatcacccc ggccggcgta tcgccaggg cggggctcag 180
 ctgaaagccg tggccggagc cgcctcccag gagccagacg ttggaggccc gcggatggcg 240
 gtcgaggagg aggtggccgt cggggctggt ctcgtactgg cagacgcggg tctcgaccag 300
 cggcgcgctc ttcagggccg ggaaccggcg ggccacctcg gcccgggccg cttccagcag 360
 ggccgggggtg atcgtccgct cgcgcccgct gggatcgatg ggctcgcccc ggggtgtcgtc 420
 cgccaccttg aagccgcggt gctcgttgcc ggggatgccg tagtagatcc gctcgccgag 480
 atcgaccag accggacagc cgccctcctg gaagcgcggg tcgccggcg gcgtgccgaa 540
 gaagaacacc tcctggcggg tgttgccgag gaaccgctca ccgatcacgt ccgggaacag 600
 cccggccagc cagggaccgc aggcgaagac gtagaggtcg gccgcgagag tggagccgtc 660
 cgaaaaggtag agccgctcca agggcccccg gaccatggcg gcctgccggt actccccgcc 720
 ctgcgccctg aacagctcca ccacggtccg gcaggcgcg cgggcgaaca gggcgccggc 780
 ttctctctcg taccagatcg tgcggacgcc gtcgaaatcg acctggggga agcggtcccg 840
 ggctcccccc tgagacagct cggcgaccgg cagcccccg tcctccagaa aagggaaggga 900
 gtcgcggacg tagctgtcgt cctcgccgca catccagagg accccggtcc tttgttacag 960
 ccggtaaccc gactggactt cggcgctccc caagagctcg aaggagcggg cgaccactc 1020
 cagctacaga cggtcgggtc cgtaggcgcc gcggatgata cgctctcgc caccggagct 1080
 ggagcgggag tgccccggac cccaggcgct caggagggtc acccgggctc cgcggcggag 1140
 gagatgcagg gcggtccagc cgccgaaggc gccggcgccg acgacggcga tatggggatg 1200
 ggagggcatg gcgggcgtaa ggttatcgca gcccgatcct tcgctggcat cccatctccg 1260
 accggagtat cctggaataa tcgaagaagg agatcgacat gcaatcgaac ggaaacgtga 1320

<210> 138
 <211> 439
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 138
 Val Gly Arg Ala Gly Leu Ala His His Ser Asn Thr Ser Ala Lys Gly
 1 5 10 15
 Thr Tyr Gly Ser Pro Leu Glu Leu Arg Pro Asp Arg Pro Ala Val Ala
 20 25 30
 Gly Ala Val Glu Leu Glu Asp Val Gln Arg Gly Ala Ala Ala Glu Asp
 35 40 45
 His Pro Gly Gly Val Leu Ala Gln Gly Gly Ala Gln Leu Glu Ala Val
 50 55 60
 Ala Gly Ala Ala Ser Gln Glu Pro Asp Val Gly Gly Pro Arg Met Ala
 65 70 75 80
 Val Glu Glu Glu Val Ala Val Gly Ala Val Leu Val Leu Ala Asp Ala
 85 90 95
 Gly Leu Asp Gln Arg Arg Val Leu Gln Gly Arg Glu Pro Ala Gly His
 100 105 110
 Leu Gly Pro Gly Arg Phe Gln Gln Gly Arg Gly Asp Arg Pro Leu Ala
 115 120 125
 Arg Arg Gly Ile Asp Gly Leu Ala Pro Gly Val Val Arg His Leu Glu
 130 135 140
 Ala Ala Val Leu Val Ala Gly Asp Ala Val Val Asp Pro Leu Ala Glu
 145 150 155 160
 Ile Asp Pro Asp Arg Thr Ala Ala Leu Leu Glu Ala Arg Val Ala Arg
 165 170 175
 Arg Arg Ala Glu Glu Glu His Leu Leu Ala Gly Val Ala Glu Glu Pro
 180 185 190
 Leu Thr Asp His Val Arg Glu Gln Pro Gly Gln Pro Gly Thr Ala Gly
 195 200 205
 Glu Asp Val Glu Val Gly Arg Glu Ser Gly Ala Val Arg Lys Val Lys

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210 215 220
 Pro Leu Gln Gly Pro Arg Asp His Gly Gly Leu Val Leu Pro Ala
 225 230 235 240
 Leu Ala Leu Glu Gln Leu His His Gly Pro Ala Gly Ala Pro Gly Glu
 245 250 255
 Gln Gly Ala Gly Phe Leu Leu Val Pro Asp Arg Ala Asp Ala Val Glu
 260 265 270
 Ile Asp Leu Gly Glu Ala Ala Pro Gly Leu Pro Leu Arg Gln Leu Gly
 275 280 285
 Asp Arg Gln Pro Arg Val Leu Gln Lys Arg Lys Gly Val Ala Asp Val
 290 295 300
 Ala Val Val Leu Ala Ala His Pro Glu Asp Pro Gly Pro Phe Val Gln
 305 310 315 320
 Pro Val Thr Gly Leu Asp Phe Gly Val Pro Glu Leu Glu Gly Ala
 325 330 335
 Gly Asp Pro Leu His Val Gln Thr Val Gly Ser Val Gly Ala Ala Asp
 340 345 350
 Asp Pro Arg Leu Ala Thr Gly Ala Gly Val Pro Arg Thr Pro
 355 360 365
 Gly Val Gln Glu Gly His Pro Gly Ser Ala Ala Glu Met Gln Gly
 370 375 380
 Gly Pro Ala Ala Glu Gly Ala Gly Ala Asp Asp Gly Asp Met Gly Met
 385 390 395 400
 Gly Gly His Gly Gly Arg Lys Val Ile Ala Ala Arg Ser Phe Ala Gly
 405 410 415
 Ile Pro Ser Pro Thr Gly Val Ser Trp Lys Ile Arg Arg Arg Ser
 420 425 430
 Thr Cys Asn Arg Thr Glu Thr
 435

<210> 139
 <211> 1524
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 139
 atgaaaacat tcaaccttaa acccacactt ttaccttttaa ctttgctgct gagttcgccg 60
 gtattggcgg cacaaaatgg aactatgatg cagtatttcc attggtatgt gccaaatgac 120
 ggcgcactct ggacacaagt tgaaaacaat gcgccagcac tatccgacaa cggttttaca 180
 gcgctgtggt tgccaccagc atataaaggc gcagggtgga gcaacgacgt tggttacggt 240
 gtttacgata tgtatgactt aggggagttt gatcaaaaag gatcggtagc aactaagtac 300
 ggcaccaaaag accaatatct aaatgccatc aaagcagcac acaaaaacaa tatccaaatt 360
 tatggtgacg tagtggtcaa ccatcgtagc ggtgcagatg gcaagtcgtg ggtcgatacc 420
 aagcgtgtgg attggaataa ccgcaatatt gaacttggcg ataaatggat tgaagcatgg 480
 gttgaattta gcttcccagg acgtaacgat aaatactcag acttccattg gacgtggtat 540
 cactttgatg gcgtcgattg ggatgacgca ggtaaagaga aagcgatctt taaattcaaa 600
 ggtgatggta aagcatggga ttgggaagtc agttctgaaa aaggcaacta tgactacctc 660
 atgtacgcag acttagacat ggatcaccca gaagtgaagc aagagctgaa agattggggg 720
 gaatggtact taaacatgac ggggtgtgat ggcttccgaa tggatgcagt gaaacacatc 780
 aaatatcagt acctacaaga gtggatcgat tacttgcgta agaaaacggg caaagagctc 840
 tttaccgttg gtgagtactg gaactacgac gtgaacaatc tgcacaactt tatgactaag 900
 acttctggca gcatgtcatt gtttgatgag cctttacata tgaacttcta taacgcttca 960
 cgctctgggt gcaactttga tatgcgccga atcatggatg gcaccttgat gaaagacaac 1020
 ccagtgaag cagtaacact ggttgagaac catgatacgc aaccactaca ggccttagag 1080
 tctccggtgg attggtgggt caaaccactt gcgtacgcgt tcattttgct tcgtgaggaa 1140
 ggttatccgt cagtcttcta cgcagattac tacggtgcgc aatacagcga taaagggcac 1200
 gatatcaaca tggtgaaagt gccttacatt gagcaattgg tgaaagcgcg taaagattat 1260
 gcttatggta aacaacattc ttaccttgac cactgggatg tgattgggtg gacacgagaa 1320
 ggggatgcgg aacatccgaa ctctatggcg gttatcatga gtgatgggtc tggcggaaca 1380
 aagtggatgt acacaggttc accgagcaca cgttatgtcg ataaactagg tattcgtacc 1440
 gaagaagtat ggactaacgc tagtggatgg gccgaattcc cagtgaacgg cggatcggtt 1500
 tctgtttggg ttggcgtaa ataa 1524

<210> 140
 <211> 507

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<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

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<400> 140
Met Lys Thr Phe Asn Leu Lys Pro Thr Leu Leu Pro Leu Thr Leu Leu
1      5      10      15
Leu Ser Ser Pro Val Leu Ala Ala Gln Asn Gly Thr Met Met Gln Tyr
20      25      30
Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu
35      40
Asn Asn Ala Pro Ala Leu Ser Asp Asn Gly Phe Thr Ala Leu Trp Leu
50      55      60
Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly
65      70      75      80
Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ser Val
85
Arg Thr Lys Tyr Gly Thr Lys Asp Gln Tyr Leu Asn Ala Ile Lys Ala
100      105      110
Ala His Lys Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His
115      120      125
Arg Gly Gly Ala Asp Gly Lys Ser Trp Val Asp Thr Lys Arg Val Asp
130      135      140
Trp Asn Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp
145      150      155      160
Val Glu Phe Ser Phe Pro Gly Arg Asn Asp Lys Tyr Ser Asp Phe His
165      170      175
Trp Thr Trp Tyr His Phe Asp Gly Val Asp Trp Asp Asp Ala Gly Lys
180      185      190
Glu Lys Ala Ile Phe Lys Phe Lys Asp Gly Lys Ala Trp Asp Trp
195      200      205
Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp
210      215      220
Leu Asp Met Asp His Pro Glu Val Lys Gln Glu Leu Lys Asp Trp Gly
225      230      235      240
Glu Trp Tyr Leu Asn Met Thr Gly Val Asp Gly Phe Arg Met Asp Ala
245      250      255
Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp Tyr Leu
260      265      270
Arg Lys Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn
275      280      285
Tyr Asp Val Asn Asn Leu His Asn Phe Met Thr Lys Thr Ser Gly Ser
290      295      300
Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala Ser
305      310      315      320
Arg Ser Gly Gly Asn Phe Asp Met Arg Arg Ile Met Asp Gly Thr Leu
325      330      335
Met Lys Asp Asn Pro Val Lys Ala Val Thr Leu Val Glu Asn His Asp
340      345      350
Thr Gln Pro Leu Gln Ala Leu Glu Ser Pro Val Asp Trp Trp Phe Lys
355      360      365
Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Tyr Pro Ser
370      375      380
Val Phe Tyr Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly His
385      390      395      400
Asp Ile Asn Met Val Lys Val Pro Tyr Ile Glu Gln Leu Val Lys Ala
405      410      415
Arg Lys Asp Tyr Ala Tyr Gly Lys Gln His Ser Tyr Leu Asp His Trp
420      425      430
Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser
435      440      445
Met Ala Val Ile Met Ser Asp Gly Pro Gly Gly Thr Lys Trp Met Tyr
450      455      460
Thr Gly Ser Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr
465      470      475      480
Glu Glu Val Trp Thr Asn Ala Ser Gly Trp Ala Glu Phe Pro Val Asn

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485 490 495
 Gly Gly Ser Val Ser Val Trp Val Gly Val Lys
 500 505

<210> 141
 <211> 1401
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 141
 atgaaaccaa taaataccct actcatatcc gcccttgctg tttgttcttt cagttccgcg 60
 acitacgccg atactatttt gcacgcgttc aattggaagt attcagatgt gacggccaac 120
 gcgaatcaaa ttgctcaagc tggttataag aaagtgcctg ttgcgcctgc aatgaaatcg 180
 agtggcagcc aatggtgggc tcgctatcaa cctcaagatc tacgcactat cgattctcct 240
 ttgggcaata aacaagattt agccgcaatg attgccgcac tcaaagggtg gggcgtcgat 300
 gtgtatgccg atgtggtact caaccatatg gcgaatgaaa gctggaagcg aagtgacttg 360
 aattaccctg gcacagaagt gctaaacgat tatgctagcc gttcaagcta ctatgctgac 420
 cagactctgt ttggcaacct agcacaaggt tatgtgtcag cgaacgactt tcatccagcg 480
 ggctgtattt cagattggaa cgaccctggt catgttcagt attggcgttt gtgtggcgca 540
 gatggtgatg taggtttacc tgaccttgat ccaaacaact ggggtggtttc acaacagcgt 600
 ttgtatctga aagcgctaaa agatatgggc atcaaagggt tccgaattga tgcagtgaag 660
 cacatgagcc aataccaaat cgatcaggta ttcacgtctg aaattactgc gaacatgcat 720
 gtgtttggcg aagtgattac tagcggtgga gcagggaata gcggctatga atcgttctta 780
 gcgccttacc tgaataatac taatcactct gcctacgatt tcccgtgttt tgcatcgatt 840
 cgctcggcat tttctatggg gggcgggtta aatcaactgc atgatcctaa agcgtacggt 900
 caggcacttg atgataatcg ctcgatcacc tttgcgatca cacatgatat tccaaccaat 960
 gacggcttcc gctaccaaat tatggacca caagacgagc agcttgctta cgcgatatatc 1020
 cttggtaaa acggtggcac gccgctgatc tacagtgatg atcttcctga ttctgaagac 1080
 aaggataacg gtcgttgggg caatgtttgg aacagttcga caatgaaaaa catgttgagc 1140
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 ttgtttaagc gtggcaaaga aggtgtttgt ggtattaaca agtgtggtga aacgcgtggc 1260
 gtgacggttg atacctacca acatgagttt aattggcatg ttcaatacaa agacgtgtta 1320
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<210> 142
 <211> 466
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 142
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 Phe Ser Ser Ala Thr Tyr Ala Asp Thr Ile Leu His Ala Phe Asn Trp
 20 25 30
 Lys Tyr Ser Asp Val Thr Ala Asn Ala Asn Gln Ile Ala Gln Ala Gly
 35 40 45
 Tyr Lys Lys Val Leu Val Ala Pro Ala Met Lys Ser Ser Gly Ser Gln
 50 55 60
 Trp Trp Ala Arg Tyr Gln Pro Gln Asp Leu Arg Thr Ile Asp Ser Pro
 65 70 75 80
 Leu Gly Asn Lys Gln Asp Leu Ala Ala Met Ile Ala Ala Leu Lys Gly
 85 90 95
 Val Gly Val Asp Val Tyr Ala Asp Val Leu Asn His Met Ala Asn
 100 105 110
 Glu Ser Trp Lys Arg Ser Asp Leu Asn Tyr Pro Gly Thr Glu Val Leu
 115 120 125
 Asn Asp Tyr Ala Ser Arg Ser Ser Tyr Tyr Ala Asp Gln Thr Leu Phe
 130 135 140
 Gly Asn Leu Ala Gln Gly Tyr Val Ser Ala Asn Asp Phe His Pro Ala
 145 150 155 160
 Gly Cys Ile Ser Asp Trp Asn Asp Pro Gly His Val Gln Tyr Trp Arg

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Leu	Cys	Gly	Ala	165	Asp	Gly	Asp	Val	Gly	170	Leu	Pro	Asp	Leu	Asp	175	Pro	Asn
Asn	Trp	Val	Val	180	Ser	Gln	Gln	Arg	Leu	185	Tyr	Leu	Lys	Ala	190	Leu	Lys	Asp
Met	Gly	Ile	Lys	195	Gly	Phe	Arg	Ile	Asp	200	Ala	Val	Lys	His	Met	Ser	Gln	
Tyr	Gln	Ile	Asp	210	Gln	Val	Phe	Thr	Ser	215	Glu	Ile	Thr	Ala	Asn	Met	His	
Val	Phe	Gly	Glu	225	Val	Ile	Thr	Ser	Gly	230	Gly	Ala	Gly	Asn	Ser	Gly	Tyr	
Glu	Ser	Phe	Leu	245	Ala	Pro	Tyr	Leu	Asn	250	Asn	Thr	Asn	His	Ser	Ala	Tyr	
Asp	Phe	Pro	Leu	260	Phe	Ala	Ser	Ile	Arg	265	Ser	Ala	Phe	Ser	Met	Gly	Gly	
Gly	Leu	Asn	Gln	275	Leu	His	Asp	Pro	Lys	280	Ala	Tyr	Gly	Gln	Ala	Leu	Asp	
Asp	Asn	Arg	Ser	290	Ile	Thr	Phe	Ala	Ile	295	Thr	His	Asp	Ile	Pro	Thr	Asn	
Asp	Gly	Phe	Arg	305	Tyr	Gln	Ile	Met	Asp	310	Pro	Gln	Asp	Glu	Gln	Leu	Ala	
Tyr	Ala	Tyr	Ile	325	Leu	Gly	Lys	Asp	Gly	330	Gly	Thr	Pro	Leu	Ile	Tyr	Ser	
Asp	Asp	Leu	Pro	340	Asp	Ser	Glu	Asp	Lys	345	Asp	Asn	Gly	Arg	Trp	Gly	Asn	
Val	Trp	Asn	Ser	355	Ser	Thr	Met	Lys	Asn	360	Met	Leu	Ser	Phe	His	Asn	Ala	
Met	Gln	Gly	Lys	370	Thr	Met	Thr	Met	Ile	375	Ser	Ser	Asp	His	Cys	Thr	Leu	
Leu	Phe	Lys	Arg	385	Gly	Lys	Glu	Gly	Val	390	Val	Gly	Ile	Asn	Lys	Cys	Gly	
Glu	Thr	Arg	Gly	405	Val	Thr	Val	Asp	Thr	410	Tyr	Gln	His	Glu	Phe	Asn	Trp	
His	Val	Gln	Tyr	420	Lys	Asp	Val	Leu	Ser	425	Ser	Ala	Thr	Glu	Thr	Val	Thr	
Ser	Arg	Tyr	His	435	Thr	Phe	Asn	Leu	Pro	440	Pro	Arg	Ser	Ala	Arg	Met	Phe	
Lys	Leu			450						455								
				465														

<210> 143
 <211> 1422
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 143					
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tacagcgaca	ttaccgcgcca	agcagagcaa	attgcgcaag	ctgggttataa	aaaagtactg
atttcaccgc	cgctgaagtc	cacaggccca	caatggtggg	cacgtttacca	accacaggac
attcgagtga	ttgactcccc	tgtcggcaac	aagcaagatt	tacaagccct	cattgcagcc
ttaaaggcac	aaggcgttga	agtatacgca	gacatcgta	tcaaccacat	ggccaacgaa
agctggaaac	gagacgatct	gaactaccgc	ggaagtgatt	tacttaccca	atacagccaa
aatatggctt	acatgaacca	gcaaaaattg	tttggagatt	tagagcaaaa	tcagttctct
gccaatgatt	ttcaccgggc	tggctgcatt	actgattgga	gtaaccggg	gcatgttcaa
tactggcgct	tatgtggtgg	taatggtgac	actgggttac	ctgatcttga	tcctaactcg
tgggtgatcg	atcaacaaaa	acgttattta	cgtgctttga	aagacatggg	aataaagggc
ttccgagttg	atgcggtaaa	acacatgagc	gattacaaa	tcaaccaagt	gtttacggca
gacatcatcg	caggcttaca	tgtatttggt	gaagtgatca	ccagtgggtg	caagggcagc
aatgactacc	actcttttct	ggaaccgtat	ttaaataaca	ccaatcacgc	cgcgatgac
ttcccgctat	ttgcctctat	ccgaaatgca	tttagttatc	atggcagctt	gtctcaatta
catgatccac	aagcttacgg	gcaagcactt	cctaacgaca	gagccattac	tttcaccatc
actcacgaca	ttccaaccaa	tgatggtttc	cgttacaaa	tcattggatcc	aaccagtga
aaactcgct	acgcgtacat	tctaggcaaa	gatgggggta	gccacttat	ctatagcgat
gctttagacc	caagtgaaga	taaagataag	ggccgctggc	gtgatgtatg	gaaccaagaa

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tacatgggta	acatgatcag	cttccacaac	aaggtgcaag	gtaaaagcat	ggaggtcatg	1200
tacagcgatc	aatgcttgct	ggtctttaaa	cgtgaaaaac	aaggcttagt	cggtattaat	1260
aagtgcgctg	aaagccgtac	ctacaccata	gatacccatc	gttttgaatt	taactgggtac	1320
caaccgtaca	acgacacatt	aagccagcac	agcgagacct	ttagcagccg	ttatcatgct	1380
ctgaccattc	cggcgcaaac	agcacgaatg	ttggcgctat	aa		1422

<210> 144
 <211> 473
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 144

Met	Pro	Lys	Ser	Thr	Phe	Thr	Lys	Ser	Ile	Thr	Lys	Ser	Leu	Leu	Ala
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Thr	Ser	Val	Val	Val	Ser	Leu	Leu	Pro	Ala	Tyr	Ala	Gln	Ala	Asp	Thr
			20					25					30		
Ile	Leu	His	Ala	Phe	Asn	Trp	Lys	Tyr	Ser	Asp	Ile	Thr	Arg	Gln	Ala
		35					40					45			
Glu	Gln	Ile	Ala	Gln	Ala	Gly	Tyr	Lys	Lys	Val	Leu	Ile	Ser	Pro	Pro
	50					55					60				
Leu	Lys	Ser	Thr	Gly	Pro	Gln	Trp	Trp	Ala	Arg	Tyr	Gln	Pro	Gln	Asp
65					70					75					80
Ile	Arg	Val	Ile	Asp	Ser	Pro	Val	Gly	Asn	Lys	Gln	Asp	Leu	Gln	Ala
				85					90					95	
Leu	Ile	Ala	Ala	Leu	Lys	Ala	Gln	Gly	Val	Glu	Val	Tyr	Ala	Asp	Ile
			100					105					110		
Val	Leu	Asn	His	Met	Ala	Asn	Glu	Ser	Trp	Lys	Arg	Asp	Asp	Leu	Asn
		115					120					125			
Tyr	Pro	Gly	Ser	Asp	Leu	Leu	Thr	Gln	Tyr	Ser	Gln	Asn	Met	Ala	Tyr
	130					135					140				
Met	Asn	Gln	Gln	Lys	Leu	Phe	Gly	Asp	Leu	Glu	Gln	Asn	Gln	Phe	Ser
145					150					155					160
Ala	Asn	Asp	Phe	His	Pro	Ala	Gly	Cys	Ile	Thr	Asp	Trp	Ser	Asn	Pro
				165					170					175	
Gly	His	Val	Gln	Tyr	Trp	Arg	Leu	Cys	Gly	Gly	Asn	Gly	Asp	Thr	Gly
			180					185					190		
Leu	Pro	Asp	Leu	Asp	Pro	Asn	Ser	Trp	Val	Ile	Asp	Gln	Gln	Lys	Arg
		195					200					205			
Tyr	Leu	Arg	Ala	Leu	Lys	Asp	Met	Gly	Ile	Lys	Gly	Phe	Arg	Val	Asp
	210					215					220				
Ala	Val	Lys	His	Met	Ser	Asp	Tyr	Gln	Ile	Asn	Gln	Val	Phe	Thr	Pro
225					230					235					240
Asp	Ile	Ile	Ala	Gly	Leu	His	Val	Phe	Gly	Glu	Val	Ile	Thr	Ser	Gly
				245					250					255	
Gly	Lys	Gly	Ser	Asn	Asp	Tyr	His	Ser	Phe	Leu	Glu	Pro	Tyr	Leu	Asn
			260					265					270		
Asn	Thr	Asn	His	Ala	Ala	Tyr	Asp	Phe	Pro	Leu	Phe	Ala	Ser	Ile	Arg
		275					280					285			
Asn	Ala	Phe	Ser	Tyr	His	Gly	Ser	Leu	Ser	Gln	Leu	His	Asp	Pro	Gln
	290					295					300				
Ala	Tyr	Gly	Gln	Ala	Leu	Pro	Asn	Asp	Arg	Ala	Ile	Thr	Phe	Thr	Ile
305					310					315					320
Thr	His	Asp	Ile	Pro	Thr	Asn	Asp	Gly	Phe	Arg	Tyr	Gln	Ile	Met	Asp
				325					330					335	
Pro	Thr	Ser	Glu	Lys	Leu	Ala	Tyr	Ala	Tyr	Ile	Leu	Gly	Lys	Asp	Gly
			340					345					350		
Gly	Ser	Pro	Leu	Ile	Tyr	Ser	Asp	Ala	Leu	Asp	Pro	Ser	Glu	Asp	Lys
		355					360					365			
Asp	Lys	Gly	Arg	Trp	Arg	Asp	Val	Trp	Asn	Gln	Glu	Tyr	Met	Val	Asn
	370					375					380				
Met	Ile	Ser	Phe	His	Asn	Lys	Val	Gln	Gly	Lys	Ser	Met	Glu	Val	Met
385					390					395					400
Tyr	Ser	Asp	Gln	Cys	Leu	Leu	Val	Phe	Lys	Arg	Glu	Lys	Gln	Gly	Leu
				405					410					415	
Val	Gly	Ile	Asn	Lys	Cys	Ala	Glu	Ser	Arg	Thr	Tyr	Thr	Ile	Asp	Thr

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			420						425				430				
His	Arg	Phe	Glu	Phe	Asn	Trp	Tyr	Gln	Pro	Tyr	Asn	Asp	Thr	Leu	Ser		
		435					440					445					
Gln	His	Ser	Glu	Thr	Phe	Ser	Ser	Arg	Tyr	His	Ala	Leu	Thr	Ile	Pro		
	450					455					460						
Ala	Gln	Thr	Ala	Arg	Met	Leu	Ala	Leu									
465					470												

<210> 145
 <211> 1542
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 145
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 gggaggaata aggcggaagc agcaacgata aataatggaa cattaatgca gtattttgag 120
 tggtagctc cgaatgatgg gaatcattgg aatcgtttgc gttatgatgc tgaaagtta 180
 gtcataagg gaatcacatc tgtatggata ccacctgcat ataaaggac ttcgcaaat 240
 gatgtagggt atggggccta tgatttatac gatttagggg agttcaatca aaaaggaacg 300
 gtgcggacga aatatgggac aaaggcacag ttgaaatctg caattgacgc tttacataag 360
 caaaacatcg acgtatacgg tgatgtagtt atgaatcata aagggtggggc tgattatact 420
 gaaaccgtaa cagctgttga ggtagaccgt aacaatcgaa atattgaagt atcagggtgat 480
 tatgaaatta gtgcgtggac gggttttaac tttccagggc gcagagatgc ttattctaata 540
 ttcaaatgga aatggtatca ttttgacgga acggattggg atgaaggaag gaaattaaac 600
 cgaatttata aatttagggg tataggtaaa gcgtgggact ggggaagtgtc tagcgaaaat 660
 ggaaattatg attatttgat gtatgcagat cttgattttg atcatccaga tgttgcaaat 720
 gaaatgaaaa gttggggaac gtggtatgct aatgaattaa atttagatgg atttcgttta 780
 gatgctgtta aacatattga tcatgaatat ttacgcgatt gggtaaatca tgtcagacag 840
 caaacgggga aagaaatggt tacgggtggct aaagataaaa aaatgatata ccagacttta 900
 aacaattatt tggcgaaagt caattataat caatctgtat ttgatgcacc gcttcattac 960
 aattttcatt atgcttcaac aggaaatggg aattatgata tgagaaatat tttaaatgga 1020
 acagtaatga aaaatcatcc tgcactcgca gttactctcg ttgagaatca tgattctcaa 1080
 cctggggcaat cattggaatc tgtagtaagt ccgtggttta agccgctggc atatgcattt 1140
 attttaactc gtgcagaggg ctatccttca gttttttatg gtgattacta tgggacaagc 1200
 ggaaatagta gttatgaaat tccagcggtta aaagataaaa ttgatccaat tttgacggca 1260
 cgaaaaaact ttgcatatgg tacgcagcgt gattatttag accatccaga tgtgattggc 1320
 tggacaagag aaggagatag tgtacatgct aagtcctggt tagcggcatt aatctccgat 1380
 ggaccaggag gatcaaagt gatggatggt ggaaagaata acgctgggga agtatggtac 1440
 gatattacgg gtaatcaaac aaatactgta acaattaata aagatggatc ggggcaattc 1500
 catgtaagtg gaggtctctg ttctatatat gttcaacagt aa 1542

<210> 146
 <211> 513
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 146
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 Gly Thr Leu Met Gln Tyr Phe Glu Trp Tyr Ala Pro Asn Asp Gly Asn
 35 40 45
 His Trp Asn Arg Leu Arg Tyr Asp Ala Glu Ser Leu Ala His Lys Gly
 50 55 60
 Ile Thr Ser Val Trp Ile Pro Pro Ala Tyr Lys Gly Thr Ser Gln Asn
 65 70 75 80
 Asp Val Gly Tyr Gly Ala Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn
 85 90 95
 Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ala Gln Leu Lys
 100 105 110
 Ser Ala Ile Asp Ala Leu His Lys Gln Asn Ile Asp Val Tyr Gly Asp

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Val	Val	115	Met	Asn	His	Lys	Gly	120	Gly	Ala	Asp	Tyr	Thr	125	Glu	Thr	Val	Thr
Ala	Val	130	Glu	Val	Asp	Arg	Asn	135	Asn	Arg	Asn	Ile	Glu	140	Val	Ser	Gly	Asp
Tyr	Glu	145	Ile	Ser	Ala	Trp	Thr	150	Gly	Phe	Asn	Phe	Pro	155	Gly	Arg	Arg	Asp
Ala	Tyr	160	Ser	Asn	Phe	Lys	Trp	165	Lys	Trp	Lys	Trp	His	170	Phe	Asp	Gly	Thr
Trp	Asp	175	Glu	Gly	Arg	Lys	Leu	180	Asn	Arg	Ile	Tyr	Lys	185	Phe	Arg	Gly	Ile
Gly	Lys	190	Ala	Trp	Asp	Trp	Glu	195	Val	Ser	Ser	Glu	Asn	200	Gly	Asn	Tyr	Asp
Tyr	Leu	205	Met	Tyr	Ala	Asp	Leu	210	Asp	Phe	Asp	His	Pro	215	Asp	Val	Ala	Asn
Glu	Met	220	Lys	Ser	Trp	Gly	Thr	225	Trp	Tyr	Ala	Asn	Glu	230	Leu	Asn	Leu	Asp
Gly	Phe	235	Arg	Leu	Asp	Ala	Val	240	Lys	His	Ile	Asp	His	245	Glu	Tyr	Leu	Arg
Asp	Trp	250	Val	Asn	His	Val	Arg	255	Gln	Thr	Gly	Lys	Glu	260	Met	Phe	Thr	
Val	Ala	265	Glu	Tyr	Trp	Gln	Asn	270	Asp	Ile	Gln	Thr	Leu	275	Asn	Asn	Tyr	Leu
Ala	Lys	280	Val	Asn	Tyr	Asn	Gln	285	Ser	Val	Phe	Asp	Ala	290	Pro	Leu	His	Tyr
Asn	Phe	295	His	Tyr	Ala	Ser	Thr	300	Gly	Asn	Gly	Asn	Tyr	305	Asp	Met	Arg	Asn
Ile	Leu	310	Asn	Gly	Thr	Val	Met	315	Lys	Asn	His	Pro	Ala	320	Leu	Ala	Val	Thr
Leu	Val	325	Glu	Asn	His	Asp	Ser	330	Gln	Pro	Gly	Gln	Ser	335	Leu	Glu	Ser	Val
Val	Ser	340	Pro	Trp	Phe	Lys	Pro	345	Leu	Ala	Tyr	Ala	Phe	350	Ile	Leu	Thr	Arg
Ala	Glu	355	Gly	Tyr	Pro	Ser	Val	360	Phe	Tyr	Gly	Asp	Tyr	365	Tyr	Gly	Thr	Ser
Gly	Asn	370	Ser	Ser	Tyr	Glu	Ile	375	Pro	Ala	Leu	Lys	Asp	380	Lys	Ile	Asp	Pro
Ile	Leu	385	Thr	Ala	Arg	Lys	Asn	390	Phe	Ala	Tyr	Gly	Thr	395	Gln	Arg	Asp	Tyr
Leu	Asp	400	His	Pro	Asp	Val	Ile	405	Gly	Trp	Thr	Arg	Glu	410	Gly	Asp	Ser	Val
His	Ala	415	Lys	Ser	Gly	Leu	Ala	420	Ala	Leu	Ile	Ser	Asp	425	Gly	Pro	Gly	Gly
Ser	Lys	430	Trp	Met	Asp	Val	Gly	435	Lys	Asn	Asn	Ala	Gly	440	Glu	Val	Trp	Tyr
Asp	Ile	445	Thr	Gly	Asn	Gln	Thr	450	Asn	Thr	Val	Thr	Ile	455	Asn	Lys	Asp	Gly
Ser	Gly	460	Gln	Phe	His	Val	Ser	465	Gly	Gly	Val	Ser	Ile	470	Tyr	Val	Gln	
Gln		475						480						485				
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		505						510										

<210> 147
 <211> 2343
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 147																		
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ggatttgacg	gtgtacagat	ttcccagcca	gcggaacata	agcgggctga	aggagtattg													240
tgggccgtat	atcagccggg	taattataag	aattttacaa	ccatgaccgg	taacgaggag													300
cagcttaagg	caatgatcaa	gacctgtaag	gatgcagggtg	ttaagggtgt	cgctgacgct													360
gttttcaacc	aaaaggctac	agacgggtgta	ggctggggcg	gttcaacttg	gagttataag													420

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aactaccctg acggattctc cggatcagat ttccatggag actgtttccat tgacaaaagc 480
tatactgatg caaataatgt cagaacctgt gcactctcag gtatgccgga cgttgccaca 540
gataactccg ctactcagga aaagattgca gattacctcg cttctttaat gaatatgggg 600
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ctttcaaaaa ctgcacagaa gactggaaga agacctcctg catatctgga agtaatcgga 720
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aaggcactgg aactcagtac ctggctcggt gcaaattcag aaacattcgt aaacaatcat 900
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gattataatc tggctcagtc ctggcttgct gtatggcctg taggtacagt aagacagata 1020
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gcacagaata ctaccaacag tcctataaacc cagacattct ctggtgaagt acctgacgga 1320
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gcctgtgttt gtaaggggga aaccaccggt aatgggtgtgt gcgtcagctg gtgtaatgct 1560
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gatgaacctg tgtctcttgt tggtgattat gttctttcca ttaacgataa gaccatggaa 1920
tatacattca ccaaggcaga tgaagtaact aatcagccac cggttgcacg atttacgcg 1980
acagttaacg gtctgaccgt ttcttttgcc aataattcat ccgaccctga gaatgatgaa 2040
ttaacctaca gctggaattt cggtaatggt aaaacatcat ccgagaaaagc tcctagcata 2100
acctatgaag aatccggtaa gtatactgtt actttaaagg ttactgattc agctaataac 2160
actgatacat ttactaaaga tataactgta acagcacctt ctagtggcaa gtacttaag 2220
gttgacgtca gaggttcgca tgataattac ggaactgacg tgtaaaccaa gaacggttct 2280
gattggaccg gcgtctttga attctttgga tccactagtg tcgacctgca ggcgcgcgag 2340
ctc

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<210> 148
 <211> 781
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 148
Met Ser Leu Asn Asn Phe Lys Val Lys Leu Leu Ser Phe Ala Val Ser
 1          5          10          15
Ser Ala Val Leu Ser Leu Ala Pro Asn Leu Ala Asn Ala Ala Asn Phe
          20          25          30
Glu Ser Glu Met Val Ile Ile His Pro Phe Gln Trp Thr Tyr Asp Asn
          35          40          45
Ile Ala Lys Glu Cys Thr Glu Tyr Leu Gly Pro Ala Gly Phe Asp Gly
          50          55          60
Val Gln Ile Ser Gln Pro Ala Glu His Lys Arg Ala Glu Gly Val Trp
 65          70          75          80
Trp Ala Val Tyr Gln Pro Val Asn Tyr Lys Asn Phe Thr Thr Met Thr
          85          90          95
Gly Asn Glu Glu Gln Leu Lys Ala Met Ile Lys Thr Cys Asn Asp Ala
          100          105          110
Gly Val Lys Val Phe Ala Asp Ala Val Phe Asn Gln Lys Ala Thr Asp
          115          120          125
Gly Val Gly Trp Gly Gly Ser Thr Trp Ser Tyr Lys Asn Tyr Pro Asp
          130          135          140
Gly Phe Ser Gly Ser Asp Phe His Gly Asp Cys Ser Ile Asp Lys Ser
 145          150          155          160
Tyr Thr Asp Ala Asn Asn Val Arg Thr Cys Ala Leu Ser Gly Met Pro
          165          170          175
Asp Val Ala Thr Asp Asn Ser Ala Thr Gln Glu Lys Ile Ala Asp Tyr
          180          185          190
Leu Ala Ser Leu Met Asn Met Gly Val Tyr Gly Phe Arg Ile Asp Ala

```

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Ala	Lys	195	Met	Gly	Tyr	Asn	200	Ile	Asn	Ser	Ile	205	Leu	Ser	Lys	Thr
Ala	Gln	210	Lys	Thr	Gly	Arg	215	Pro	Pro	Ala	Tyr	220	Leu	Glu	Val	Ile
Ala	Gly	225	Asn	Glu	Ala	230	Asp	Ile	Gln	Pro	235	Asp	Lys	Tyr	Thr	Phe
Glu	Asn	245	Ala	Val	Thr	245	Asp	Phe	Gly	250	Tyr	Val	Trp	Asp	Ala	255
Ser	Phe	260	Gly	Lys	Gly	260	Asn	Tyr	Gly	265	Lys	Ala	Leu	Glu	Leu	270
Leu	Gly	275	Ala	Asn	Ser	280	Thr	Phe	Val	Asn	285	Asn	His	Asp	Asp	290
Gly	Arg	295	Cys	Ser	Ala	300	Gly	Ser	Cys	305	Met	Lys	Thr	Gln	Asn	310
Asp	Tyr	315	Asn	Leu	Ala	320	Gln	Ser	Trp	325	Leu	Ala	Val	Trp	Pro	330
Val	Arg	335	Gln	Ile	Tyr	340	Ser	Gly	Tyr	345	Ser	Phe	Pro	Val	Lys	350
Pro	Tyr	355	Arg	Val	Ser	360	Asp	Ala	Thr	365	His	Asp	Gln	Gly	Gly	370
Ala	Asp	375	Arg	Cys	Glu	380	Gly	Gly	Trp	385	Cys	Gln	His	Arg	Val	390
Val	Leu	395	Asn	Ser	Pro	400	Arg	Phe	Ala	405	Arg	Ala	Thr	Arg	Gly	410
Ser	Thr	415	Lys	Gly	Phe	420	Asp	Asn	Gly	425	Ala	Leu	Trp	Phe	Asn	430
Lys	Gly	435	Phe	Tyr	Ala	440	Gln	Asn	Thr	445	Thr	Asn	Ser	Pro	Ile	450
Phe	Ser	455	Val	Glu	Val	460	Pro	Asp	Gly	465	Asn	Tyr	Cys	Asp	Ile	470
Ser	Asp	475	Pro	Lys	Ser	480	Asn	Pro	Cys	485	Gly	Ala	Asp	Val	Val	490
Gly	Lys	495	Ala	Thr	Phe	500	Thr	Ile	Pro	505	Ala	Lys	Thr	Ala	Val	510
Thr	Asp	515	Ser	Asp	Trp	520	Cys	Gly	Lys	525	Gly	Val	Asp	Pro	Cys	530
Pro	Thr	535	Gly	Ala	Cys	540	Val	Cys	Lys	545	Gly	Glu	Thr	Thr	Val	550
Val	Cys	555	Val	Ser	Trp	560	Asn	Ala	His	565	Ser	Ser	Asn	Glu	Glu	570
Cys	Val	575	Leu	Asn	Pro	580	Asp	Ala	Asn	585	Cys	Gln	Ala	Asp	Ile	590
Thr	Lys	595	Gly	Lys	Leu	600	Cys	Tyr	Ala	605	Gly	Thr	Ser	Asn	Gly	610
Asp	Pro	615	Leu	Thr	Tyr	620	Asn	Arg	Lys	625	Thr	Gly	Phe	Trp	Thr	630
Thr	Leu	635	Asp	Gly	Ala	640	Gly	Asp	Thr	645	Ser	Gly	Ala	Gln	Arg	650
Thr	Asp	655	Gly	Cys	Ser	660	Trp	Thr	Gly	665	Thr	Val	Tyr	Gly	Ser	670
Ala	Gly	675	Lys	Leu	Asp	680	Val	Asn	Thr	685	Ser	Ser	Thr	Gly	Asp	690
Ser	Leu	695	Val	Gly	Asp	700	Val	Leu	Ser	705	Ile	Asn	Asp	Lys	Thr	710
Tyr	Thr	715	Phe	Thr	Lys	720	Ala	Asp	Glu	725	Val	Thr	Asn	Gln	Pro	730
Ser	Phe	735	Thr	Ala	Thr	740	Val	Asn	Gly	745	Leu	Thr	Val	Ser	Phe	750
Ser	Ser	755	Asp	Pro	Glu	760	Asn	Asp	Glu	765	Leu	Thr	Tyr	Ser	Trp	770
Asn	Gly	775	Lys	Thr	Ser	780	Ser	Glu	Lys	785	Ala	Pro	Ser	Ile	Thr	790
Ser	Gly	795	Lys	Tyr	Thr	800	Val	Thr	Leu	805	Lys	Val	Thr	Asp	Ser	810
Thr	Asp	815	Thr	Phe	Thr	820	Lys	Asp	Ile	825	Thr	Val	Thr	Ala	Pro	830
Lys	Tyr	835	Leu	Lys	Val	840	Ala	Val	Arg	845	Gly	Ser	His	Asp	Asn	850

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Asp	Leu	Leu	740	Thr	Lys	Asn	Gly	Ser	745	Asp	Trp	Thr	Gly	Val	750	Phe	Glu	Phe
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Phe	Gly	Ser	770	Thr	Ser	Val	Asp	Leu	775	Gln	Ala	Arg	Glu	Leu	780			

<210> 149
<211> 2502
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 149

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ggatttgacg	gggtgcagat	ttcccaggcg	gctgagcata	aagatgccgg	tggtgcatgg	240
tggggtagct	accagcctgt	aaacttcaag	agttttacta	ccatggttgg	taatgaagaa	300
cagcttagag	caatgattaa	aacctgtaac	gaggcagggt	ttaaggctct	tgccgatgcc	360
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aattatcctg	acggattttac	cagtgatgat	tttcatcata	ataactgcag	tataggtaat	480
aattattcag	atgcatgggt	agtaagattc	tgtgacctca	gtggcatgcc	ggatatagca	540
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gggttatagc	gattccgtat	tgatgctgcc	aagcacttta	gctatgatga	tatagacgct	660
attgtagaga	aaacagcaac	caaagcaggc	aggagacctc	ctgtctatat	ggaggttatc	720
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gccgttgtaa	cagattttac	ttatgctaata	agcatgcata	atatttttaa	cggaaagcgg	840
tatgccaaag	ctttgaacat	ggggctaggg	catgtttgat	ctgaaaatgc	cgaagtcttt	900
ataagtaatc	atgataatga	atggggaaga	aagtctgccg	gttctctgct	aataagaacc	960
cagaataatc	cggattacca	tctggctcag	tcctggctcg	cagtttgacc	tttaggcaag	1020
gtagacacag	tttattctgc	atatcagttc	ccggctcttg	aagatagttg	tgagcgggtc	1080
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ttgtgtcagc	accgtgtacc	gtttgtgctc	aattctccta	gatttgcaag	agcaaccaga	1200
gggacagtcg	ttactactaa	aggttttgat	gacggagctt	tgtggtttaa	cagaggaagc	1260
aagggcttct	atgcccagaa	tactaccggc	agttctataa	ctcatacatt	ctcagttgaa	1320
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ggagcggatg	tcactgtaag	cggaggtaaa	gcaaccttta	ccattccggc	aaagaccgcc	1440
gtagctatct	gtactgatga	aaagtgggtg	ggcaaggggg	ttgacctttg	tgaaagcgat	1500
cctaccgggt	ccgcctgtgt	atgtaagggt	gaaaccacag	ttaacggcgt	atgtgtaagc	1560
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tcagccaccg	gagatgaacc	ggtttacttg	acaggtaaat	atgttctttc	cataaatgat	1920
aagaccatgg	aatacacatt	cattcctgca	ggcagtgtaa	acaagcctcc	ggttgcgtca	1980
tttactccga	ctgttaaaga	tctgactgta	tcttttgtca	ataattcatc	cgaccttgag	2040
aatgatgaat	taacctacag	ctggaatttc	ggtaacggta	aaacctcatc	tgaaaagaat	2100
ccgagtgtta	catatgataa	agccggtaaa	tatactgttt	caactcaaagt	aaccgatact	2160
gcaaacaaca	ctgataccaa	aacactggaa	atcgatttaa	catctcctgt	taacggaaaa	2220
tattccaagg	ttgcagtcag	aggttcacat	gataactacg	gaacaaatct	gttaaccagg	2280
aatggttcag	aatggaccgg	tatctttgaa	ttcagtaaga	caaccaaatt	caagcttgaa	2340
gctctgcttc	ctgcagctga	ccagtgtatc	ttcctcggcg	gtaatcgagg	tgaggcattg	2400
actgcctccg	gtggatttat	atctcttcct	gccggaaggt	atactataaa	gtttaatgag	2460
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<210> 150
<211> 834
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 150

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1	Ser	Ala	Val	Leu	5 Thr	Leu	Ala	Ala	10 Asn	Val	Ala	Asn	Ala	15 Lys	Asn	Tyr
	Glu	Ser	Glu	Met	20 Val	Ile	Ile	His	25 Pro	Phe	Gln	Trp	Thr	30 Tyr	Asp	Asn
	Ile	Ala	Lys	Glu	35 Cys	Thr	Glu	Tyr	40 Leu	Gly	Pro	Ala	Gly	45 Phe	Asp	Gly
	Val	Gln	Ile	Ser	50 Gln	Ala	Ala	Glu	55 His	Lys	Asp	Ala	Gly	60 Gly	Ala	Trp
65	Trp	Gly	Thr	Tyr	70 Gln	Pro	Val	Asn	75 Phe	Lys	Ser	Phe	Thr	80 Thr	Met	Val
	Gly	Asn	Glu	Glu	85 Gln	Leu	Arg	Ala	90 Met	Ile	Lys	Thr	Cys	95 Asn	Glu	Ala
	Gly	Val	Lys	Val	100 Phe	Ala	Asp	Ala	105 Val	Ile	Asn	Gln	Lys	110 Ala	Gly	Asp
	Gly	Val	Gly	Ile	115 Gly	Gly	Ser	Thr	120 Phe	Gly	Asn	Tyr	Asn	125 Tyr	Pro	Asp
	Gly	Phe	Thr	Ser	130 Phe	Asp	Asp	Phe	135 His	His	Asn	Asn	Cys	140 Ser	Ile	Gly
145	Asn	Tyr	Ser	Asp	145 Trp	Val	Val	Arg	150 Phe	Cys	Asp	Leu	Ser	155 Gly	Met	
	Pro	Asp	Ile	Ala	165 Thr	Asp	Asn	Asp	170 Ser	Thr	Arg	Asn	Lys	175 Ile	Ala	Asp
	Tyr	Phe	Ala	Ser	180 Leu	Met	Asn	Met	185 Gly	Val	Tyr	Gly	Phe	190 Arg	Ile	Asp
	Ala	Ala	Lys	His	195 Phe	Ser	Tyr	Asp	200 Asp	Ile	Asp	Ala	Ile	205 Val	Glu	Lys
	Thr	Ala	Thr	Lys	210 Ala	Gly	Arg	Arg	215 Pro	Pro	Val	Tyr	Met	220 Glu	Val	Ile
225	Gly	Asn	Pro	Gly	225 Gln	Glu	Ala	Asp	230 Asp	Ile	Gln	Pro	Asn	235 Lys	Tyr	Thr
	Trp	Ile	Asp	Asn	240 Ala	Val	Val	Thr	245 Asp	Phe	Thr	Tyr	Ala	250 Asn	Ser	Met
	His	Asn	Ile	Phe	255 Asn	Gly	Ser	Gly	260 Tyr	Ala	Lys	Ala	Leu	265 Asn	Met	Gly
	Leu	Gly	His	Val	270 Asp	Ala	Glu	Asn	275 Ala	Glu	Val	phe	Ile	280 Ser	Asn	His
	Asp	Asn	Glu	Trp	285 Gly	Arg	Lys	Ser	290 Ala	Gly	Ser	Cys	Ser	300 Ile	Arg	Thr
305	Gln	Asn	Asn	Pro	305 Asp	Tyr	His	Leu	310 Ala	Gln	Ser	Trp	Leu	315 Ala	Val	Trp
	Pro	Leu	Gly	Lys	320 Val	Arg	Gln	Ile	325 Tyr	Ser	Ala	Tyr	Gln	330 Phe	Pro	Val
	Phe	Glu	Asp	Ser	335 Cys	Glu	Arg	Val	340 Ser	Gln	Gln	Ala	His	345 Asp	Gln	Gly
	Gly	Pro	Ile	Gly	350 Ala	Ala	Arg	Cys	355 Glu	Gly	Gly	Trp	Leu	360 Cys	Gln	His
	Arg	Val	Pro	Phe	365 Val	Leu	Asn	Ser	370 Phe	Arg	Phe	Ala	Arg	375 Ala	Thr	Arg
385	Gly	Thr	Val	Val	380 Thr	Thr	Lys	Gly	385 Phe	Asp	Asp	Gly	Ala	390 Leu	Trp	Phe
	Asn	Arg	Gly	Ser	405 Lys	Gly	Phe	Tyr	410 Ala	Gln	Asn	Thr	Thr	415 Gly	Ser	Ser
	Ile	Thr	His	Thr	420 Phe	Ser	Val	Glu	425 Leu	Pro	Asp	Gly	Asn	430 Tyr	Cys	Asp
	Ile	Leu	Gly	Ala	435 Thr	Asp	Pro	Lys	440 Asn	Asn	Pro	Cys	Gly	445 Ala	Asp	Val
	Thr	Val	Ser	Gly	450 Gly	Lys	Ala	Thr	455 Phe	Thr	Ile	Pro	Ala	460 Lys	Thr	Ala
465	Val	Ala	Ile	Cys	465 Thr	Asp	Glu	Lys	470 Trp	Cys	Gly	Lys	Gly	475 Val	Asp	Pro
	Cys	Glu	Ser	Asp	480 Pro	Thr	Gly	Ser	485 Ala	Cys	Val	Cys	Lys	490 Gly	Glu	Thr
	Thr	Val	Asn	Gly	500 Val	Cys	Val	Ser	505 Trp	Cys	Asn	Ala	His	510 Ser	Ser	Asn
	Glu	Glu	Cys	Ala	515 Cys	Val	Leu	Asn	520 Pro	Asn	Asp	Ala	Glu	525 Cys	Gln	Ala
	Asp	Ile	Glu	Pro	530 Thr	Lys	Gly	Lys	535 Leu	Cys	Tyr	Val	Gly	540 Thr	Ser	Asn

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545 Lys Trp Thr Gln Glu 550 Pro Leu Thr Tyr Asn 555 Lys Thr Gly Phe 560
 Thr Leu Asn Val Glu 565 Leu Asp Gly Lys 570 Gly Asp Thr Ser Gly Ala Gln
 Arg Phe Lys Val Thr Asp Gly Cys 585 Ser Trp Gln Gly Thr Val Tyr Gly
 Ser Ser Gly Val Glu Gly Arg 600 Asp Val Asn Thr Ser Ala Thr Gly
 Asp Glu Pro Val Ser Leu Thr Gly Lys Tyr Val Leu Ser Ile Asn Asp
 Lys Thr Met Glu Tyr Thr Phe Ile Pro Ala Gly Ser Gly Asn Lys Pro
 Pro Val Ala Ser Phe Thr Pro Thr Val Lys Asp Leu Thr Val Ser Phe
 Val Asn Asn Ser Ser Asp Pro Glu Asn Asp Glu Leu Thr Tyr Ser Trp
 Asn Phe Gly Asn Gly Lys Thr Ser Ser Glu Lys Asn Pro Ser Val Thr
 Tyr Asp Lys Ala Gly Lys Tyr Thr Val Ser Leu Lys Val Thr Asp Thr
 Ala Asn Asn Thr Asp Thr Lys Thr Leu Glu Ile Asp Leu Thr Ser Pro
 Val Asn Gly Lys Tyr Ser Lys Val Ala Val Arg Gly Ser His Asp Asn
 Tyr Gly Thr Asn Leu Leu Thr Arg Asn Gly Ser Glu Trp Thr Gly Ile
 Phe Glu Phe Ser Lys Thr Thr Lys Phe Lys Leu Glu Ala Leu Pro Pro
 Ala Ala Asp Gln Cys Ile Phe Leu Gly Gly Asn Arg Gly Glu Ala Leu
 Thr Ala Ser Gly Gly Phe Ile Ser Leu Pro Ala Gly Arg Tyr Thr Ile
 Lys Phe Asn Glu Glu Ser Lys Val Leu Thr Ala Gly Asp Val Asp Cys
 Thr Gly

<210> 151
 <211> 2112
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 151
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 tattcagcca ccaagtggac taaactggaa gcacaggctg acgagatctg caactatttc 180
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 aagaatgtct caaactgggt ggatttcccg gccgagacct acaacggtgt aacctatcag 420
 atggtaagca ccgacatcgt ttcgaacgat gacggcggaa aaacagccac ttgggcaaat 480
 caaaacggct acagtctctc ctccaatgcc gacgaaggcg aaggctggga cggcatgcgc 540
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 gttgacgact taggctatac cggattccgc tacgatattg taaagggatt tgacggatcg 660
 catgtagccg actacaacac caatgccggc gtgcagttct ctgtcggcga atattgggac 720
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 ctggctcgga ttacacaacc gagcacctat gccgaatgga tgagcgggtg agcctacatc 1140
 ggacgtaccg taacaggtac gaacggcacc ttacgtgttc tgtgcggctc ttatcagtat 1200
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gaacggaaaag aagatactca tcagataaac agaggttccg aaccattctc ctattatgaa 2040
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ctcctcgtcg gg 2112

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<210> 152
 <211> 704
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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20     25     30
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35     40     45
Leu Glu Ala Gln Ala Asp Glu Ile Cys Asn Tyr Phe Ser Leu Val Trp
50     55     60
Val Pro Gln Ser Ala Tyr Thr Gly Ser Ser Thr Ser Met Gly Tyr Asp
65     70     75     80
Pro Leu Tyr Tyr Phe Asp Gln His Ser Ser Phe Gly Thr Glu Glu Gln
85     90     95
Leu Arg Ser Phe Ile Ser Thr Tyr Lys Gln Lys Gly Thr Gly Ile Ile
100    105    110
Ala Asp Val Val Val Asn His Arg Lys Asn Val Ser Asn Trp Val Asp
115    120    125
Phe Pro Ala Glu Thr Tyr Asn Gly Val Thr Tyr Gln Met Val Ser Thr
130    135    140
Asp Ile Val Ser Asn Asp Asp Gly Gly Lys Thr Ala Thr Trp Ala Asn
145    150    155    160
Gln Asn Gly Tyr Ser Leu Ser Ser Asn Ala Asp Glu Gly Glu Gly Trp
165    170    175
Asp Gly Met Arg Asp Leu Asp His Lys Ser Gln Asn Val Gln Lys Ser
180    185    190
Val Leu Ala Tyr Thr Lys Tyr Leu Val Asp Asp Leu Gly Tyr Thr Gly
195    200    205
Phe Arg Tyr Asp Met Val Lys Gly Phe Asp Gly Ser His Val Ala Asp
210    215    220
Tyr Asn Thr Asn Ala Gly Val Gln Phe Ser Val Gly Glu Tyr Trp Asp
225    230    235    240
Gly Thr Ala Ser Lys Val Tyr Ser Trp Ile Asn Ser Thr Lys Lys Ser
245    250    255
Asp Val Pro Gln Ser Ala Ala Phe Asp Phe Ala Phe Arg Tyr Thr Cys
260    265    270
Arg Asp Ala Val Asn Asn Lys Asn Trp Ala Asn Leu Lys Asn Thr Ser
275    280    285
Gly Ile Ser Asp Ala Asp Tyr Arg Arg Tyr Ser Val Thr Phe Val Glu
290    295    300
Asn His Asp Thr Glu Tyr Arg Ser Ala Thr Ala Ser Gln Asp Pro Ile
305    310    315    320
Lys Gly Asp Thr Val Ala Leu Asn Ala Trp Met Leu Ala Met Pro Gly
325    330    335
Thr Pro Cys Val Phe Leu Lys His Trp Thr Asp Cys Lys Glu Glu Ile

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Lys Asn Leu 340 Glu Ala Arg Arg 345 Leu Val Gly Ile His 350 Asn Gln Ser
Thr Tyr Ala 355 Glu Trp Met Ser 360 Gly Ala Ala Tyr Ile Gly Arg Thr Val
Thr Gly Thr Asn Gly Thr 375 Leu Arg Val Leu Cys 380 Gly Ser Tyr Gln Tyr
385 Asn Val Ala Ala Asn Tyr Ile Gln Ile Leu Ser Gly Lys Asn Tyr 400
405 Tyr Tyr Val Leu Asn Thr Leu Glu Ala Pro Trp Ile Gly Lys Gly Ser
420 Gly Ser Tyr Thr Glu Gly Glu Thr Val Thr Val Pro Leu Ile Ala Ile
435 Ser Ala Asp Ala Asn Ala Lys 440 Leu Val Tyr Thr Thr Asp Gly Thr Asp
450 Pro Thr Ala Thr Ser Thr Ala Val Thr Ser Gly Thr Glu Leu Thr Ile
465 Thr Ser Asp Ala Val Leu Lys Val Gly Leu Leu Ser Gly Gly Ile Val
485 Arg Asn Ile Gln Ser Arg Thr Phe Thr Phe Gln Ala Ala Asn Thr Ser
500 Glu Tyr Tyr Thr Ala Thr Met His Val Cys Asn Gln Ser Gly Ala Leu
515 Asn Pro Leu Phe Ala Tyr Val Trp Ala Gly Pro Asp Asn Glu Gln Ile
530 Asn Gly Asn Trp Pro Gly Thr Lys Leu Thr Ala Thr Ile Thr Glu Asn
545 Asn Leu Thr Trp Tyr Thr Gln Ser Phe Gln Ile Pro Lys Asn Val Asp
565 Tyr Val Val Asn Phe Val Phe Thr Thr Thr Gly Gly Gly Thr Gln Thr
580 Val Asp Val Thr Gly Met Lys Ala Asp Val Trp Tyr Ile Ile Asn Ser
595 Thr Lys Ser Gly Asn Lys Tyr Thr Val Thr Asp Val Thr Ser Gln Tyr
610 Ser Ser Leu Glu Ala Ile Phe Asp Glu Glu Asn Ser Gly Ser Phe Pro
625 Val Tyr Asp Leu Gln Gly Arg Arg Val Ser Glu Ile Arg Asn Arg Thr
645 Ile Ile Ser Ser Glu Arg Lys Glu Asp Thr His Gln Ile Asn Arg Gly
660 Ser Glu Pro Phe Ser Tyr Tyr Glu Asn Gln Thr Leu Ser Asn Leu Ser
675 Thr Ala Gly Phe Gly Gly Leu Val His His Gln Leu Leu Val Gly
690

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<210> 153
 <211> 1620
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 153
atgccttcaa ttaatgcaag cgattgcaaa aaaaagggag ataggagtat gaagaggaaa      60
aaatggactg cgtttagcact atctttacca ctagttatga gcttatcaac aaacatacaa      120
gcagaaacat tacataataa taagggtcaa aaggcgcaaa caggaaataa agacggaatt      180
ttttatgaac tgtatgttaa ttctttttat gatactgata gcaatggaca tgggtgattta      240
aaaggcgta caaagaaact tgattattta aatgatggaa atccaagaac aaataatgat      300
cttcaaataa acggtatctg gatgatgcct attaacacct ctcctagtta tcacaaatat      360
gatgtaacag attactataa tatcgcctcct cagtatggaa gtttacaaga tttccgtgaa      420
ctaacaacag aagcgcataa acgcaacgta aaggtagtaa tagatcttgt tattaatcat      480
acaagcagtg agcatccttg gtttgtcgat gcattaaaaa ataaaaacag taagtatcga      540
gattactata tttgggctga taaaaataca gacttaaatg aaaaaggccc atgggggtcaa      600
caagtatggc acaaagcgtc gaacggagag tattttctacg caacgttctg ggaagggatg      660
ccggacttaa actatgacaa ccctaaagta agagaagaaa tgattaaaat cgggaaattt      720
tggctcaaac aaggagctga tggctttcgt ctagatgcag ccatgcacat ctttaaaggg      780

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caaacacctg aaggagcaaa gaaaaatatt gaatggtgga atgaattccg cgacgcgatg 840
agagaaacga atccaaatac gtatctagtt ggtgaaatat gggatcaacc agaagtagtt 900
gctccgtatt atcaatcggt agattctaca ttttaacttcg acttagcata taaaatcggt 960
aattccgtta aaaatggtac tgatcaaggg gtagccgcgg cagctgttgc aacggatgag 1020
ttatataaaa catataatcc aaataaaatt gatggaacgt ttttaacgaa tcatgaccaa 1080
aatcgtgtaa tgagttagtt aaatggtgat gtaaacaag caaaatcagc agcctctatt 1140
ctgttgacac tccctggtaa tccgttcatt tattatggcg aagaaatcgg catgacaggc 1200
caaaaaccag atgagttgat tcgtgagcct ttccgttggg atgaagatga taaagaaggt 1260
caaacgagct gggagactcc agtatataac attgatcata atggtgtttc agttgaagca 1320
caagataaac aaaaagcttc tcttctaagc cattatcgta aaatgattcg tgttcgtcag 1380
caacacgatg aacttgtcaa aggtaattta gaacctattt ctgtcaataa ttcacagggt 1440
gttgccata atcgtacgta taaaaataaa tcaattcaag tgtaccataa tatttcagac 1500
aagccggtta cattaactgt ttcaaacaaa ggaaaactga ttttttctag tgaattagga 1560
gcaaaaaagg aaaaatcaac attagtaatt ccagcgaata cgacagtgtc agtaaagtaa 1620

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<210> 154
 <211> 539
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 154
Met Pro Ser Ile Asn Ala Ser Asp Cys Lys Lys Lys Gly Asp Arg Ser
1      5      10      15
Met Lys Arg Lys Lys Trp Thr Ala Leu Ala Leu Ser Leu Pro Leu Val
20     25     30
Met Ser Leu Ser Thr Asn Ile Gln Ala Glu Thr Leu His Asn Asn Lys
35     40     45
Gly Gln Lys Ala Gln Thr Gly Asn Lys Asp Gly Ile Phe Tyr Glu Leu
50     55     60
Tyr Val Asn Ser Phe Tyr Asp Thr Asp Ser Asn Gly His Gly Asp Leu
65     70     75     80
Lys Gly Val Thr Lys Lys Leu Asp Tyr Leu Asn Asp Gly Asn Pro Arg
85     90     95
Thr Asn Asn Asp Leu Gln Ile Asn Gly Ile Trp Met Met Pro Ile Asn
100    105    110
Thr Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile
115    120    125
Asp Pro Gln Tyr Gly Ser Leu Gln Asp Phe Arg Glu Leu Thr Thr Glu
130    135    140
Ala His Lys Arg Asn Val Lys Val Val Ile Asp Leu Val Ile Asn His
145    150    155    160
Thr Ser Ser Glu His Pro Trp Phe Val Asp Ala Leu Lys Asn Lys Asn
165    170    175
Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu
180    185    190
Asn Glu Lys Gly Pro Trp Gly Gln Val Trp His Lys Ala Ser Asn
195    200    205
Gly Glu Tyr Phe Tyr Ala Thr Phe Trp Glu Gly Met Pro Asp Leu Asn
210    215    220
Tyr Asp Asn Pro Lys Val Arg Glu Glu Met Ile Lys Ile Gly Lys Phe
225    230    235    240
Trp Leu Lys Gln Gly Ala Asp Gly Phe Arg Leu Asp Ala Ala Met His
245    250    255
Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Glu Trp
260    265    270
Trp Asn Glu Phe Arg Asp Ala Met Arg Glu Thr Asn Pro Asn Thr Tyr
275    280    285
Leu Val Gly Glu Ile Trp Asp Gln Pro Glu Val Val Ala Pro Tyr Tyr
290    295    300
Gln Ser Leu Asp Ser Thr Phe Asn Phe Asp Leu Ala Tyr Lys Ile Val
305    310    315    320
Asn Ser Val Lys Asn Gly Thr Asp Gln Gly Val Ala Ala Ala Val
325    330    335
Ala Thr Asp Glu Leu Tyr Lys Thr Tyr Asn Pro Asn Lys Ile Asp Gly
340    345    350

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Thr	Phe	Leu	Thr	Asn	His	Asp	Gln	Asn	Arg	Val	Met	Ser	Glu	Leu	Asn
		355					360					365			
Gly	Asp	Val	Asn	Lys	Ala	Lys	Ser	Ala	Ala	Ser	Ile	Leu	Leu	Thr	Leu
	370					375					380				
Pro	Gly	Asn	Pro	Phe	Ile	Tyr	Tyr	Gly	Glu	Glu	Ile	Gly	Met	Thr	Gly
	385				390					395					400
Gln	Lys	Pro	Asp	Glu	Leu	Ile	Arg	Glu	Pro	Phe	Arg	Trp	Tyr	Glu	Asp
			405					410						415	
Asp	Lys	Glu	Gly	Gln	Thr	Ser	Trp	Glu	Thr	Pro	Val	Tyr	Asn	Ile	Asp
			420					425					430		
His	Asn	Gly	Val	Ser	Val	Glu	Ala	Gln	Asp	Lys	Gln	Lys	Ala	Ser	Leu
		435					440					445			
Leu	Ser	His	Tyr	Arg	Lys	Met	Ile	Arg	Val	Arg	Gln	Gln	His	Asp	Glu
	450					455					460				
Leu	Val	Lys	Gly	Asn	Leu	Glu	Pro	Ile	Ser	Val	Asn	Asn	Ser	Gln	Val
	465				470					475					480
Val	Ala	Tyr	Asn	Arg	Thr	Tyr	Lys	Asn	Lys	Ser	Ile	Gln	Val	Tyr	His
			485					490						495	
Asn	Ile	Ser	Asp	Lys	Pro	Val	Thr	Leu	Thr	Val	Ser	Asn	Lys	Gly	Lys
			500					505					510		
Leu	Ile	Phe	Ser	Ser	Glu	Leu	Gly	Ala	Lys	Lys	Glu	Lys	Ser	Thr	Leu
	515						520					525			
Val	Ile	Pro	Ala	Asn	Thr	Thr	Val	Leu	Val	Lys					
	530					535									

<210> 155
 <211> 1773
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 155

gtgtcaagaa	tgtttgcaaa	acgattcaaa	acctctttac	tgccggttatt	cgctggattt	60
ttattgctgt	ttcatttggg	tctggcagga	ccaacggctg	cgaatgctga	aacggctaac	120
aaatcaaatg	agcttacagc	accgtcgatc	aaaagcggaa	ccattcttca	tgcttggaat	180
tggtcgttca	atacgttaaa	acacaatatg	aaggatattc	atgatgcagg	atatacagcg	240
attcagacgt	ctccgattaa	ccaagtcaag	gaagggaacc	aaggaaataa	aaacatgtcg	300
aactgggtact	ggctctatca	gccgacatcg	taccaaattg	gcaaccgtta	cttaggtact	360
gaacaagaat	ttaaagaaat	gtgtgcagcc	gctgaagaat	atggcataaa	ggttattggt	420
gacgcggtca	tcaatcatac	caccagtgcg	tatgccgcga	tttccaatga	gattaagagt	480
attccaaact	ggacacatgg	aaacacacaa	attaaaaact	ggctctgatcg	atgggatgtc	540
acgcagaaatg	cattgctcgg	gctgtatgac	tggaatacac	aaaatacac	agtacagtc	600
tatttgaaac	ggttcttaga	aagagcattg	aatgacgggg	cagacgggtt	tcgatttgat	660
gccgccaaac	atatagagct	tccggatgat	ggcagttacg	gcagtcaatt	ttggccgaat	720
atcacaataa	catctgcaga	gttccaatac	ggagaaatcc	tgcaggatag	tgcttcaaga	780
gatgcttcat	atgcgaatta	tatgaatgtg	acagcgtcta	actatgggca	ttccataagg	840
tccgctttaa	agaatcgtaa	tctgggcgtg	tcgaatatct	cccactatgc	atcagatgtg	900
tctgcggaca	agctagtgcg	atgggtagaa	tcgcatgata	cgtatgccaa	tgatgatgaa	960
gagtcgacat	ggatgagcga	tgatgatata	cgtttaggct	gggcgggtgat	agcttctcgt	1020
tcaggcagta	cgctctttt	cttttccaga	cctgagggag	gcggaaatgg	tgtgagattc	1080
ccggggaaaa	gccaaatagg	cgatcgcggg	agtgccttat	ttgaagatca	ggctatcact	1140
gcgggtcaata	gatttcacaa	tgtgatggct	ggacagcctg	aggaactctc	gaacccaaat	1200
ggaaacaacc	agatatattat	gaatcagcgc	ggctcacatg	gcgttggtgt	ggcaaatgca	1260
ggttcatcct	ctgtttctat	caatacgcca	acaaaattgc	ctgatggcag	gtatgataat	1320
aaagctgggg	caggttcatt	tcaagtaaat	gacggtaaac	tgacaggcac	gatcaatgcc	1380
aggctctgtg	ctgtgcttta	tcctgatgat	attgcaaaag	cgcctcatgt	tttccttgag	1440
aattacaaaa	caggtgtaac	acattctttc	aatgatcaac	tgacgattac	actgcgtgca	1500
gatgcgaata	caacaaaagc	cgtttatcaa	atcaataatg	gaccagagac	ggcgtttaag	1560
gatggagatc	aattcacaat	cggaaaagga	gatccatttg	gcaaaacata	caccatcatg	1620
ttaaaaggaa	cgaacagtga	tggtgtaacg	aggaccgagg	aatacagttt	tgttaaaaga	1680
gatccagctt	cggccaaaac	catcggctat	caaaatccga	atcattggag	ccaggtaaat	1740
gcttatatct	ataaacatga	tgggggccgg	gca			1773

<210> 156
 <211> 591
 <212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 156

Val 1	Ser	Arg	Met	Phe 5	Ala	Lys	Arg	Phe	Lys 10	Thr	Ser	Leu	Leu	Pro 15	Leu
Phe	Ala	Gly	Phe 20	Leu	Leu	Leu	Phe	His 25	Leu	Val	Leu	Ala	Gly 30	Pro	Thr
Ala	Ala	Asn 35	Ala	Glu	Thr	Ala	Asn 40	Lys	Ser	Asn	Glu	Leu	Thr	Ala	Pro
Ser	Ile 50	Lys	Ser	Gly	Thr	Ile 55	Leu	His	Ala	Trp	Asn 60	Trp	Ser	Phe	Asn
Thr 65	Leu	Lys	His	Asn 70	Met	Lys	Asp	Ile	His 75	Asp	Ala	Gly	Tyr	Thr	Ala 80
Ile	Gln	Thr	Ser	Pro 85	Ile	Asn	Gln	Val	Lys 90	Glu	Gly	Asn	Gln	Gly 95	Asn
Lys	Asn	Met	Ser 100	Asn	Trp	Tyr	Trp	Leu 105	Tyr	Gln	Pro	Thr	Ser	Tyr	Gln
Ile	Gly	Asn 115	Arg	Tyr	Leu	Gly	Thr 120	Glu	Gln	Glu	Phe	Lys 125	Glu	Met	Cys
Ala 130	Ala	Ala	Glu	Glu	Tyr	Gly 135	Ile	Lys	Val	Ile	Val 140	Asp	Ala	Val	Ile
Asn 145	His	Thr	Thr	Ser	Asp 150	Tyr	Ala	Ala	Ile	Ser 155	Asn	Glu	Ile	Lys	Ser 160
Ile	Pro	Asn	Trp	Thr 165	His	Gly	Asn	Thr	Gln 170	Ile	Lys	Asn	Trp	Ser 175	Asp
Arg	Trp	Asp	Val 180	Thr	Gln	Asn	Ala	Leu 185	Leu	Gly	Leu	Tyr	Asp 190	Trp	Asn
Thr	Gln	Asn 195	Thr	Gln	Val	Gln	Ser 200	Tyr	Leu	Lys	Arg	Phe 205	Leu	Glu	Arg
Ala 210	Leu	Asn	Asp	Gly	Ala	Asp 215	Gly	Phe	Arg	Phe	Asp 220	Ala	Ala	Lys	His
Ile 225	Glu	Leu	Pro	Asp 230	Asp	Gly	Ser	Tyr	Gly	Ser 235	Gln	Phe	Trp	Pro	Asn 240
Ile	Thr	Asn	Thr	Ser 245	Ala	Glu	Phe	Gln	Tyr 250	Gly	Glu	Ile	Leu	Gln 255	Asp
Ser	Ala	Ser	Arg 260	Asp	Ala	Ser	Tyr	Ala 265	Asn	Tyr	Met	Asn	Val 270	Thr	Ala
Ser	Asn	Tyr 275	Gly	His	Ser	Ile	Arg 280	Ser	Ala	Leu	Lys	Asn	Arg	Asn	Leu
Gly 290	Val	Ser	Asn	Ile	Ser	His 295	Tyr	Ala	Ser	Asp	Val 300	Ser	Ala	Asp	Lys
Leu 305	Val	Thr	Trp	Val	Glu 310	Ser	His	Asp	Thr	Tyr 315	Ala	Asn	Asp	Asp	Glu 320
Glu	Ser	Thr	Trp	Met 325	Ser	Asp	Asp	Asp	Ile 330	Arg	Leu	Gly	Trp	Ala 335	Val
Ile	Ala	Ser	Arg 340	Ser	Gly	Ser	Thr	Pro 345	Leu	Phe	Phe	Ser	Arg 350	Pro	Glu
Gly	Gly	Gly 355	Asn	Gly	Val	Arg	Phe 360	Pro	Gly	Lys	Ser	Gln 365	Ile	Gly	Asp
Arg 370	Gly	Ser	Ala	Leu	Phe	Glu 375	Asp	Gln	Ala	Ile	Thr 380	Ala	Val	Asn	Arg
Phe 385	His	Asn	Val	Met	Ala 390	Gly	Gln	Pro	Glu	Glu 395	Leu	Ser	Asn	Pro	Asn 400
Gly	Asn	Asn	Gln 405	Ile	Phe	Met	Asn	Gln	Arg 410	Gly	Ser	His	Gly	Val 415	Val
Leu	Ala	Asn	Ala 420	Gly	Ser	Ser	Ser	Val 425	Ser	Ile	Asn	Thr	Pro 430	Thr	Lys
Leu	Pro	Asp 435	Gly	Arg	Tyr	Asp	Asn 440	Lys	Ala	Gly	Ala	Gly 445	Ser	Phe	Gln
Val 450	Asn	Asp	Gly	Lys	Leu	Thr 455	Gly	Thr	Ile	Asn	Ala	Arg	Ser	Val	Ala
Val 465	Leu	Tyr	Pro	Asp	Asp 470	Ile	Ala	Lys	Ala	Pro 475	His	Val	Phe	Leu	Glu 480
Asn	Tyr	Lys	Thr	Gly 485	Val	Thr	His	Ser	Phe 490	Asn	Asp	Gln	Leu	Thr 495	Ile

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Thr	Leu	Arg	Ala	Asp	Ala	Asn	Thr	Thr	Lys	Ala	Val	Tyr	Gln	Ile	Asn
			500					505					510		
Asn	Gly	Pro	Glu	Thr	Ala	Phe	Lys	Asp	Gly	Asp	Gln	Phe	Thr	Ile	Gly
		515					520					525			
Lys	Gly	Asp	Pro	Phe	Gly	Lys	Thr	Tyr	Thr	Ile	Met	Leu	Lys	Gly	Thr
	530				535						540				
Asn	Ser	Asp	Gly	Val	Thr	Arg	Thr	Glu	Glu	Tyr	Ser	Phe	Val	Lys	Arg
545					550					555					560
Asp	Pro	Ala	Ser	Ala	Lys	Thr	Ile	Gly	Tyr	Gln	Asn	Pro	Asn	His	Trp
				565					570					575	
Ser	Gln	Val	Asn	Ala	Tyr	Ile	Tyr	Lys	His	Asp	Gly	Gly	Arg	Ala	
			580					585					590		

<210> 157
 <211> 1596
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 157

atgcaaacga	ttgcaaaaaa	aggggatgaa	acgatgaaag	ggaaaaaatg	gacagcatta	60
gctctaacac	tgccgctggc	tgctagctta	tcaacaggcg	ttcacgccga	aaccgtacat	120
aaaggtaaag	ctccaacagc	agataaaaaa	ggtgtctttt	atgaggtgta	tgtaaactct	180
ttttacgatg	caaataaaga	tggacatggt	gattttaaag	gtcttacaca	aaagctggat	240
tatttgaatg	acggcaattc	tcataccaaa	aatgatcttc	aagtaaacgg	aatttggatg	300
atgccggtaa	acccttctcc	tagctatcat	aaatatgatg	taacggacta	ttataacatt	360
gatccgcagt	acggaaatct	gcaagatttt	cgcaagctga	tgaaagaagc	agataaacga	420
gacgtaaagg	ttattatgga	cctcgtttgt	aatcatacaa	gcagtgaaca	tccttggttt	480
caagctgcat	taaaagataa	aaacagcaag	tacagagatt	actatatatt	ggccgataaa	540
aatactgatt	taaatgaaaa	aggatcttgg	gggcagcaag	tatggcataa	agctccaaac	600
ggagagtatt	tttatgggtac	gttttgggaa	ggaatgcctg	acttaaatta	cgataatccc	660
gaagtaagaa	aagaaatgat	taacgtcggg	aaattttggc	taaagcaagg	cgttgacggg	720
ttccgcttag	atgctgcgct	tcataattttt	aaagggtcaa	cacctgaagg	cgctaagaaa	780
aatatcgtgt	ggtggaatga	gttttagagat	gcaatgaaaa	aagaaaaccc	taacgtatat	840
ctaaccgggtg	aagtatggga	tcaaccggaa	gtagttagctc	cttactatca	atcgcttgat	900
cttttattta	actttgattt	agcaggaaag	attgtaaact	ctgtaaaatc	aggaaatgat	960
caaggaatcg	cgactgcagc	agccgcaact	gatgagctgt	tcaaatcata	caatccaaat	1020
aaaattgacg	gcattttctt	aaccaaccat	gaccaaatac	gcgtcatgag	tgagctaagc	1080
ggcgatgtga	ataaagcaaa	gtcagctgcc	tctatcttac	ttacgcttcc	tggcaaccgc	1140
tatatttatt	acggtgaaga	aattggaatg	accggtgaaa	agcctgatga	gttaatccgt	1200
gaaccgttcc	gctggtacga	aggcaatgga	cttggacaaa	ccagctggga	aacatccgta	1260
tacaacaaaag	gcggcaatgg	tgtgtcagta	gagacacaaa	caaaacaaaa	ggattctttg	1320
ttaaatcatt	accgtgaaat	gattcgcgtg	cgtcagcagc	atgaagagtt	agtaaaagga	1380
acccttcaat	ctatttcagt	agacagtaaa	gaagtcgttg	cctatagccg	cacgtataaa	1440
ggcaaatcga	ttagcgtgta	tcataaatatt	tcaaatcaac	cggtaaaagt	atctgtaaca	1500
gcgaaaggta	aattgatttt	tgctagttaa	aaagggtgaa	aaaaagtcaa	aaatcagctt	1560
gtggttccag	ctaatacaac	ggttttaata	aaataa			1596

<210> 158
 <211> 531
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 158

Met	Gln	Thr	Ile	Ala	Lys	Lys	Gly	Asp	Glu	Thr	Met	Lys	Gly	Lys	Lys
1				5					10					15	
Trp	Thr	Ala	Leu	Ala	Leu	Thr	Leu	Pro	Leu	Ala	Ala	Ser	Leu	Ser	Thr
			20					25					30		
Gly	Val	His	Ala	Glu	Thr	Val	His	Lys	Gly	Lys	Ala	Pro	Thr	Ala	Asp
		35					40					45			
Lys	Asn	Gly	Val	Phe	Tyr	Glu	Val	Tyr	Val	Asn	Ser	Phe	Tyr	Asp	Ala
	50					55					60				
Asn	Lys	Asp	Gly	His	Gly	Asp	Leu	Lys	Gly	Leu	Thr	Gln	Lys	Leu	Asp

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65	Tyr	Leu	Asn	Asp	Gly	70	Asn	Ser	His	Thr	Lys	75	Asn	Asp	Leu	Gln	Val	80	Asn
					85						90						95		
	Gly	Ile	Trp	Met	Met	Pro	Val	Asn	Pro	105	Ser	Pro	Ser	Tyr	His	110	Lys	Tyr	
	Asp	Val	Thr	Asp	Tyr	Tyr	Asn	Ile	Asp	120	Pro	Gln	Tyr	Gly	Asn	125	Leu	Gln	
	Asp	Phe	Arg	Lys	Leu	Met	Lys	Glu	Ala	135	Asp	Lys	Arg	Asp	Val	140	Lys	Val	
	Ile	Met	Asp	Leu	Val	Val	Asn	His	Thr	150	Ser	Ser	Glu	His	Pro	155	Trp	Phe	
	Gln	Ala	Ala	Leu	Lys	Asp	Lys	Asn	Ser	165	Lys	Tyr	Arg	Asp	Tyr	170	Tyr	Ile	
	Trp	Ala	Asp	Lys	Asn	Thr	Asp	Leu	Asn	180	Glu	Lys	Gly	Ser	Trp	185	Gly	Gln	
	Gln	Val	Trp	His	Lys	Ala	Pro	Asn	Gly	195	Glu	Tyr	Phe	Tyr	Gly	200	Thr	Phe	
	Trp	Glu	Gly	Met	Pro	Asp	Leu	Asn	Tyr	210	Asp	Asn	Pro	Glu	Val	215	Arg	Lys	
	Glu	Met	Ile	Asn	Val	Gly	Lys	Phe	Trp	225	Leu	Lys	Gln	Gly	Val	230	Asp	Gly	
	Phe	Arg	Leu	Asp	Ala	Ala	Leu	His	Ile	245	Phe	Lys	Gly	Gln	Thr	250	Pro	Glu	
	Gly	Ala	Lys	Lys	Asn	Ile	Val	Trp	Trp	260	Asn	Glu	Phe	Arg	Asp	265	Ala	Met	
	Lys	Lys	Glu	Asn	Pro	Asn	Val	Tyr	Leu	275	Thr	Gly	Glu	Val	Trp	280	Asp	Gln	
	Pro	Glu	Val	Val	Ala	Pro	Tyr	Tyr	Gln	290	Ser	Leu	Asp	Ser	Leu	300	Phe	Asn	
	Phe	Asp	Leu	Ala	Gly	Lys	Ile	Val	Asn	310	Ser	Val	Lys	Ser	Gly	315	Asn	Asp	
	Gln	Gly	Ile	Ala	Thr	Ala	Ala	Ala	Ala	325	Thr	Asp	Glu	Leu	Phe	330	Lys	Ser	
	Tyr	Asn	Pro	Asn	Lys	Ile	Asp	Gly	Ile	340	Phe	Leu	Thr	Asn	His	345	Asp	Gln	
	Asn	Arg	Val	Met	Ser	Glu	Leu	Ser	Gly	355	Asp	Val	Asn	Lys	Ala	360	Lys	Ser	
	Ala	Ala	Ser	Ile	Leu	Leu	Thr	Leu	Pro	370	Gly	Asn	Pro	Tyr	Ile	375	Tyr	Tyr	
	Gly	Glu	Glu	Ile	Gly	Met	Thr	Gly	Glu	385	Lys	Pro	Asp	Glu	Leu	390	Ile	Arg	
	Glu	Pro	Phe	Arg	Trp	Tyr	Glu	Gly	Asn	405	Gly	Leu	Gly	Gln	Thr	410	Ser	Trp	
	Glu	Thr	Ser	Val	Tyr	Asn	Lys	Gly	Gly	420	Asn	Gly	Val	Ser	Val	425	Glu	Thr	
	Gln	Thr	Lys	Gln	Lys	Asp	Ser	Leu	Leu	435	Asn	His	Tyr	Arg	Glu	440	Met	Ile	
	Arg	Val	Arg	Gln	Gln	His	Glu	Glu	Leu	450	Val	Lys	Gly	Thr	Leu	455	Gln	Ser	
	Ile	Ser	Val	Asp	Ser	Lys	Glu	Val	Val	465	Ala	Tyr	Ser	Arg	Thr	470	Tyr	Lys	
	Gly	Lys	Ser	Ile	Ser	Val	Tyr	His	Asn	485	Ile	Ser	Asn	Gln	Pro	490	Val	Lys	
	Val	Ser	Val	Thr	Ala	Lys	Gly	Lys	Leu	500	Ile	Phe	Ala	Ser	Glu	505	Lys	Gly	
	Ala	Lys	Lys	Val	Lys	Asn	Gln	Leu	Val	515	Val	Pro	Ala	Asn	Thr	520	Thr	Val	
	Leu	Ile	Lys							530									

<210> 159

<211> 1587

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

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<400> 159
atgcaaaaaa aaggggatga aacgatgaaa gggaaaaaat ggacagcttt agctctaaca      60
ctgccgctgg ctgctagctt atcaacaggc gttcacgccg aaaccgtaca taaaggtaaa      120
tctccaacag cagataaaaa cgggtgatttt tatgaggtgt atgtaaactc tttttacgat      180
gcaaataaag atggacatgg tgatttaaaa ggtcttacac aaaagttgga ttattttaat      240
gatggcaatt ctcatacaaa gaatgatctt caagtaaacg ggatttggtat gatgccggtc      300
aacccttctc ccagctatca taaatatgat gtaacggact attataatat tgatccgcag      360
tatggaaatc tgcaagattt tcgcaaaactg atgaaagaag cagataaacg agatgtaaaa      420
gtcattatgg acctcgttgt gaatcatacg agcagtgaac acccttggtt tcaagctgca      480
ttaaagataa aaaacagcaa gtacagagat tactatatct gggctgataa aaataccgac      540
ttgaatgaaa aaggatcttg gggacagcaa gtatggcata aagctccaaa cggagagtat      600
ttttacggaa cgttttggga aggaatgccg gacttaaatt acgataatcc tgaagtaaga      660
aaagaaatga ttaacgtagg aaagttttgg ctaaagcaag gagttgatgg gttccgtcta      720
gatgtgcgcg ttcatatttt taaaggccaa acacctgaag gcgctaagaa aaatctcctg      780
tggtggaaatg aatttagaga tgcaatgaaa aaggaaaacc ctaacgtata tctaacgggt      840
gaagtatggg atcaaccgga agtagtagct cttactatc aatcgcttga ttctttattt      900
aactttgatt tagcaggaaa gattgtaaac tctgtaaaat caggaaatga tcaaggaatc      960
gcgactgcag cagcggcaac ggatgaactg ttcaaatacat acaatccaaa taaaattgac     1020
ggtattttct taaccaacca tgaccaaata cgcgctcatga gtgagctaaa cggcgatgtg     1080
aataaaagcaa agtcagctgc ctctatctta cttacgcttc ctggcaaccg gtatatttat     1140
tacggtgaag aaatcggcat accgggtgaa aagcctgatg agttaatccg tgaaccgttc     1200
ccctgggtacg aaggaaacgg acttggaaca accagctggg aaacacctgt atataacaaa     1260
ggcggcaacg gcgtgtctgt agaagcacaa acaaaacaaa aggactcttt gttaaatcat     1320
taccgtgaaa tgattcgcgt gcgtcagcag cacgaagagt tagtaaaagg aacgcttcaa     1380
tctatttcag tagacagtaa agaagtcggt gcctatagcc gtacgtataa aggcaaatcg     1440
attagcgtgt atcataatat ttcaaatacaa ccggtaaaag tatctgtagc agcaaaaggt     1500
aaattgattt ttgctagtga aaaaggtgct aagaaagtca aaaatcagct tgtgattccg     1560
gcgaatacaa cggttttaat aaaataa                                     1587

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<210> 160

<211> 528

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 160

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Met Gln Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala
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Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His
          20          25          30
Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr Ala Asp Lys Asn Gly
          35          40          45
Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp
          50          55          60
Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn
 65          70          75          80
Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp
          85          90          95
Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr
          100          105          110
Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg
          115          120          125
Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val Ile Met Asp
          130          135          140
Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala
          145          150          155          160
Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp
          165          170          175
Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp
          180          185          190
His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly
          195          200          205
Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile
          210          215          220
Asn Val Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu
          225          230          235          240

```

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Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys
 245 250 255
 Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu
 260 270
 Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val
 275 280 285
 Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu
 290 300
 Ala Gly Lys Ile Val Asn Ser Val Lys Ser Gly Asn Asp Gln Gly Ile
 305 310 315 320
 Ala Thr Ala Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro
 325 330 335
 Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val
 340 345 350
 Met Ser Glu Leu Asn Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser
 355 360 365
 Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu
 370 375 380
 Ile Gly Met Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe
 385 390 395 400
 Pro Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr Ser Trp Glu Thr Pro
 405 410 415
 Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys
 420 425 430
 Gln Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg
 435 440 445
 Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser Ile Ser Val
 450 455 460
 Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser
 465 470 475 480
 Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val
 485 490 495
 Ala Ala Lys Gly Lys Leu Ile Phe Ala Ser Glu Lys Gly Ala Lys Lys
 500 505 510
 Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys
 515 520 525

<210> 161
 <211> 1623
 <212> DNA
 <213> Bacterial

<400> 161
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 atgaaagggga aaaaatggac agcttttagct ctaacactgc cgctggctgc tagcttatca 120
 acagggtgttc acgccgaaac cgtacataaa ggtaaagctc caacagcaga taaaaacggt 180
 gtcttttatg aggtatatgt aaactctttt tacgatgcaa ataaagatgg acatgggtgat 240
 ttaaaaggcc ttacacaaaa gttggactat ttaaatgacg gaaattctca tacaagaat 300
 gatcttcaag taaacgggat ttggatgatg ccggtcaacc cttctcctag ctatcataaa 360
 tatgatgtaa cggactatta taatattgat ccgcagtatg gaaatctgca agattttcgc 420
 aaacttatga aagaagcaga taaacgagac gtaaaagtca ttatggacct tgttgtgaat 480
 catacgagca gtgaacaccc ttggtttcaa gctgcgttga aagataaaaa cagcaagtac 540
 agagattact atatttgggc tgataaaaat actgacttga atgaaaaagg atcttgggga 600
 caacaagtat ggcataaagc tccaaacgga gagtattttt acggaacggt ctgggaagga 660
 atgcctgact taaattacga taacctgaa gtaagaaaag aaatgattaa cgtcggaaag 720
 ttttggtctaa aacaaggcgt tgacggcttc cgcttagatg ctgcccttca tatttttaaa 780
 ggtcaaacgc ctgaaggcgc taagaaaaac attctatggt ggaatgagtt tagagatgcg 840
 atgaaaaaag aaaacccgaa cgtatatcta acgggtgaag tgtgggacca gccagaagta 900
 gtagccctt actatcaatc acctgattct ctatttaatt ttgatttagc agggaaaaatt 960
 gtcagctctg taaaagcagg aaatgatcaa ggaatcgcca ctgcagcagc ggcaactgat 1020
 gagctgttca aatcatacaa tccaaataaa attgacggca ttttcttaac caaccatgac 1080
 caaaatcgcg tcatgagtga gttaagcggc gatgtgaata aagcaaaatc agccgcctct 1140
 atcttactta cgcttcctgg aaatccgtat atttattacg gtgaagaaat tggcatgaca 1200
 ggtgaaaagc ctgatgaatt aatccgtgaa ccgttccgct ggtacgaagg caacggaatt 1260
 ggacaaaacta gctgggaaac acctgtatat aacaaaggcg gtaacggcgt gtctgtagaa 1320
 gcacaaaaca aacaaaagga ttccttgtaa aatcattacc gtgaaatgat tcgtgtgcgc 1380
 cagcagcacg aagagttagt aaaaggaacg cttcaatcca tttcagtaga cagtaaagaa 1440
 gtcgttgcct atagccgcac gtacaaaggc aaatcgatta gcgtgtatca taatatttca 1500

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aatcaacctg taaaagtatc tgtagcagcg aaaggtaact tgatttttgc tagtgaaaaa 1560
 ggtgctaaga aagtcaaaaa tcagcttgtg attccggcga atgcgacggt ttttaataaaa 1620
 taa 1623

<210> 162
 <211> 540
 <212> PRT
 <213> Bacterial

<400> 162
 Val Asp Pro Lys Asn Cys Ser Gln Phe Met Gln Thr Ile Ala Lys Lys
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 Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
 20 25 30
 Leu Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val
 35 40 45
 His Lys Gly Lys Ala Pro Thr Ala Asp Lys Asn Gly Val Phe Tyr Glu
 50 55 60
 Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp
 65 70 75 80
 Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser
 85 90 95
 His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val
 100 105 110
 Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn
 115 120 125
 Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys
 130 135 140
 Glu Ala Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn
 145 150 155 160
 His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys
 165 170 175
 Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 180 185 190
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro
 195 200 205
 Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly Met Pro Asp Leu
 210 215 220
 Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly Lys
 225 230 235 240
 Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu
 245 250 255
 His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys Lys Asn Ile Leu
 260 265 270
 Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu Asn Pro Asn Val
 275 280 285
 Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr
 290 295 300
 Tyr Gln Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile
 305 310 315 320
 Val Ser Ser Val Lys Ala Gly Asn Asp Gln Gly Ile Ala Thr Ala Ala
 325 330 335
 Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp
 340 345 350
 Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu
 355 360 365
 Ser Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile Leu Leu Thr
 370 375 380
 Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
 385 390 395 400
 Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu
 405 410 415
 Gly Asn Gly Ile Gly Gln Thr Ser Trp Glu Thr Pro Val Tyr Asn Lys
 420 425 430
 Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys Asp Ser
 435 440 445
 Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu
 450 455 460

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Glu	Leu	Val	Lys	Gly	Thr	Leu	Gln	Ser	Ile	Ser	Val	Asp	Ser	Lys	Glu
465					470				475						480
Val	Val	Ala	Tyr	Ser	Arg	Thr	Tyr	Lys	Gly	Lys	Ser	Ile	Ser	Val	Tyr
				485					490					495	
His	Asn	Ile	Ser	Asn	Gln	Pro	Val	Lys	Val	Ser	Val	Ala	Ala	Lys	Gly
			500					505					510		
Asn	Leu	Ile	Phe	Ala	Ser	Glu	Lys	Gly	Ala	Lys	Lys	Val	Lys	Asn	Gln
		515					520					525			
Leu	Val	Ile	Pro	Ala	Asn	Ala	Thr	Val	Leu	Ile	Lys				
	530					535					540				

<210> 163
 <211> 1422
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 163
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 gatgtcgggt atggggcgta cgatttatac gatctcggcg aattcaacca aaaagggacg 180
 acccggacga agtacgggac gaaagcgcag ctccagaccg ccatctcgaa cttgcgcggt 240
 aaagggatcg gtgtgtacgg cgacgtcgtc atgaatcaca agggcggggc cgattatacc 300
 gaatccgttc aggcgatcga ggtcaatccg tcgaaccgga accaagaaac gtccggtgag 360
 tatggcatct cggcctggac tgggttcaac ttcgcggggc gcaacaatac atactcgccg 420
 ttcaaatggc gctggtacca ttttgacggg accgattggg atcagtcacg cagcttgagc 480
 cgcatctata agttcaagag cacaggcaag gcgtgggaca cggacgtgtc gaacgagaac 540
 ggcaactatg attatcttat gtatgccgac gtcgatttcg agcatcccga ggtccgcca 600
 gagatgaaga actggggcaa atggtacgcc gactcgctcg ggctcgacgg tttccggttg 660
 gatgcggtca aacatatcag ccactcgta tgaaggagt gggtgacgag cgtgcgccag 720
 acgaccggga aagagatggt cacggtcgcc gagtattgga agaacgatct cggtgccatc 780
 aacgactatc tgtataagac gggctacacg cactccgtct tcgatgtgcc gctccattat 840
 aacttccaag cggccggtaa cggcggcggg tattacgata tgcgcaacat cttgaaaggc 900
 accgtcaccg aacagcatcc gtcgctgtcc gtgacgattg tcgataacca cgactcacag 960
 ccggggcagt cgctcgagtc gacggtcgcc aactggttca aaccgctcgc ctacgcgacg 1020
 atcatgacgc gcggtcaggg ttatccggcc ctcttctatg gagactatta tggcacgaaa 1080
 gggacgacga accgcgaaat cccgaacatg tcgggcacgc tccaaccgat tttgaaggca 1140
 cgaaaagact tcgcctacgg gacgcagcat gactacctcg atcatcagga cgtcatcggc 1200
 tggacacgtg aagggtgtgac cgaccgtgcc aaatcgggtc tcgacgacgat tctatcggac 1260
 ggtccgggag gctcgaagtg gatgtacgtc ggcaaacaga acgccggcga ggtatggaaa 1320
 gacatgacga acaacaacgc ccgtctcgtc acgatcaatg ctgacggctg gggtcagttc 1380
 ttcgtcaacg gaggtcgggt ctcgatttat acgcaacaat aa 1422

<210> 164
 <211> 473
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 164
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 Leu Ala Ala Leu Glu Arg His Trp Val Thr Thr Val Trp Ile Pro Pro
 20 25 30
 Ala Tyr Lys Gly Thr Ser Gln Asn Asp Val Gly Tyr Gly Ala Tyr Asp
 35 40 45
 Leu Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Thr Arg Thr Lys
 50 55 60
 Tyr Gly Thr Lys Ala Gln Leu Gln Thr Ala Ile Ser Asn Leu Arg Gly
 65 70 75 80
 Lys Gly Ile Gly Val Tyr Gly Asp Val Val Met Asn His Lys Gly Gly
 85 90 95
 Ala Asp Tyr Thr Glu Ser Val Gln Ala Ile Glu Val Asn Pro Ser Asn
 100 105 110

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Arg	Asn	Gln	Glu	Thr	Ser	Gly	Glu	Tyr	Gly	Ile	Ser	Ala	Trp	Thr	Gly
		115					120					125			
Phe	Asn	Phe	Ala	Gly	Arg	Asn	Asn	Thr	Tyr	Ser	Pro	Phe	Lys	Trp	Arg
	130					135					140				
Trp	Tyr	His	Phe	Asp	Gly	Thr	Asp	Trp	Asp	Gln	Ser	Arg	Ser	Leu	Ser
145					150					155					160
Arg	Ile	Tyr	Lys	Phe	Lys	Ser	Thr	Gly	Lys	Ala	Trp	Asp	Thr	Asp	Val
				165					170					175	
Ser	Asn	Glu	Asn	Gly	Asn	Tyr	Asp	Tyr	Leu	Met	Tyr	Ala	Asp	Val	Asp
			180					185					190		
Phe	Glu	His	Pro	Glu	Val	Arg	Gln	Glu	Met	Lys	Asn	Trp	Gly	Lys	Trp
		195					200					205			
Tyr	Ala	Asp	Ser	Leu	Gly	Leu	Asp	Gly	Phe	Arg	Leu	Asp	Ala	Val	Lys
	210					215					220				
His	Ile	Ser	His	Ser	Tyr	Leu	Lys	Glu	Trp	Val	Thr	Ser	Val	Arg	Gln
225					230					235					240
Thr	Thr	Gly	Lys	Glu	Met	Phe	Thr	Val	Ala	Glu	Tyr	Trp	Lys	Asn	Asp
				245					250					255	
Leu	Gly	Ala	Ile	Asn	Asp	Tyr	Leu	Tyr	Lys	Thr	Gly	Tyr	Thr	His	Ser
			260					265					270		
Val	Phe	Asp	Val	Pro	Leu	His	Tyr	Asn	Phe	Gln	Ala	Ala	Gly	Asn	Gly
		275					280				285				
Gly	Gly	Tyr	Tyr	Asp	Met	Arg	Asn	Ile	Leu	Lys	Gly	Thr	Val	Thr	Glu
	290					295					300				
Gln	His	Pro	Ser	Leu	Ser	Val	Thr	Ile	Val	Asp	Asn	His	Asp	Ser	Gln
305					310					315					320
Pro	Gly	Gln	Ser	Leu	Glu	Ser	Thr	Val	Ala	Asn	Trp	Phe	Lys	Pro	Leu
				325					330					335	
Ala	Tyr	Ala	Thr	Ile	Met	Thr	Arg	Gly	Gln	Gly	Tyr	Pro	Ala	Leu	Phe
			340					345					350		
Tyr	Gly	Asp	Tyr	Tyr	Gly	Thr	Lys	Gly	Thr	Thr	Asn	Arg	Glu	Ile	Pro
		355					360					365			
Asn	Met	Ser	Gly	Thr	Leu	Gln	Pro	Ile	Leu	Lys	Ala	Arg	Lys	Asp	Phe
	370					375					380				
Ala	Tyr	Gly	Thr	Gln	His	Asp	Tyr	Leu	Asp	His	Gln	Asp	Val	Ile	Gly
385					390					395					400
Trp	Thr	Arg	Glu	Gly	Val	Thr	Asp	Arg	Ala	Lys	Ser	Gly	Leu	Ala	Thr
				405					410					415	
Ile	Leu	Ser	Asp	Gly	Pro	Gly	Gly	Ser	Lys	Trp	Met	Tyr	Val	Gly	Lys
			420					425					430		
Gln	Asn	Ala	Gly	Glu	Val	Trp	Lys	Asp	Met	Thr	Asn	Asn	Asn	Ala	Arg
		435					440					445			
Leu	Val	Thr	Ile	Asn	Ala	Asp	Gly	Trp	Gly	Gln	Phe	Phe	Val	Asn	Gly
	450					455					460				
Gly	Ser	Val	Ser	Ile	Tyr	Thr	Gln	Gln							
465					470										

<210> 165
 <211> 1437
 <212> DNA
 <213> Bacterial

<400> 165	
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gatgctgaaa	atttagctca
ggaacttcac	aaaatgatgt
aatcaaaaag	gaacgatacg
gaggctttac	ataatcaaaa
ggggcagatt	atactgaggt
gaaacatcga	gtgattatca
gactcctatt	ctaattttta
ggaaggaaat	taaatagaat
gtgtctagtg	agaatggtaa
cctgaagttg	caaatgaaat
gatggctttc	gttttagacgc
aatcatgtta	gaaagcaaac
gatattcgta	ctttaaacaa
gcacctcttc	attataattt
	tcattatgct
	gatggggaac
	acatctgtat
	gtgtatgatg
	gggacaaaag
	tacggtgatg
	gttgaggtag
	tggacgggat
	tttcattttg
	aaaggcgtag
	ttaatgtatg
	ggaacctggt
	attgaccatg
	atgtttacag
	aaagtaaat
	tcaacagga
	attggaatcg
	ggataccacc
	tatatgattt
	cacaattaaa
	ttgttatgaa
	accgtaacaa
	ttgattttcc
	atggaacaga
	gtaaagcttg
	cagatcttga
	atgcggaacg
	agtatcttcg
	tagctgaata
	ataatcaatc
	atggaatata
	tttgcaaaat
	tggtgtcgat
	tgatatgagg
	60
	120
	180
	240
	300
	360
	420
	480
	540
	600
	660
	720
	780
	840
	900

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aatatttttaa	aggggtacggt	agtagaaagt	catcctacac	ttgctgttac	tcttgttgag	960
aatcatgatt	ctcagcctgg	acagtcatta	gaatctgttg	tgagtccttg	gtttaagccg	1020
ttggcctatg	catttatttt	aacgcgtgca	gaagggtatc	cttctgtttt	ttatggagat	1080
tactatggca	caaatggaaa	tagtagttat	gaaattccaa	cgttaaagga	taaaattgat	1140
ccaattctga	cggcacgaaa	aaactttgca	tatggtacgc	aacatgatta	tttagaccat	1200
ccagatgtga	ttggctggac	aagagaaggg	gatagtatac	atgctaattc	tggttttagca	1260
acattaatct	ctgatggacc	aggaggatca	aaatggatga	atgttgga	gaacaacgca	1320
ggggaaaat	ggtacgat	tacgggcaat	caaacaata	ctgtaacgat	taataaagat	1380
ggatgggggc	agttccatgt	aaatgggggc	tctgtttcaa	tatatgttca	gaagtaa	1437

<210> 166

<211> 478

<212> PRT

<213> Bacterial

<400> 166

Met	Gln	Tyr	Phe	Glu	Trp	Tyr	Val	Pro	Asn	Asp	Gly	Glu	His	Trp	Asn
1				5					10					15	
Arg	Leu	Arg	Asn	Asp	Ala	Glu	Asn	Leu	Ala	His	Lys	Gly	Ile	Thr	Ser
			20					25					30		
Val	Trp	Ile	Pro	Pro	Val	Tyr	Lys	Gly	Thr	Ser	Gln	Asn	Asp	Val	Gly
		35					40					45			
Tyr	Gly	Val	Tyr	Asp	Val	Tyr	Asp	Leu	Gly	Glu	Phe	Asn	Gln	Lys	Gly
	50					55					60				
Thr	Ile	Arg	Thr	Lys	Tyr	Gly	Thr	Lys	Ala	Gln	Leu	Lys	Ser	Ala	Ile
65					70					75					80
Glu	Ala	Leu	His	Asn	Gln	Asn	Ile	Asp	Val	Tyr	Gly	Asp	Val	Val	Met
				85					90					95	
Asn	His	Lys	Gly	Gly	Ala	Asp	Tyr	Thr	Glu	Val	Val	Thr	Ala	Val	Glu
			100					105					110		
Val	Asp	Arg	Asn	Asn	Arg	Asn	Ile	Glu	Thr	Ser	Ser	Asp	Tyr	Gln	Ile
		115					120					125			
Asp	Ala	Trp	Thr	Gly	Phe	Asp	Phe	Pro	Gly	Arg	Arg	Asp	Ser	Tyr	Ser
	130					135					140				
Asn	Phe	Lys	Trp	Arg	Trp	Phe	His	Phe	Asp	Gly	Thr	Asp	Trp	Asp	Glu
145					150					155					160
Gly	Arg	Lys	Leu	Asn	Arg	Ile	Tyr	Lys	Phe	Lys	Gly	Val	Gly	Lys	Ala
				165					170					175	
Trp	Asp	Trp	Glu	Val	Ser	Ser	Glu	Asn	Gly	Asn	Tyr	Asp	Tyr	Leu	Met
			180					185					190		
Tyr	Ala	Asp	Leu	Asp	Phe	Asp	His	Pro	Glu	Val	Ala	Asn	Glu	Met	Lys
		195					200					205			
Asn	Trp	Gly	Thr	Trp	Tyr	Ala	Asp	Glu	Leu	Asn	Leu	Asp	Gly	Phe	Arg
	210					215					220				
Leu	Asp	Ala	Val	Lys	His	Ile	Asp	His	Glu	Tyr	Leu	Arg	Asp	Trp	Val
225					230					235					240
Asn	His	Val	Arg	Lys	Gln	Thr	Gly	Lys	Glu	Met	Phe	Thr	Val	Ala	Glu
				245					250					255	
Tyr	Trp	Gln	Asn	Asp	Ile	Arg	Thr	Leu	Asn	Asn	Tyr	Leu	Gly	Lys	Val
			260					265					270		
Asn	Tyr	Asn	Gln	Ser	Val	Phe	Asp	Ala	Pro	Leu	His	Tyr	Asn	Phe	His
		275					280					285			
Tyr	Ala	Ser	Thr	Gly	Asn	Gly	Asn	Tyr	Asp	Met	Arg	Asn	Ile	Leu	Lys
	290					295					300				
Gly	Thr	Val	Val	Glu	Ser	His	Pro	Thr	Leu	Ala	Val	Thr	Leu	Val	Glu
305					310					315					320
Asn	His	Asp	Ser	Gln	Pro	Gly	Gln	Ser	Leu	Glu	Ser	Val	Val	Ser	Pro
				325					330					335	
Trp	Phe	Lys	Pro	Leu	Ala	Tyr	Ala	Phe	Ile	Leu	Thr	Arg	Ala	Glu	Gly
			340					345					350		
Tyr	Pro	Ser	Val	Phe	Tyr	Gly	Asp	Tyr	Tyr	Gly	Thr	Asn	Gly	Asn	Ser
		355					360					365			
Ser	Tyr	Glu	Ile	Pro	Thr	Leu	Lys	Asp	Lys	Ile	Asp	Pro	Ile	Leu	Thr
	370					375					380				
Ala	Arg	Lys	Asn	Phe	Ala	Tyr	Gly	Thr	Gln	His	Asp	Tyr	Leu	Asp	His
385					390					395					400
Pro	Asp	Val	Ile	Gly	Trp	Thr	Arg	Glu	Gly	Asp	Ser	Ile	His	Ala	Asn
				405					410					415	

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Ser Gly Leu Ala Thr Leu Ile Ser Asp Gly Pro Gly Gly Ser Lys Trp
 420 425 430
 Met Asn Val Gly Lys Asn Asn Ala Gly Glu Ile Trp Tyr Asp Ile Thr
 435 440 445
 Gly Asn Gln Thr Asn Thr Val Thr Ile Asn Lys Asp Gly Trp Gly Gln
 450 455 460
 Phe His Val Asn Gly Gly Ser Val Ser Ile Tyr Val Gln Lys
 465 470 475

<210> 167
 <211> 1596
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 167
 atgcaaacga ttgcaaaaaa aggggatgaa acgatgaaag ggaaaaaatg gacagcttta 60
 gctctaacac tgccgctggc tgctagctta tcaacaggcg ttcacgccga aaccgtacat 120
 aaagggtaaat ctccaacagc agataaaaac ggtgtatttt atgaggtgta tgtaactctt 180
 ttttacgatg caaataaaga tggacatggt gatttaaaag gtccttacaca aaagttggat 240
 tatttaaatg atggcaattc tcatacaaaag aatgatcttc aagtaaacgg gatttgatg 300
 atgccggtca acccttctcc cagctatcat aaatatgatg taacggacta ttataatatt 360
 gatccgcagt atggaaatct gcaagatttt cgcaaactga tgaaagaagc agataaacga 420
 gatgtaaaag tcattatgga cctcgtttgt aatcatacga gcagtgaaca cccttggttt 480
 caagctgcat taaaagataa aaacagcaag tacagagatt actatatctg ggctgataaa 540
 aataccgact tgaatgaaaa aggatcttgg ggacagcaag tatggcataa agccccaac 600
 ggagagtatt ttacggaac gttttgggaa ggaatgccgg acttaaatta cgataatcct 660
 gaagtaagaa aagaaatgat taacgtagga aagttttggc taaagcaagg agttgacggg 720
 ttccgtctag atgctgcgct tcataattttt aaaggccaaa cacctgaagg cgctaagaaa 780
 aatctcctgt ggtggaatga atttagagat gcaatgaaaa aggaaaacc taacgtatat 840
 ctaacgggtg aagtatggga tcaaccggaa gtagtagctc cttactatca atcgcttgat 900
 tctttattta actttgattt agcaggaag attgtaaact ctgtaaaatc aggaaatgat 960
 caaggaatcg cgactgcagc agcggcaacg gatgaactgt tcaaatcata caatccaaat 1020
 aaaattgacg gtattttctt aaccaacctt gaccaaactc gcgtcatgag tgagctaagc 1080
 ggcatgtga ataaagcaaa gtcagctgcc tctatcttac ttacgcttcc tggcaaccg 1140
 tatatttatt acggtgaaga aatcggcatg accggtgaaa agcctgatga gttaatccgt 1200
 gaaccgttcc gctggtacga aggaaacgga cttggacaaa ccagctggga aacacctgta 1260
 tacaacaaag gcggcaacgg cgtgtctgta gaagcacaaa caaaacaaa ggactctttg 1320
 ttaaatcatt accgtgaaat gattcgcgtg cgtcagcagc acgaagagtt agtaaaagga 1380
 acgcttcaat ctatttcagt agacagtaaa gaagtcgttg cctatagccg cacgtataaa 1440
 ggcaaatcga ttagcgtgta tcataatatt tcaaatcaac cggtaaaagt atctgtagca 1500
 gcaaaaggta aattgatttt tggtagtgaa aaaggtgcta agaaagtcaa aaatcagctt 1560
 gtgattccgg cgaatacaac ggttttaata aaataa 1596

<210> 168
 <211> 531
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 168
 Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
 1 5 10 15
 Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr
 20 25 30
 Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr Ala Asp
 35 40 45
 Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala
 50 55 60
 Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp
 65 70 75 80
 Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn
 85 90 95
 Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr

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Asp	Val	Thr	100	Asp	Tyr	Tyr	Asn	Ile	105	Asp	Pro	Gln	Tyr	Gly	110	Asn	Leu	Gln
		115						120						125				
Asp	Phe	Arg	Lys	Leu	Met	Lys	Glu	Ala	Asp	Lys	Arg	Asp	Val	Lys	Val			
	130					135					140							
Ile	Met	Asp	Leu	Val	Val	Asn	His	Thr	Ser	Ser	Glu	His	Pro	Trp	Phe			
145					150					155					160			
Gln	Ala	Ala	Leu	Lys	Asp	Lys	Asn	Ser	Lys	Tyr	Arg	Asp	Tyr	Tyr	Ile			
			165						170					175				
Trp	Ala	Asp	Lys	Asn	Thr	Asp	Leu	Asn	Glu	Lys	Gly	Ser	Trp	Gly	Gln			
			180					185					190					
Gln	Val	Trp	His	Lys	Ala	Pro	Asn	Gly	Glu	Tyr	Phe	Tyr	Gly	Thr	Phe			
		195					200					205						
Trp	Glu	Gly	Met	Pro	Asp	Leu	Asn	Tyr	Asp	Asn	Pro	Glu	Val	Arg	Lys			
	210					215					220							
Glu	Met	Ile	Asn	Val	Gly	Lys	Phe	Trp	Leu	Lys	Gln	Gly	Val	Asp	Gly			
225					230					235					240			
Phe	Arg	Leu	Asp	Ala	Ala	Leu	His	Ile	Phe	Lys	Gly	Gln	Thr	Pro	Glu			
				245					250					255				
Gly	Ala	Lys	Lys	Asn	Leu	Leu	Trp	Trp	Asn	Glu	Phe	Arg	Asp	Ala	Met			
			260					265					270					
Lys	Lys	Glu	Asn	Pro	Asn	Val	Tyr	Leu	Thr	Gly	Glu	Val	Trp	Asp	Gln			
		275					280					285						
Pro	Glu	Val	Val	Ala	Pro	Tyr	Tyr	Gln	Ser	Leu	Asp	Ser	Leu	Phe	Asn			
	290					295					300							
Phe	Asp	Leu	Ala	Gly	Lys	Ile	Val	Asn	Ser	Val	Lys	Ser	Gly	Asn	Asp			
305				310						315					320			
Gln	Gly	Ile	Ala	Thr	Ala	Ala	Ala	Ala	Thr	Asp	Glu	Leu	Phe	Lys	Ser			
				325					330					335				
Tyr	Asn	Pro	Asn	Lys	Ile	Asp	Gly	Ile	Phe	Leu	Thr	Asn	His	Asp	Gln			
			340					345					350					
Asn	Arg	Val	Met	Ser	Glu	Leu	Ser	Gly	Asp	Val	Asn	Lys	Ala	Lys	Ser			
		355					360					365						
Ala	Ala	Ser	Ile	Leu	Leu	Thr	Leu	Pro	Gly	Asn	Pro	Tyr	Ile	Tyr	Tyr			
		370				375					380							
Gly	Glu	Glu	Ile	Gly	Met	Thr	Gly	Glu	Lys	Pro	Asp	Glu	Leu	Ile	Arg			
385				390						395					400			
Glu	Pro	Phe	Arg	Trp	Tyr	Glu	Gly	Asn	Gly	Leu	Gly	Gln	Thr	Ser	Trp			
				405					410					415				
Glu	Thr	Pro	Val	Tyr	Asn	Lys	Gly	Gly	Asn	Gly	Val	Ser	Val	Glu	Ala			
			420					425					430					
Gln	Thr	Lys	Gln	Lys	Asp	Ser	Leu	Leu	Asn	His	Tyr	Arg	Glu	Met	Ile			
		435					440					445						
Arg	Val	Arg	Gln	Gln	His	Glu	Glu	Leu	Val	Lys	Gly	Thr	Leu	Gln	Ser			
						455					460							
Ile	Ser	Val	Asp	Ser	Lys	Glu	Val	Val	Ala	Tyr	Ser	Arg	Thr	Tyr	Lys			
465					470					475					480			
Gly	Lys	Ser	Ile	Ser	Val	Tyr	His	Asn	Ile	Ser	Asn	Gln	Pro	Val	Lys			
				485					490					495				
Val	Ser	Val	Ala	Ala	Lys	Gly	Lys	Leu	Ile	Phe	Gly	Ser	Glu	Lys	Gly			
			500					505					510					
Ala	Lys	Lys	Val	Lys	Asn	Gln	Leu	Val	Ile	Pro	Ala	Asn	Thr	Thr	Val			
		515					520					525						
Leu	Ile	Lys																
	530																	

<210> 169
 <211> 1524
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 169	
atgaaaacat tcaaatataa acgcactttt ttaccgctaa ctttgctgct cagtgtctct	60
gcctttgctg ggcaaatgg caccatgatg cagtattttc attggtatgt acctaataatgat	120
ggcgcattat ggacgcaggt tgaaagcaat gctccagcac tcgctgaaaa cggttttaca	180

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gcgctctggc	taccgccagc	ttacaaaggc	gcgggcggca	gtaatgacgt	cggttatggc	240
gtctatgata	tgtacgattt	aggtgagttt	gatcaaaaag	gctcagtagc	aaccaaatat	300
ggcaccaagg	ctcagtagat	ctctgcaatc	aatgccgcgc	acaacaacaa	tatccaaatc	360
tacggcgatg	ttgtgtttta	ccaccgaggt	ggtgctgatg	ggaagtcgtg	ggtcgatacc	420
aagcgcggtg	attgggacaa	ccgtaacatt	gaactgggcg	acaaatggat	tgaagcttgg	480
gttgagttta	attttcctgg	ccgcaacgac	aaatactcaa	acttccattg	gacttgggtat	540
cactttgacg	gtgttgactg	ggatgatgcc	ggcaaagaaa	aagcgatctt	taaattcaaa	600
ggcgaaggaa	aagcatggga	ttgggaagtc	agctctgaaa	aaggcaatta	cgactaccta	660
atgtacgccg	atttagacat	ggatcaccaa	gaagttaaac	aagagctgaa	agattgggggt	720
gagtggtaca	tcaacatgac	cggcgttgat	ggcttttagaa	tggatgccgt	gaagcacatt	780
aaatatcagt	atctacaaga	gtggattgat	catttacgtt	ggaaaacagg	caaagagctt	840
ttaccggttg	gtgagtattg	gaattacgac	gtaaatacaac	tgcataactt	tattactaag	900
acctctggca	gtatgtcgtt	gttcgatgcg	ccgcttcaca	tgaactttcta	caacgcgtca	960
aaatctggcg	gcaattacga	tatgcgccaa	atcatgaatg	gcacgttgat	gaaggacaac	1020
ccagtcaaag	ctgtgactct	cgtagaaaac	cacgatacac	agccattgca	ggcgttagag	1080
tcgacagtgg	attggtgggtt	caagcctctt	gcttacgcac	tcatttttatt	gcgtgaagaa	1140
ggttatccat	cagtgttcta	cgcagattac	tacggcgcg	agtacagcga	caaaggctac	1200
aacatcaata	tggccaaagt	tccttacatt	gaagaacttg	taacactgcg	taaagagtat	1260
gcgtatggca	aacagaattc	ttatctcgac	cactgggatg	tgattggctg	gacccgagag	1320
ggcgatgctg	aacatccaaa	ctcaatggcg	gtgatcgacc	gtgatggacc	aggtggcaaa	1380
aaatggatgt	ataccggtaa	cccaagcacg	cgctatgtcg	acaagctggg	tatccgaact	1440
gaagaagttt	ggaccgatac	caatggctgg	gcagaatttc	ctgtcaatgg	tggttcagtc	1500
tcggtttggg	tgggcgttaa	gtaa				1524

<210> 170
 <211> 507
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 170
 Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu
 1 5 10 15
 Leu Ser Ala Pro Ala Phe Ala Gly Gln Asn Gly Thr Met Met Gln Tyr
 20 25 30
 Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu
 35 40 45
 Ser Asn Ala Pro Ala Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu
 50 55 60
 Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly
 65 70 75 80
 Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ser Val
 85 90 95
 Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile Ser Ala Ile Asn Ala
 100 105 110
 Ala His Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His
 115 120 125
 Arg Gly Gly Ala Asp Gly Lys Ser Trp Val Asp Thr Lys Arg Val Asp
 130 135 140
 Trp Asp Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp
 145 150 155 160
 Val Glu Phe Asn Phe Pro Gly Arg Asn Asp Lys Tyr Ser Asn Phe His
 165 170 175
 Trp Thr Trp Tyr His Phe Asp Gly Val Asp Trp Asp Asp Ala Gly Lys
 180 185 190
 Glu Lys Ala Ile Phe Lys Phe Lys Gly Glu Gly Lys Ala Trp Asp Trp
 195 200 205
 Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp
 210 215 220
 Leu Asp Met Asp His Gln Glu Val Lys Gln Glu Leu Lys Asp Trp Gly
 225 230 235 240
 Glu Trp Tyr Ile Asn Met Thr Gly Val Asp Gly Phe Arg Met Asp Ala
 245 250 255
 Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp His Leu
 260 265 270
 Arg Trp Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn

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Tyr Asp 275 Val Asn Gln Leu His 280 Asn Phe Ile Thr Lys 285 Thr Ser Gly Ser
 290 Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala Ser
 305 Lys Ser Gly Gly Asn Tyr Asp Met Arg Gln Ile Met Asn Gly Thr Leu
 325 Met Lys Asp Asn Pro Val Lys Ala Val Thr Leu Val Glu Asn His Asp
 340 Thr Gln Pro Leu Gln Ala Leu Glu Ser Thr Val Asp Trp Trp Phe Lys
 355 Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser
 370 Val Phe Tyr Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly Tyr
 385 Asn Ile Asn Met Ala Lys Val Pro Tyr Ile Glu Glu Leu Val Thr Leu
 405 Arg Lys Glu Tyr Ala Tyr Gly Lys Gln Asn Ser Tyr Leu Asp His Trp
 420 Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser
 435 Met Ala Val Ile Met Ser Asp Gly Pro Gly Gly Lys Lys Trp Met Tyr
 450 Thr Gly Lys Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr
 465 Glu Glu Val Trp Thr Asp Thr Asn Gly Trp Ala Glu Phe Pro Val Asn
 485 Gly Gly Ser Val Ser Val Trp Val Gly Val Lys
 500

<210> 171
 <211> 1431
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 171
 gtgtatgtaa actctttttta cgatgcaa ataatgatggc atgggtgattt aaaaggtctt 60
 acacaaaagt tggattatatt aaatgatggc aattctcata caaagaatga tcttcaagta 120
 aacgggattt ggatgatgcc ggtcaaccct tctcccagct atcataaata tgatgtaacg 180
 gactattata atattgatcc gcagtatgga aatctgcaag attttcgcaa actgatgaaa 240
 gaagcagata aacgagatgt aaaagtcatt atggacctcg ttgtgaatca tacgagcagt 300
 gaacaccctt ggtttcaagc tgcattaaaa gataaaaaca gcaagtacag agattactat 360
 atctgggctg ataaaaatac cgacttgaat gaaaaaggat cttggggaca gcaagtatgg 420
 cataaagccc caaacggaga gtattttttac ggaacgtttt gggaaggaaat gccggactta 480
 aattacgata atcctgaagt aagaaaagaa atgattaacg taggaaagt ttggctaaag 540
 caaggagtgt acgggttccg tctagatgct gcgcttcata tttttaaaagg ccaaaccct 600
 gaagcgcta agaaaaatct cctgtggtgg aatgaattta gagatgcaat gaaaaaggaa 660
 aaccctaacg tatatctaac gggatgaagta tgggatcaac cggaagtagt agctccttac 720
 tatcaatcgc ttgattcttt atttaacttt gatttagcag gaaagattgt aaactctgta 780
 aaatcaggaa atgatcaagg aatcgcgact gcagcagcgg caacggatga actgttcaaa 840
 tcatacaatc caaataaaat tgacgggtatt ttcttaacca accatgacca aaatcgcgtc 900
 atgagtggc taagcggcga tgtgaataaa gcaaagtcag ctgcctctat cttacttacg 960
 cttcctggca acccgatat ttattacggg gaagaaatcg gcatgaccgg tgaaaagcct 1020
 gatgagttaa tccgtgaacc gttccgctgg tacgaaggaa acggacttgg acaaaccagc 1080
 tgggaaacac ctgtatacaa caaaggcggc aacggcgtgt ctgtagaagc acaaacaaaa 1140
 caaaaggact ctttgttaaa tcattaccgt gaaatgattc gcgtgcgtca gcagcacgaa 1200
 gagttagtaa aaggaacgct tcaatctatt tcagtagaca gtaaagaagt cgttgccat 1260
 agcgcacgt ataaaggcaa atcgattagc gtgtatcata atatttcaaa tcaaccggt 1320
 aaagtatctg tagcagcaaa aggtaaattg atttttggta gtgaaaaagg tgctaagaaa 1380
 gtcaaaaatc agcttgtgat tccggcgaat acaacggttt taataaaata a 1431

<210> 172
 <211> 476
 <212> PRT
 <213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 172

Val Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp
 1 5 10 15
 Leu Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser
 20 25 30
 His Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val
 35 40 45
 Asn Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn
 50 55 60
 Ile Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys
 65 70 75 80
 Glu Ala Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn
 85 90 95
 His Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys
 100 105 110
 Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp
 115 120 125
 Leu Asn Glu Lys Gly Ser Trp Gly Gln Gln Val Trp His Lys Ala Pro
 130 135 140
 Asn Gly Glu Tyr Phe Tyr Gly Thr Phe Trp Glu Gly Met Pro Asp Leu
 145 150 155 160
 Asn Tyr Asp Asn Pro Glu Val Arg Lys Glu Met Ile Asn Val Gly Lys
 165 170 175
 Phe Trp Leu Lys Gln Gly Val Asp Gly Phe Arg Leu Asp Ala Ala Leu
 180 185 190
 His Ile Phe Lys Gly Gln Thr Pro Glu Gly Ala Lys Lys Asn Leu Leu
 195 200 205
 Trp Trp Asn Glu Phe Arg Asp Ala Met Lys Lys Glu Asn Pro Asn Val
 210 215 220
 Tyr Leu Thr Gly Glu Val Trp Asp Gln Pro Glu Val Val Ala Pro Tyr
 225 230 235 240
 Tyr Gln Ser Leu Asp Ser Leu Phe Asn Phe Asp Leu Ala Gly Lys Ile
 245 250 255
 Val Asn Ser Val Lys Ser Gly Asn Asp Gln Gly Ile Ala Thr Ala Ala
 260 265 270
 Ala Ala Thr Asp Glu Leu Phe Lys Ser Tyr Asn Pro Asn Lys Ile Asp
 275 280 285
 Gly Ile Phe Leu Thr Asn His Asp Gln Asn Arg Val Met Ser Glu Leu
 290 295 300
 Ser Gly Asp Val Asn Lys Ala Lys Ser Ala Ala Ser Ile Leu Leu Thr
 305 310 315 320
 Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr
 325 330 335
 Gly Glu Lys Pro Asp Glu Leu Ile Arg Glu Pro Phe Arg Trp Tyr Glu
 340 345 350
 Gly Asn Gly Leu Gly Gln Thr Ser Trp Glu Thr Pro Val Tyr Asn Lys
 355 360 365
 Gly Gly Asn Gly Val Ser Val Glu Ala Gln Thr Lys Gln Lys Asp Ser
 370 375 380
 Leu Leu Asn His Tyr Arg Glu Met Ile Arg Val Arg Gln Gln His Glu
 385 390 395 400
 Glu Leu Val Lys Gly Thr Leu Gln Ser Ile Ser Val Asp Ser Lys Glu
 405 410 415
 Val Val Ala Tyr Ser Arg Thr Tyr Lys Gly Lys Ser Ile Ser Val Tyr
 420 425 430
 His Asn Ile Ser Asn Gln Pro Val Lys Val Ser Val Ala Ala Lys Gly
 435 440 445
 Lys Leu Ile Phe Gly Ser Glu Lys Gly Ala Lys Lys Val Lys Asn Gln
 450 455 460
 Leu Val Ile Pro Ala Asn Thr Thr Val Leu Ile Lys
 465 470 475

<210> 173

<211> 1596

<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 173
 atgcaaacga ttgcaaaaaa aggggatgaa acgatgaaag ggaaaaaatg gacagcttta 60
 gctctaacac tgccgctggc tgctagctta tcaacaggcg ttcacgcaga aactgtacat 120
 aaaggtaaag ctccaacagc agataaaaaac ggtgtttttt atgaggtgta tgtaaactct 180
 ttttacgatg caaataaaga tggacatggg gatttaaaag gtctgacaca aaagttggat 240
 tatttaaatg acggcaattc tcatacaaaag aatgatcttc aagtaaaccg gattttggatg 300
 atgccggtaa acccttctcc tagctatcat aaatatgatg taacggacta ttataacatt 360
 gatccctcagt acggaagtct gcaagatttc cgcaaactga tgaaagaagc agataaacga 420
 gacgtaaaaag ttattatgga ccttggtgtg aatcatatcga gcagtgaaca cccttggttt 480
 caagctgcac taaaagataa aaacagcaag tacagagatt actatatattg ggctgataaa 540
 aataccgatt tgaatgaaaa aggatcttgg ggacagcaag tatggcataa agctccaaac 600
 ggagagtatt tttacggaac gttctgggaa ggaatgcctg acttaaatta cgataaccct 660
 gaagtaagaa aagaaatgat taacgtcggg aagttttggc taaagcaagg cgttgatggc 720
 ttccgcttag atgtgccc tcatatcttt aaaggtcaaa ctctgaagg cgctaagaaa 780
 aatctcctgt ggtggaatga gtttagagat gcaatgaaaa aagaaaacc taacgtatat 840
 ctaacgggtg aagtatggga tcagccggaa gtagtagctc cttattatca atcgcttgat 900
 tccctattta actttgattt agcaggaaaa attgtcagct ctgtaaaagc aggaaatgat 960
 caaggaatcg ccactgcagc agcggcaacg gatgagctgt tcaaatcata caatccaaat 1020
 aaaattgacg gcattttctt aaccaaccat gaccaaacc gcgtcatgag tgagctaagc 1080
 ggagatgtga ataaagcaaa atcagctgct tctatcttac ttacgcttcc tggaaatccg 1140
 tatatttatt acggtgaaga aattggcatg accggtgaaa agcctgatga attaatccgt 1200
 gaaccgttcc gctggtacga aggcaacgga attggacaaa ctagctggga aacacctgta 1260
 tataacaaag gcggcaatgg tgtgtctgta gaagcacaaa ccaaacaaaa ggattctttg 1320
 ttaaatcatt accgtgaaat gattcgcgtg cgtcagcagc acgaagagtt agtaaaagga 1380
 acgcttcagt ctatttcagt agacagtaaa gaagttgtcg cttatagccg tacgtataaa 1440
 ggcaactcca ttagtgtgta tcataatatt tcaaatcaac ctgtaaaagt atctgtagcg 1500
 gcgaaaggta aattgatttt tgctagttaa aaaggtgcta aaaaaggcaa aaatcagctt 1560
 gtgattccgg cgaatgcgac ggttttaata aaataa 1596

<210> 174
<211> 531
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 174
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 Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr
 20 25 30
 Gly Val His Ala Glu Thr Val His Lys Gly Lys Ala Pro Thr Ala Asp
 35 40 45
 Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala
 50 55 60
 Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp
 65 70 75 80
 Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn
 85 90 95
 Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr
 100 105 110
 Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Ser Leu Gln
 115 120 125
 Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val
 130 135 140
 Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
 145 150 155 160
 Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile
 165 170 175
 Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp Gly Gln
 180 185 190

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Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe
 195 200 205
 Trp Glu Gly Met Pro Asp Leu Tyr Asp Asn Pro Glu Val Arg Lys
 210 215 220
 Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly
 225 230 235 240
 Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu
 245 250 255
 Gly Ala Lys Lys Asn Leu Leu Trp Trp Asn Glu Phe Arg Asp Ala Met
 260 265 270
 Lys Lys Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln
 275 280 285
 Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn
 290 295 300
 Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp
 305 310 315 320
 Gln Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser
 325 330 335
 Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His Asp Gln
 340 345 350
 Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys Ser
 355 360 365
 Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr
 370 375 380
 Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg
 385 390 395 400
 Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Ile Gly Gln Thr Ser Trp
 405 410 415
 Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala
 420 425 430
 Gln Thr Lys Gln Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile
 435 440 445
 Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser
 450 455 460
 Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys
 465 470 475 480
 Gly Asn Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys
 485 490 495
 Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala Ser Glu Lys Gly
 500 505 510
 Ala Lys Lys Gly Lys Asn Gln Leu Val Ile Pro Ala Asn Ala Thr Val
 515 520 525
 Leu Ile Lys
 530

<210> 175
 <211> 1398
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 175
 atgaaaaata taatacgact ttgtgctgcc agcgctatcc tcacggtgtc ccacgccagt 60
 tacgccgacg caattttaca cgcgtttaac tggcaatata ccgatgtaac cgccaatgca 120
 aatcaaatgg ccgcaaatgg ctttaaaaaa gtcctcattt caccgcgaat gaaatccagc 180
 ggcagtcaat ggtgggcccg ctatcaaccg caagacttgc gtgtcattga ttctccgctg 240
 ggcaacaaac aagatttagt cgcgatgatc aatgcgctca acagcgttgg ggtcgacgtg 300
 tatgctgacg tgggtgctta ccatatggct aacgagtcac ggaagcgcag tgacctgaac 360
 taccgggga gtgaggtgct caacgactat caatcccga gtgcttacta tcaaaggcaa 420
 acacttttcg gcaatttaca ggagaacctt ttttccgaga atgatttcca tccggcaggc 480
 tgtattacca attggaatga tcctggccac gtccagtatt ggcgcttggt cggcggacag 540
 ggcgatactg ggctaccgga tctcgatcct aatcaatggg ttgtgagtca gcagaagagt 600
 tacttgaacg cactcaaadc aatgggaatc aaagggttcc gtatcgatgc ggtcaaacat 660
 atgagtcaat atcaaataga ccaagtgttt accccagaca ttaccgctgg tatgcatata 720
 ttcggaag tcattaccga tgggtgggcaa gctatgaggc ttttcttgcc 780
 ccttacctta ataataccga tcacgccgct tatgacttcc cgctatttgc atcgattcga 840

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gccgcgtttt	cattctcttg	tgggttaa	cagctacaca	atccacaagc	ctatggccaa	900
gcgttacagg	actcacgtgc	gataccttt	acgattaccc	acgacattcc	aaccaatgac	960
ggtttcgcgt	accagatcat	ggatccaacc	gatgaacagc	tcgcctatgc	ctacatcttg	1020
ggcaaagatg	gaggaacgcc	acttgtctat	agtgatgacc	tacctgacag	cgaagacaaa	1080
gacagtgggtc	gttgggccga	tgtgtggcaa	gatccgaaca	tgattaacat	gcttgccttc	1140
cacaacgcga	tgcaaggaca	aagcatgact	gtagtggcta	gcgatcaatg	taccttgcta	1200
tttaagcgcg	gcaagcaagg	cgtggtagga	atcaataaat	gtggcgagag	taagtcggtg	1260
actgtcgata	cttaccagca	tgagtttaac	tggtacaccc	cgtaccaaga	cgtattgagc	1320
ggcgacatca	ccacagttag	ttctcgttat	caccaatttg	ttttgccagc	gcgcagtgca	1380
aggatgtgga	aactataa					1398

<210> 176

<211> 465

<212> PRT

<213> Unknown

<220>

<223> obtained from an environmental sample.

<400> 176

Met	Lys	Asn	Ile	Ile	Arg	Leu	Cys	Ala	Ala	Ser	Ala	Ile	Leu	Thr	Val	
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Ser	His	Ala	Ser	Tyr	Ala	Asp	Ala	Ile	Leu	His	Ala	Phe	Asn	Trp	Gln	
			20					25				30				
Tyr	Thr	Asp	Val	Thr	Ala	Asn	Ala	Asn	Gln	Ile	Ala	Ala	Asn	Gly	Phe	
		35					40				45					
Lys	Lys	Val	Leu	Ile	Ser	Pro	Ala	Met	Lys	Ser	Ser	Gly	Ser	Gln	Trp	
	50					55					60					
Trp	Ala	Arg	Tyr	Gln	Pro	Gln	Asp	Leu	Arg	Val	Ile	Asp	Ser	Pro	Leu	
65					70				75						80	
Gly	Asn	Lys	Gln	Asp	Leu	Val	Ala	Met	Ile	Asn	Ala	Leu	Asn	Ser	Val	
			85						90					95		
Gly	Val	Asp	Val	Tyr	Ala	Asp	Val	Val	Leu	Asn	His	Met	Ala	Asn	Glu	
			100					105					110			
Ser	Trp	Lys	Arg	Ser	Asp	Leu	Asn	Tyr	Pro	Gly	Ser	Glu	Val	Leu	Asn	
		115					120					125				
Asp	Tyr	Gln	Ser	Arg	Ser	Ala	Tyr	Tyr	Gln	Arg	Gln	Thr	Leu	Phe	Gly	
	130					135					140					
Asn	Leu	Gln	Glu	Asn	Leu	Phe	Ser	Glu	Asn	Asp	Phe	His	Pro	Ala	Gly	
145					150					155					160	
Cys	Ile	Thr	Asn	Trp	Asn	Asp	Pro	Gly	His	Val	Gln	Tyr	Trp	Arg	Leu	
			165						170					175		
Cys	Gly	Gly	Gln	Gly	Asp	Thr	Gly	Leu	Pro	Asp	Leu	Asp	Pro	Asn	Gln	
			180					185					190			
Trp	Val	Val	Ser	Gln	Gln	Lys	Ser	Tyr	Leu	Asn	Ala	Leu	Lys	Ser	Met	
		195					200					205				
Gly	Ile	Lys	Gly	Phe	Arg	Ile	Asp	Ala	Val	Lys	His	Met	Ser	Gln	Tyr	
	210					215					220					
Gln	Ile	Asp	Gln	Val	Phe	Thr	Pro	Asp	Ile	Thr	Ala	Gly	Met	His	Ile	
225					230					235					240	
Phe	Gly	Glu	Val	Ile	Thr	Ser	Gly	Gly	Gln	Gly	Asp	Ser	Gly	Tyr	Glu	
			245						250					255		
Ala	Phe	Leu	Ala	Pro	Tyr	Leu	Asn	Asn	Thr	Asp	His	Ala	Ala	Tyr	Asp	
			260					265					270			
Phe	Pro	Leu	Phe	Ala	Ser	Ile	Arg	Ala	Ala	Phe	Ser	Phe	Ser	Gly	Gly	
		275					280					285				
Leu	Asn	Gln	Leu	His	Asn	Pro	Gln	Ala	Tyr	Gly	Gln	Ala	Leu	Gln	Asp	
	290					295					300					
Ser	Arg	Ala	Ile	Thr	Phe	Thr	Ile	Thr	His	Asp	Ile	Pro	Thr	Asn	Asp	
305					310					315				320		
Gly	Phe	Arg	Tyr	Gln	Ile	Met	Asp	Pro	Thr	Asp	Glu	Gln	Leu	Ala	Tyr	
			325						330					335		
Ala	Tyr	Ile	Leu	Gly	Lys	Asp	Gly	Gly	Thr	Pro	Leu	Val	Tyr	Ser	Asp	
			340					345					350			
Asp	Leu	Pro	Asp	Ser	Glu	Asp	Lys	Asp	Ser	Gly	Arg	Trp	Ala	Asp	Val	
		355					360					365				
Trp	Gln	Asp	Pro	Asn	Met	Ile	Asn	Met	Leu	Ala	Phe	His	Asn	Ala	Met	
	370					375					380					

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Gln Gly Gln Ser Met Thr Val Val Ala Ser Asp Gln Cys Thr Leu Leu
385 390 395 400
Phe Lys Arg Gly Lys Gln Gly Val Val Gly Ile Asn Lys Cys Gly Glu
405 410 415
Ser Lys Ser Val Thr Val Asp Thr Tyr Gln His Glu Phe Asn Trp Tyr
420 425 430
Thr Pro Tyr Gln Asp Val Leu Ser Gly Asp Ile Thr Thr Val Ser Ser
435 440 445
Arg Tyr His Gln Phe Val Leu Pro Ala Arg Ser Ala Arg Met Trp Lys
450 455 460
Leu
465

<210> 177
<211> 1524
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 177
atgaaaacat tcaaattaaa acgcactttt ttaccgctga ccttgctgct cagtgcctct 60
gcctttgctg ggcaaaatgg caccatgatg cagtattttc attggtacgt acctaattgat 120
ggcgcatat ggacgcaggt tgaaagcaat gctccagtac tcgctgaaaa cggttttaca 180
gcgctctggc taccgcccgc atacaaaggc gcgggcggca gtaatgacgt cggttatggc 240
gtctatgata tgtacgattt aggtgagttt gaccaaaaag gctcagtacg aaccaaatac 300
ggcaccaagg ctacgtacat ctctgcaatc aatgccgcgc acaacaacaa tatccaaatt 360
tacggcgacg ttgtgtttaa ccaccgaggt ggcgctgatg ggaagtcgtg ggtcgatacc 420
aagcgcgttg attgggacaa ccgcaatatt gaactgggcg acaaatggat tgaagcttgg 480
gttgagttaa attttcctgg ccgcaacgac aaatactcga acttccattg gacttggtat 540
cactttgacg gtgttgactg ggatgatgcc ggcaaagaaa aagcgatctt taaattcaaa 600
ggcgaaggaa aagcatggga ttgggaagtc agctctgaaa aaggcaatta cgactaccta 660
atgtacgccg atttagacat ggatcaccca gaagttaaac aagagctgaa agattggggg 720
gagtggtaga tcaacatgac cggcgttgat ggcttttaga tggatgccgt gaagcacatt 780
aaatatcagt atctacaaga gtggattgat catttacgtt ggaaaacagg caaagagctt 840
ttcaccgttg gtgagtattg gaattacgac gtaaatcaac tgcacaactt tattactaag 900
acctctggca gtatgtcgtt gttcgtatgc ccgcttcaca tgaatttcta caacgcgtca 960
aaatctggcg gcacttacga tatgcgccaa atcatgaatg gcacgttgat gaaggacaac 1020
ccagtcaaag cagtgactct cgtagaaaac cagcgtacgc agccattgca ggcgttagag 1080
tcgacagtag attggtgggt caagcctctt gcttacgcat tcattttatt gcgtgaagaa 1140
ggttatccat cgggtgttcta cgcagattac tacggcgcgc agtacagcga caaaggttac 1200
aacattaata tggccaaagt gccttacatt gaagaacttg taacactgcg taaagagtat 1260
gcgtatggca aacagaattc ttatctcgac cattgggatg tgattggctg gacccgagag 1320
ggcgtatgctg aacatccaaa ctcaatggcg gtgatcatga gtgatggacc gggcggcaca 1380
aaatggatgt ataccggtaa gccaagtacg cgctatgtcg acaagctggg tatccgaact 1440
gaagatgttt ggaccgatgc caatggctgg gcagaatttc ctgtcaatgg tggttcagtc 1500
tcggtttggg tgggcgttaa gtaa 1524

<210> 178
<211> 507
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 178
Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu
1 5 10 15
Leu Ser Ala Pro Ala Phe Ala Gly Gln Asn Gly Thr Met Met Gln Tyr
20 25 30
Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu
35 40 45
Ser Asn Ala Pro Val Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu
50 55 60
Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly
65 70 75 80

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Val	Tyr	Asp	Met	Tyr	Asp	Leu	Gly	Glu	Phe	Asp	Gln	Lys	Gly	Ser	Val
Arg	Thr	Lys	Tyr	Gly	Thr	Lys	Ala	Gln	Tyr	Ile	Ser	Ala	Ile	Asn	Ala
Ala	His	Asn	Asn	Asn	Ile	Gln	Ile	Tyr	Gly	Asp	Val	Val	Phe	Asn	His
Arg	Gly	Gly	Ala	Asp	Gly	Lys	Ser	Trp	Val	Asp	Thr	Lys	Arg	Val	Asp
Trp	Asp	Asn	Arg	Asn	Ile	Glu	Leu	Gly	Asp	Lys	Trp	Ile	Glu	Ala	Trp
Val	Glu	Phe	Asn	Phe	Pro	Gly	Arg	Asn	Asp	Lys	Tyr	Ser	Asn	Phe	His
Trp	Thr	Trp	Tyr	His	Phe	Asp	Gly	Val	Asp	Trp	Asp	Asp	Ala	Gly	Lys
Glu	Lys	Ala	Ile	Phe	Lys	Phe	Lys	Gly	Glu	Gly	Lys	Ala	Trp	Asp	Trp
Glu	Val	Ser	Ser	Glu	Lys	Gly	Asn	Tyr	Asp	Tyr	Leu	Met	Tyr	Ala	Asp
Leu	Asp	Met	Asp	His	Pro	Glu	Val	Lys	Gln	Glu	Leu	Lys	Asp	Trp	Gly
Glu	Trp	Tyr	Ile	Asn	Met	Thr	Gly	Val	Asp	Gly	Phe	Arg	Met	Asp	Ala
Val	Lys	His	Ile	Lys	Tyr	Gln	Tyr	Leu	Gln	Glu	Trp	Ile	Asp	His	Leu
Arg	Trp	Lys	Thr	Gly	Lys	Glu	Leu	Phe	Thr	Val	Gly	Glu	Tyr	Trp	Asn
Tyr	Asp	Val	Asn	Gln	Leu	His	Asn	Phe	Ile	Thr	Lys	Thr	Ser	Gly	Ser
Met	Ser	Leu	Phe	Asp	Ala	Pro	Leu	His	Met	Asn	Phe	Tyr	Asn	Ala	Ser
Lys	Ser	Gly	Gly	Thr	Tyr	Asp	Met	Arg	Gln	Ile	Met	Asn	Gly	Thr	Leu
Met	Lys	Asp	Asn	Pro	Val	Lys	Ala	Val	Thr	Leu	Val	Glu	Asn	His	Asp
Thr	Gln	Pro	Leu	Gln	Ala	Leu	Glu	Ser	Thr	Val	Asp	Trp	Trp	Phe	Lys
Pro	Leu	Ala	Tyr	Ala	Phe	Ile	Leu	Leu	Arg	Glu	Glu	Gly	Tyr	Pro	Ser
Val	Phe	Tyr	Ala	Asp	Tyr	Tyr	Gly	Ala	Gln	Tyr	Ser	Asp	Lys	Gly	Tyr
Asn	Ile	Asn	Met	Ala	Lys	Val	Pro	Tyr	Ile	Glu	Glu	Leu	Val	Thr	Leu
Arg	Lys	Glu	Tyr	Ala	Tyr	Gly	Lys	Gln	Asn	Ser	Tyr	Leu	Asp	His	Trp
Asp	Val	Ile	Gly	Trp	Thr	Arg	Glu	Gly	Asp	Ala	Glu	His	Pro	Asn	Ser
Met	Ala	Val	Ile	Met	Ser	Asp	Gly	Pro	Gly	Gly	Thr	Lys	Trp	Met	Tyr
Thr	Gly	Lys	Pro	Ser	Thr	Arg	Tyr	Val	Asp	Lys	Leu	Gly	Ile	Arg	Thr
Glu	Asp	Val	Trp	Thr	Asp	Ala	Asn	Gly	Trp	Ala	Glu	Phe	Pro	Val	Asn
Gly	Gly	Ser	Val	Ser	Val	Trp	Val	Gly	Val	Lys					

<210> 179
 <211> 1524
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

<400> 179
 atgaaaacat tcaaattaaa acgcactttt ttaccgctaa ccttgctgct cagtgtctct 60
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 ggcgcattat ggacgcagg tgaagcaat gctccagcac tcgctgaaaa cggttttaca 180
 gcgctctggc taccgccagc ttacaaaggc gcggcgcca gtaatgatgt cggttatggc 240

1001827087_1.txt

```

gtctacgata tgtacgattt aggtgagttt gatcaaaaag gctcagtagc aaccaaaatac 300
ggtaccaagg ctcagtagat ctcctgcaatc aatgctgctgc acaacaacaa tatccaaatt 360
tacggcgacg ttgtgtttta ccctcgtggg ggcgctgatg ggaagtcgtg ggtcgatacc 420
aagcgcgttg attgggacaa ccgtaacatt gaactgggag acaaattggat tgaagcttgg 480
gttgagttta attttcctag ccgcaacgac aaatactcga acttccattg gacttggtat 540
cactttgacg gtgttgactg ggatgatgcc ggcaaagaaa aagcgatctt taaattcaaa 600
ggcgaaggaa aagcatggga ttgggaagtc agctctgaaa aaggcaatta cgactaccta 660
atgtacgccg atttagacat ggatcaccca gaagttaaac aagagctgaa agattggggg 720
gagtgggtaca tcaacatgac cggcgttgat ggcttttaga tggatgccgt taagcacatt 780
aaatatcagt atctacaaga gtggattgat catttacgtt ggaaaacagg caaagagctt 840
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acctctggca gtatgtcgtt gttcgtatgc ccgcttcaca tgaacttcta caacgcgtca 960
aaatctggcg gcaattacga tatgcgccaa atcatgaatg gcacgttgat gaaggacaac 1020
ccagtcaaa cgtgtactct cgtagaaaa cagatagcgc agccattgca ggcgttagag 1080
tcgacagtgg attggtgggt caagcctctt gcttacgcac tcactttggt gcgtgaagaa 1140
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aacattaata tggccaaagt gccttacatt gaagaacttg taacactgcg taaagagtat 1260
gcgtatggca aacagaattc ttatctcgac cattgggatg tgattggctg gactcgagag 1320
ggcgtatgctg aacatccaaa ctcaatggcg gtgatcatga gtgatggacc gggcggaaca 1380
aaatggatgt ataccggtaa tccaagcacg cgctatgtcg acaagctggg tatccgaact 1440
gaagatgttt ggaccgatgc caatggctgg gcaagaatttc ctgtcaatgg tggttcagtc 1500
tcggtttggg tgggcgttaa gtaa 1524

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<210> 180

<211> 507

<212> PRT

<213> Unknown

<220>

<223> obtained from an environmental sample.

<400> 180

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Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu
1      5      10      15
Leu Ser Ala Pro Ala Phe Ala Gly Gln Asn Gly Thr Met Met Gln Tyr
20      25      30
Phe His Trp Tyr Val Pro Asn Asp Gly Ala Leu Trp Thr Gln Val Glu
35      40      45
Ser Asn Ala Pro Ala Leu Ala Glu Asn Gly Phe Thr Ala Leu Trp Leu
50      55      60
Pro Pro Ala Tyr Lys Gly Ala Gly Gly Ser Asn Asp Val Gly Tyr Gly
65      70      75      80
Val Tyr Asp Met Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ser Val
85      90      95
Arg Thr Lys Tyr Gly Thr Lys Ala Gln Tyr Ile Ser Ala Ile Asn Ala
100     105     110
Ala His Asn Asn Asn Ile Gln Ile Tyr Gly Asp Val Val Phe Asn His
115     120     125
Arg Gly Gly Ala Asp Gly Lys Ser Trp Val Asp Thr Lys Arg Val Asp
130     135     140
Trp Asp Asn Arg Asn Ile Glu Leu Gly Asp Lys Trp Ile Glu Ala Trp
145     150     155     160
Val Glu Phe Asn Phe Pro Ser Arg Asn Asp Lys Tyr Ser Asn Phe His
165     170     175
Trp Thr Trp Tyr His Phe Asp Gly Val Asp Trp Asp Asp Ala Gly Lys
180     185     190
Glu Lys Ala Ile Phe Lys Phe Lys Gly Glu Gly Lys Ala Trp Asp Trp
195     200     205
Glu Val Ser Ser Glu Lys Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp
210     215     220
Leu Asp Met Asp His Pro Glu Val Lys Gln Glu Leu Lys Asp Trp Gly
225     230     235
Glu Trp Tyr Ile Asn Met Thr Gly Val Asp Gly Phe Arg Met Asp Ala
245     250     255
Val Lys His Ile Lys Tyr Gln Tyr Leu Gln Glu Trp Ile Asp His Leu
260     265     270
Arg Trp Lys Thr Gly Lys Glu Leu Phe Thr Val Gly Glu Tyr Trp Asn
275     280     285

```

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Tyr Asp Val Asn Gln Leu His Asn Phe Ile Thr Lys Thr Ser Gly Ser
 290 295 300
 Met Ser Leu Phe Asp Ala Pro Leu His Met Asn Phe Tyr Asn Ala Ser
 305 310 315 320
 Lys Ser Gly Gly Asn Tyr Asp Met Arg Gln Ile Met Asn Gly Thr Leu
 325 330 335
 Met Lys Asp Asn Pro Val Lys Ala Val Thr Leu Val Glu Asn His Asp
 340 345 350
 Thr Gln Pro Leu Gln Ala Leu Glu Ser Thr Val Asp Trp Trp Phe Lys
 355 360 365
 Pro Leu Ala Tyr Ala Phe Ile Leu Leu Arg Glu Glu Gly Tyr Pro Ser
 370 375 380
 Val Phe Tyr Ala Asp Tyr Tyr Gly Ala Gln Tyr Ser Asp Lys Gly Tyr
 385 390 395 400
 Asn Ile Asn Met Ala Lys Val Pro Tyr Ile Glu Glu Leu Val Thr Leu
 405 410 415
 Arg Lys Glu Tyr Ala Tyr Gly Lys Gln Asn Ser Tyr Leu Asp His Trp
 420 425 430
 Asp Val Ile Gly Trp Thr Arg Glu Gly Asp Ala Glu His Pro Asn Ser
 435 440 445
 Met Ala Val Ile Met Ser Asp Gly Pro Gly Gly Thr Lys Trp Met Tyr
 450 455 460
 Thr Gly Asn Pro Ser Thr Arg Tyr Val Asp Lys Leu Gly Ile Arg Thr
 465 470 475 480
 Glu Asp Val Trp Thr Asp Ala Asn Gly Trp Ala Glu Phe Pro Val Asn
 485 490 495
 Gly Gly Ser Val Ser Val Trp Val Gly Val Lys
 500 505

<210> 181

<211> 1830

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 181

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ggcgtgtcgc	gcgggggcag	cctgccgatg	ccggtcctcc	actggccggc	gttcaccttc	120
gtccggcgct	tcgtcgccgg	tcattccgaac	aagcacaaga	accggagtat	tgcgatgagc	180
cacaccctgc	gtgccgccgt	attggcgggc	atcctgtctgc	cgttccccgc	cctcgctgac	240
caggccggca	agagcccggc	cggcgtgcgc	taccacggcg	gcgacgaaat	catcctccag	300
ggcttccact	ggaacgtcgt	ccgcgaagcg	cccaacgact	ggtacaacat	ccttcgccag	360
caggcctcga	cgatcgccgc	ggacggcttc	tcggcaatct	ggatgccggg	gccctggcgt	420
gacttctcca	gctggaccga	cggcggcaag	tcaggcggcg	gcgaaggcta	cttctggcac	480
gacttcaaca	agaacggccg	ctacggcagc	gacgcccagc	tgcgccaggc	cgccggcgca	540
ctcggtggcg	ccgggggtgaa	ggtgctctac	gatgtggtgc	ccaatcacat	gaaccgcggc	600
tatccggaca	aggagatcaa	cctgccggcc	ggccagggtc	tctggcgcaa	cgactgcacc	660
gacccgggca	actaccccaa	cgactgcgat	gacggtgacc	gcttcacatc	cggaagtcg	720
gacctgaaca	ccggccatcc	gcagatctac	ggcatgtttc	gcgacgagct	tgccaacctg	780
cgagcggggt	acggcgccgg	cggtctccgc	ttcgacttcg	ttcgcggtta	tgcgccgaa	840
cggttcgaca	gctggatgag	cgacagcgcc	gacagcagtt	tctgcgttgg	cgagctgtgg	900
aaaagcccg	ccgagtacc	gagctgggac	tggcgcaaca	cggcgagctg	gcagcagatc	960
atcaaggagt	ggtccgaccg	ggccaagtgc	ccggtgttcg	acttcgcgct	caaggagcgc	1020
atgcagaacg	gctcggctcg	cgactggaag	catggcctca	atggcaacc	ggaccgcgc	1080
tggcgcgagg	tggcgggtgac	ctttgtcgac	aaccacgaca	ccggctattc	gcccgggcag	1140
aacggcggcc	agcaccactg	ggcgctgcag	gacgggtga	tccgccaggc	ctacgcctac	1200
atcctcacca	gcccgggcac	gccgggtggtg	tactggctgc	acatgtacga	ctgggggtac	1260
ggcgacttca	ttcgccagct	gatccagggtg	cggcgcaccg	ctggcgtgcg	cgccgattcg	1320
gcgatcagct	tcacagcgg	ctacagcggc	ctggtcgcta	ccgtcagcgg	cagccatcag	1380
accctgggtg	tggcgctcaa	ctccgatctg	gccaaacccg	gccaggctgc	cagcggcagc	1440
ttcagcgagg	cggtcaacgc	cagcaacggc	cagggtgcgc	tctggcgcag	cggtagcggc	1500
gatggcggcg	gcaatgacgg	cggcgagggc	ggtctggtca	atgtgaactt	ccgctgcgac	1560
aacggcgtga	cgagatggg	cgacagcgtc	tacgcggtgg	gcaacgtcag	ccagctcggc	1620
aactggagcc	cggcctccgc	ggtacggctg	accgacacca	gcagctatcc	gacctggaag	1680
ggcagcatcg	ccctgcctga	cggtcagaa	gtggaaatga	agtgcctgat	ccgtaacgag	1740
gcggacgcga	cgctgggtgcg	ccagtggcaa	tcgggcggca	acaaccaggt	ccaggccgct	1800

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1830

gccggcgcgga gcaccagcgg ctcgttctga

<210> 182
<211> 609
<212> PRT
<213> Unknown

<220>
<223> obtained from an environmental sample.

<400> 182
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1 5 10 15
Gly Arg Leu Val Gly Val Ser Arg Gly Ser Leu Pro Met Pro Val
20 25 30
Leu His Trp Pro Ala Phe Ile Leu Val Arg Arg Phe Val Ala Gly His
35 40 45
Pro Asn Lys His Lys Asn Arg Ser Ile Ala Met Ser His Thr Leu Arg
50 55 60
Ala Ala Val Leu Ala Ala Ile Leu Leu Pro Phe Pro Ala Leu Ala Asp
65 70 75 80
Gln Ala Gly Lys Ser Pro Ala Gly Val Arg Tyr His Gly Gly Asp Glu
85 90 95
Ile Ile Leu Gln Gly Phe His Trp Asn Val Val Arg Glu Ala Pro Asn
100 105 110
Asp Trp Tyr Asn Ile Leu Arg Gln Ala Ser Thr Ile Ala Ala Asp
115 120 125
Gly Phe Ser Ala Ile Trp Met Pro Val Pro Trp Arg Asp Phe Ser Ser
130 135 140
Trp Thr Asp Gly Gly Lys Ser Gly Gly Gly Glu Gly Tyr Phe Trp His
145 150 155 160
Asp Phe Asn Lys Asn Gly Arg Tyr Gly Ser Asp Ala Gln Leu Arg Gln
165 170 175
Ala Ala Gly Ala Leu Gly Gly Ala Gly Val Lys Val Leu Tyr Asp Val
180 185 190
Val Pro Asn His Met Asn Arg Gly Tyr Pro Asp Lys Glu Ile Asn Leu
195 200 205
Pro Ala Gly Gln Gly Phe Trp Arg Asn Asp Cys Thr Asp Pro Gly Asn
210 215 220
Tyr Pro Asn Asp Cys Asp Asp Gly Asp Arg Phe Ile Gly Gly Lys Ser
225 230 235 240
Asp Leu Asn Thr Gly His Pro Gln Ile Tyr Gly Met Phe Arg Asp Glu
245 250 255
Leu Ala Asn Leu Arg Ser Gly Tyr Gly Ala Gly Gly Phe Arg Phe Asp
260 265 270
Phe Val Arg Gly Tyr Ala Pro Glu Arg Val Asp Ser Trp Met Ser Asp
275 280 285
Ser Ala Asp Ser Ser Phe Cys Val Gly Glu Leu Trp Lys Ser Pro Ser
290 295 300
Glu Tyr Pro Ser Trp Asp Trp Arg Asn Thr Ala Ser Trp Gln Gln Ile
305 310 315 320
Ile Lys Asp Trp Ser Asp Arg Ala Lys Cys Pro Val Phe Asp Phe Ala
325 330 335
Leu Lys Glu Arg Met Gln Asn Gly Ser Val Ala Asp Trp Lys His Gly
340 345 350
Leu Asn Gly Asn Pro Asp Pro Arg Trp Arg Glu Val Ala Val Thr Phe
355 360 365
Val Asp Asn His Asp Thr Gly Tyr Ser Pro Gly Gln Asn Gly Gly Gln
370 375 380
His His Trp Ala Leu Gln Asp Gly Leu Ile Arg Gln Ala Tyr Ala Tyr
385 390 395 400
Ile Leu Thr Ser Pro Gly Thr Pro Val Val Tyr Trp Ser His Met Tyr
405 410 415
Asp Trp Gly Tyr Gly Asp Phe Ile Arg Gln Leu Ile Gln Val Arg Arg
420 425 430
Thr Ala Gly Val Arg Ala Asp Ser Ala Ile Ser Phe His Ser Gly Tyr
435 440 445
Ser Gly Leu Val Ala Thr Val Ser Gly Ser His Gln Thr Leu Val Val

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450 455 460
 Ala Leu Asn Ser Asp Leu Ala Asn Pro Gly Gln Val Ala Ser Gly Ser
 465 470 475 480
 Phe Ser Glu Ala Val Asn Ala Ser Asn Gly Gln Val Arg Val Trp Arg
 485 490 495
 Ser Gly Ser Gly Asp Gly Gly Gly Asn Asp Gly Gly Glu Gly Gly Leu
 500 505 510
 Val Asn Val Asn Phe Arg Cys Asp Asn Gly Val Thr Gln Met Gly Asp
 515 520 525
 Ser Val Tyr Ala Val Gly Asn Val Ser Gln Leu Gly Asn Trp Ser Pro
 530 535 540
 Ala Ser Ala Val Arg Leu Thr Asp Thr Ser Ser Tyr Pro Thr Trp Lys
 545 550 555 560
 Gly Ser Ile Ala Leu Pro Asp Gly Gln Asn Val Glu Trp Lys Cys Leu
 565 570 575
 Ile Arg Asn Glu Ala Asp Ala Thr Leu Val Arg Gln Trp Gln Ser Gly
 580 585 590
 Gly Asn Asn Gln Val Gln Ala Ala Gly Ala Ser Thr Ser Gly Ser
 595 600 605
 Phe

<210> 183
 <211> 1596
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 183
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 gctctaacac tgccgctggc tgctagctta tcaacaggcg ttcacgccga aaccgtacat 120
 aaaggtaagt ctgaagcaac agataaaaac ggtgtctttt atgaggtgta tgtaaactct 180
 ttttacgata caaataaaga tggacatggg gatttaaaag gtctgacaca aaagttggat 240
 tatttaaatg acggcaattc tcatacaaag aatgatcttc aagtaaacgg gatttggatg 300
 atgccagtca acccttctcc tagctatcat aaatatgatg taacggacta ttataacatt 360
 gatcctcagt acggaaatct gcaagatttt cgcaagctga tgaaagaagc agacaaacga 420
 gacgtaaaag tcattatgga ccttgttgtg aatcatacga gcagcgaaca cccttggttt 480
 caagctgcat taaaagataa aaacagcaag tacagagatt actatatatt ggctgataaa 540
 aataccgatt tgaatgaaaa aggatcttg gggcagcaag tatggcataa agctccaaac 600
 ggagagtatt ttacggaac gttttgggaa ggaatgcctg acttaaatta cgataaccct 660
 gaagtaagaa aagaaatgat taacgtcggg aagttttggc taaagcaagg cgtaaattggc 720
 ttccgcttag atgctgcgct tcatattttt aaaggtcaaa cacctgaagg cgctaagaaa 780
 aatatcctgt ggtggaatga gttagagat gcgatgaaaa aagaaaacc taacgtatat 840
 ctaacgggtg aagtatggga tcagcctgaa gtggtagctc cttactatca atcgcttgat 900
 tctttattta attttgattt agcaggaaaa attgtcagct ctgtaaaagc aggaaatgat 960
 caaggaatcg ccactgcagc agcggcaaca gatgaactgt tcaaatcata caatccaaat 1020
 aaaattgacg gcattttctt aaccaaccat gaccaaaatc gcgtcatgag tgagctgagc 1080
 ggcgatgtga acaaaagcaaa atcagctgct tctatcttac ttacgcttcc tggcaaccgc 1140
 tatattttatt acggtgaaga aattggcatg accggtgaaa agcctgatga gttaatccgt 1200
 gaaccattcc gctggtacga aggaaacgga cttggacaaa ctagctggga aacacctgta 1260
 tataacaaag gcggcaacgg cgtgtctgta gaagtacaaa ccaaacaaaa ggattctttg 1320
 ttaaatcatt atcgtgaaat gattcgcgtg cgtcagcagc atgaagagtt agtaaaagga 1380
 acgcttcaat ctatttcagt agacagtaaa gaagtggttg cctatagtcg cacgtataaa 1440
 ggcaactcga ttagcgtgta tcataatatt tcaaatcaac ctgtaaaagt atctgtagca 1500
 gcgaaaggta aattgatttt tgctagttaa aaaggtgcta aaaaagtcaa aaatcagctt 1560
 gtaattccgg ctaatacaac ggttttaata aaataa 1596

<210> 184
 <211> 531
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 184

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Trp	Thr	Ala	Leu 20	Ala	Leu	Thr	Leu	Pro 25	Leu	Ala	Ala	Ser	Leu 30	Ser	Thr
Gly	Val	His 35	Ala	Glu	Thr	Val	His 40	Lys	Gly	Lys	Ser	Glu 45	Ala	Thr	Asp
Lys	Asn 50	Gly	Val	Phe	Tyr	Glu 55	Val	Tyr	Val	Asn 60	Ser	Phe	Tyr	Asp	Thr
Asn 65	Lys	Asp	Gly	His	Gly 70	Asp	Leu	Lys	Gly	Leu 75	Thr	Gln	Lys	Leu	Asp 80
Tyr	Leu	Asn	Asp	Gly 85	Asn	Ser	His	Thr	Lys 90	Asn	Asp	Leu	Gln	Val 95	Asn
Gly	Ile	Trp	Met 100	Met	Pro	Val	Asn	Pro 105	Ser	Pro	Ser	Tyr	His 110	Lys	Tyr
Asp	Val	Thr 115	Asp	Tyr	Tyr	Asn	Ile 120	Asp	Pro	Gln	Tyr	Gly 125	Asn	Leu	Gln
Asp	Phe 130	Arg	Lys	Leu	Met	Lys 135	Glu	Ala	Asp	Lys	Arg 140	Asp	Val	Lys	Val
Ile 145	Met	Asp	Leu	Val	Val 150	Asn	His	Thr	Ser	Ser 155	Glu	His	Pro	Trp	Phe 160
Gln	Ala	Ala	Leu	Lys 165	Asp	Lys	Asn	Ser	Lys 170	Tyr	Arg	Asp	Tyr	Tyr 175	Ile
Trp	Ala	Asp	Lys 180	Asn	Thr	Asp	Leu	Asn 185	Glu	Lys	Gly	Ser	Trp 190	Gly	Gln
Gln	Val	Trp 195	His	Lys	Ala	Pro	Asn 200	Gly	Glu	Tyr	Phe	Tyr 205	Gly	Thr	Phe
Trp	Glu 210	Gly	Met	Pro	Asp	Leu 215	Asn	Tyr	Asp	Asn	Pro 220	Glu	Val	Arg	Lys
Glu 225	Met	Ile	Asn	Val	Gly 230	Lys	Phe	Trp	Leu	Lys 235	Gln	Gly	Val	Asn	Gly 240
Phe	Arg	Leu	Asp	Ala 245	Ala	Leu	His	Ile	Phe 250	Lys	Gly	Gln	Thr	Pro 255	Glu
Gly	Ala	Lys	Lys 260	Asn	Ile	Leu	Trp	Trp 265	Asn	Glu	Phe	Arg	Asp 270	Ala	Met
Lys	Lys	Glu 275	Asn	Pro	Asn	Val	Tyr 280	Leu	Thr	Gly	Glu	Val 285	Trp	Asp	Gln
Pro	Glu 290	Val	Val	Ala	Pro	Tyr 295	Tyr	Gln	Ser	Leu	Asp 300	Ser	Leu	Phe	Asn
Phe 305	Asp	Leu	Ala	Gly	Lys 310	Ile	Val	Ser	Ser	Val 315	Lys	Ala	Gly	Asn	Asp 320
Gln	Gly	Ile	Ala	Thr 325	Ala	Ala	Ala	Ala	Thr 330	Asp	Glu	Leu	Phe	Lys 335	Ser
Tyr	Asn	Pro	Asn 340	Lys	Ile	Asp	Gly	Ile 345	Phe	Leu	Thr	Asn	His 350	Asp	Gln
Asn	Arg	Val 355	Met	Ser	Glu	Leu	Ser 360	Gly	Asp	Val	Asn	Lys 365	Ala	Lys	Ser
Ala	Ala 370	Ser	Ile	Leu	Leu	Thr 375	Leu	Pro	Gly	Asn	Pro 380	Tyr	Ile	Tyr	Tyr
Gly 385	Glu	Glu	Ile	Gly	Met 390	Thr	Gly	Glu	Lys	Pro 395	Asp	Glu	Leu	Ile	Arg 400
Glu	Pro	Phe	Arg	Trp 405	Tyr	Glu	Gly	Asn	Gly 410	Leu	Gly	Gln	Thr	Ser 415	Trp
Glu	Thr	Pro	Val 420	Tyr	Asn	Lys	Gly	Gly 425	Asn	Gly	Val	Ser	Val 430	Glu	Val
Gln	Thr	Lys 435	Gln	Lys	Asp	Ser	Leu 440	Leu	Asn	His	Tyr	Arg 445	Glu	Met	Ile
Arg	Val 450	Arg	Gln	Gln	His	Glu 455	Glu	Leu	Val	Lys	Gly 460	Thr	Leu	Gln	Ser
Ile 465	Ser	Val	Asp	Ser	Lys 470	Glu	Val	Val	Ala	Tyr 475	Ser	Arg	Thr	Tyr	Lys 480
Gly	Asn	Ser	Ile	Ser 485	Val	Tyr	His	Asn	Ile 490	Ser	Asn	Gln	Pro	Val 495	Lys
Val	Ser	Val	Ala 500	Ala	Lys	Gly	Lys	Leu 505	Ile	Phe	Ala	Ser	Glu 510	Lys	Gly
Ala	Lys	Lys 515	Val	Lys	Asn	Gln	Leu 520	Val	Ile	Pro	Ala	Asn 525	Thr	Thr	Val
Leu	Ile 530	Lys													

<210> 185
 <211> 1572
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 185
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aaaaacgggtg tcttttatga ggtgtatgta aactcttttt acgatgcaaa taaagatgga      180
catggtgatt taaaagggtc tacacaaaag ctggactatt taaatgacgg aaattctcat      240
acaaagaatg atcttcaagt aaacgggatt tggatgatgc cagtcaaccc ttctcctagc      300
tatcataaat atgatgtaac ggattattat aacattgatc cgcagtacgg aaatctgcaa      360
gattttcgca agctgatgaa agaagcagac aaacgagacg taaaagtcac tatggacctt      420
gttggtgaatc atacgagcag cgaacaccct tggtttcaag ctgctgttaa agataaaaac      480
agcaagtaca gagattacta ttttggggct gataaaaata ccgacttgaa tgaaaaagga      540
tcttggggag agcaagtatg gcataaagct ccaaacggag agtattttta cggaacgttt      600
tgggaaggaa tgcctgactt aaattacgat aacctggaag taagaaaaga aatgattaac      660
gtcggaaagt tttggctaaa gcaaggcggt gatggcttcc gcttagatgc tgcgcttcat      720
atttttaaag gtcaaagcc tgaaggcgct aagaaaaata ttctgtggtg gaatgagttt      780
agagatgcga tgaaaaaaga aaaccctaac gtatatctaa cgggtgaagt atgggatcag      840
cctgaagtgg tagctcctta ctatcaatcg cttgattccc tatttaactt tgatttagca      900
gggaaaattg tcagttctgt aaaagcagga aatgatcaag gaatcgccac tgcagcagcg      960
gcaacggatg agctgttcaa atcatacaat ccaaataaaa ttgacggcat tttcttaacc      1020
aaccatgacc aaaaccgcgt catgagtgaac ctgacggcg atgtgaacaa agcaaaatca      1080
gctgcttcta tcttacttac gcttcctggc aaccctgata tttattacgg tgaagaaatt      1140
ggcatgaccg gtgaaaagcc tgatgagtta atccgtgaac cgttccgctg gtacgaagga      1200
aacggacttg gacaaaccag ctgggaaaca cctgtatata acaaaggcgg caacggcggtg      1260
tctgtagaag cacaaaccaa acaaaaggat tctttgttaa atcattaccg tgaaatgatt      1320
cgctgtcgctc agcagcatga agagttagta aaaggaacgc ttcaatctat tttagtagac      1380
agtaaagaag ttgttgccca tagccgtacg tataaagaca actcgattag cgtgtatcat      1440
aatatttcaa atcaaccggt aaaagtatct gtagcagcaa aaggtaaatt aatttttgct      1500
agtgaaaaag gtgctaaaaa agtcaagaat cagcttgtga ttccggctaa tacaacggtt      1560
ttaataaaat aa                                     1572
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<210> 186
 <211> 523
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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<400> 186
Met Lys Leu Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr Leu
1      5      10      15
Pro Leu Ala Ala Ser Leu Ser Thr Gly Val His Ala Glu Thr Val His
20      25      30
Lys Gly Lys Ala Pro Thr Ala Asp Lys Asn Gly Val Phe Tyr Glu Val
35      40      45
Tyr Val Asn Ser Phe Tyr Asp Ala Asn Lys Asp Gly His Gly Asp Leu
50      55      60
Lys Gly Leu Thr Gln Lys Leu Asp Tyr Leu Asn Asp Gly Asn Ser His
65      70      75      80
Thr Lys Asn Asp Leu Gln Val Asn Gly Ile Trp Met Met Pro Val Asn
85      90      95
Pro Ser Pro Ser Tyr His Lys Tyr Asp Val Thr Asp Tyr Tyr Asn Ile
100      105      110
Asp Pro Gln Tyr Gly Asn Leu Gln Asp Phe Arg Lys Leu Met Lys Glu
115      120      125
Ala Asp Lys Arg Asp Val Lys Val Ile Met Asp Leu Val Val Asn His
130      135      140
Thr Ser Ser Glu His Pro Trp Phe Gln Ala Ala Leu Lys Asp Lys Asn
145      150      155      160
Ser Lys Tyr Arg Asp Tyr Tyr Ile Trp Ala Asp Lys Asn Thr Asp Leu
```

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Asn	Glu	Lys	Gly	165	Ser	Trp	Gly	Gln	Gln	170	Val	Trp	His	Lys	Ala	175	Pro	Asn
Gly	Glu	Tyr	Phe	180	Tyr	Gly	Thr	Phe	185	Trp	Glu	Gly	Met	Pro	Asp	Leu	Asn	
Tyr	Asp	Asn	Pro	195	Glu	Val	Arg	Lys	200	Glu	Met	Ile	Asn	Val	Gly	Lys	Phe	
Trp	Leu	Lys	Gln	210	Gly	Val	Asp	Gly	215	Phe	Arg	Leu	Asp	Ala	Ala	Leu	His	
Ile	Phe	Lys	Gly	225	Gln	Thr	Pro	Glu	230	Gly	Ala	Lys	Lys	Asn	Ile	Leu	Trp	
Trp	Asn	Glu	Phe	245	Arg	Asp	Ala	Met	250	Lys	Lys	Glu	Asn	Pro	Asn	Val	Tyr	
Leu	Thr	Gly	Glu	260	Val	Trp	Asp	Gln	265	Pro	Glu	Val	Val	Ala	Pro	Tyr	Tyr	
Gln	Ser	Leu	Asp	275	Ser	Leu	Phe	Asn	280	Phe	Asp	Leu	Ala	Gly	Lys	Ile	Val	
Ser	Ser	Val	Lys	290	Ala	Gly	Asn	Asp	295	Gln	Gly	Ile	Ala	Thr	Ala	Ala	Ala	
Ala	Thr	Asp	Glu	305	Leu	Phe	Lys	Ser	310	Tyr	Asn	Pro	Asn	Lys	Ile	Asp	Gly	
Ile	Phe	Leu	Thr	325	Asn	His	Asp	Gln	330	Asn	Arg	Val	Met	Ser	Glu	Leu	Ile	
Gly	Asp	Val	Asn	340	Lys	Ala	Lys	Ser	345	Ala	Ala	Ser	Ile	Leu	Leu	Thr	Leu	
Pro	Gly	Asn	Pro	355	Tyr	Ile	Tyr	Gly	360	Glu	Glu	Glu	Ile	Gly	Met	Thr	Gly	
Glu	Lys	Pro	Asp	370	Glu	Leu	Ile	Arg	375	Glu	Pro	Phe	Arg	Trp	Tyr	Glu	Gly	
Asn	Gly	Leu	Gly	385	Gln	Thr	Ser	Trp	390	Glu	Thr	Pro	Val	Tyr	Asn	Lys	Gly	
Gly	Asn	Gly	Val	405	Ser	Val	Glu	Ala	410	Gln	Thr	Lys	Gln	Lys	Asp	Ser	Leu	
Leu	Asn	His	Tyr	420	Arg	Glu	Met	Ile	425	Arg	Val	Arg	Gln	Gln	His	Glu	Glu	
Leu	Val	Lys	Gly	435	Thr	Leu	Gln	Ser	440	Ile	Leu	Val	Asp	Ser	Lys	Glu	Val	
Val	Ala	Tyr	Ser	450	Arg	Thr	Tyr	Lys	455	Asp	Asn	Ser	Ile	Ser	Val	Tyr	His	
Asn	Ile	Ser	Asn	465	Gln	Pro	Val	Lys	470	Val	Ser	Val	Ala	Ala	Lys	Gly	Lys	
Leu	Ile	Phe	Ala	485	Ser	Glu	Lys	Gly	490	Ala	Lys	Lys	Val	Lys	Asn	Gln	Leu	
Val	Ile	Pro	Ala	500	Asn	Thr	Thr	Val	505	Leu	Ile	Lys			510			
				515					520									

<210> 187

<211> 2052

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 187

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gacttgccgg	ttcttgccga	attctgcaaa	aaagccggat	ttgatcttgt	acagcttctt	180
ccggtcaatg	acaccggcac	agaaagttct	ccatacagcg	cgctttctgc	ctttgccctg	240
cacccgctgt	atatcaggct	ttccgacctg	cctgaagcag	cggttttcga	aaagcagatt	300
acagatctga	aaagccggtt	tgaggacttg	cctcgtttca	gctatacgga	gctgcgccgt	360
gccaaactgg	atatactgcg	tgcagtgttt	gataaaaaca	aggcaaccat	catcggcagt	420
gccgaactgg	aagcctggat	ttcagataac	ccctggatca	tcgaatatgc	ggtttttatg	480
aaccagaaac	accgcaactt	tgaagccggc	tggaaacatt	gggaaaagct	gcgcaacccc	540
actcataacg	aaatacaaaa	aacctggcag	ggtaaaacct	ggcaggctga	ccatcaattc	600
tttgcattgg	tgcagatgcg	gctggaccag	cagtttactg	ccgccgttac	agagtgaac	660
gccctgggtg	tctatcttaa	gggcgatata	cctataatga	tgaacgagga	ttccgcagat	720
gcctgggcga	atccggaatt	cttccgtgac	gatcttcggg	cgggaagtcc	ccctgacggt	780

1001827087_1.txt

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gaaaaccccc agggacaaaa ctggggcttc cccatttata actgggaaaa ccttgcaaat 840
gacgggtaca gctgggtgaa aaaacgtctg aagcacagcg cacgggtatta ccatgcctac 900
cgattgacc atattcttgg gtttttccgg atatgggcta taccctatgg cgaatactcc 960
ggctacctgg gatggccctt gccgcatgaa ccggtaagtg cagcagaact ggcagaacgg 1020
ggcttttcca aggaccgctt gcgctggctt accgaacccc acttgccctac acgggcagcc 1080
gaggaagcga ataactggga ctatctggga acacacggct atctgaatca gatcatgaac 1140
cgatcgggtg aagaagaact atggctgttc aagcccgaga tcacctgcga ggcagatata 1200
cgaaacacaa acctgccgga tgccctgaaa gaggttcttg tacggcagtg gaaaaaccgg 1260
ctgctgcagg ttaccggccg cgacgaaaaa ggacggacaa tctactatcc gctgtggcgt 1320
ttccgtgaca gcactgcatg gcagacgctt accgatggcg agaaacactc cctggaagag 1380
ctgttcgccc aaaaagcggc gcacaatgaa accctgtggc gagaacaggc ggtggaactt 1440
ctgggtgagc tgacgcgatc tacggatatg cttgcctgtg ctgaagatct gggaagtatt 1500
ccccacagtg taccggaagt gctttcaaac ctttcaattt acagtctgcg ggttaccgcg 1560
tgggcccgcg aatgggatgc ccccgccag ccctttcaca gactggagga gtatccgctc 1620
atgtcggtag cgaccccatc ggttcatgat tcctctacc tgcgcggatg gtgggaaacc 1680
gaaggcggcg accgggcctt tatggacgca tggcctccgg aacaggatgc atacgcagga 1740
gcaggccgcc atgagttcga aggcgcctgg ggaccccgcc aggcattctg ggtactccgt 1800
aaactctgcg aagcccgttc cgcgctctgt gttttcccca tccaggatat tttggccctg 1860
tcttcagact tttatgcaat gacagcggag gaggaacgca tcaatatcc gggcagtgta 1920
tccggattta actggacata ccggttgctt gcggcaatcg aggatttatc taaaaacagc 1980
caacttataa ccgcaatcca gaccgcgttg caggaccgcc ggcgaggaag ggcacagga 2040
gcacagcaat ga 2052

```

<210> 188
 <211> 683
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

<400> 188
 Met Tyr Leu Ile Gln Glu Gly His Met Arg Phe Pro Pro Ile Ile His
 1 5 10 15
 Pro Leu Thr Gly Leu Ala Val Pro Val Gly Ala Leu Arg Thr Ala Gln
 20 25 30
 Ser Cys Gly Ile Gly Glu Phe Ala Asp Leu Pro Val Leu Ala Glu Phe
 35 40 45
 Cys Lys Lys Ala Gly Phe Asp Leu Val Gln Leu Leu Pro Val Asn Asp
 50 55 60
 Thr Gly Thr Glu Ser Ser Pro Tyr Ser Ala Leu Ser Ala Phe Ala Leu
 65 70 75 80
 His Pro Leu Tyr Ile Arg Leu Ser Asp Leu Pro Glu Ala Ala Gly Phe
 85 90 95
 Glu Lys Gln Ile Thr Asp Leu Lys Ser Arg Phe Glu Asp Leu Pro Arg
 100 105 110
 Phe Ser Tyr Thr Glu Leu Arg Arg Ala Lys Leu Asp Ile Leu Arg Ala
 115 120 125
 Val Phe Asp Lys Asn Lys Ala Thr Ile Ile Gly Ser Ala Glu Leu Glu
 130 135 140
 Ala Trp Ile Ser Asp Asn Pro Trp Ile Ile Glu Tyr Ala Val Phe Met
 145 150 155 160
 Asn Gln Lys His Arg Asn Phe Glu Ala Gly Trp Lys His Trp Glu Lys
 165 170 175
 Leu Arg Asn Pro Thr His Asn Glu Ile Gln Lys Thr Trp Gln Gly Lys
 180 185 190
 Thr Trp Gln Ala Asp His Gln Phe Phe Ala Trp Leu Gln Met Arg Leu
 195 200 205
 Asp Gln Gln Phe Thr Ala Ala Ala Thr Glu Cys Asn Ala Leu Gly Val
 210 215 220
 Tyr Leu Lys Gly Asp Ile Pro Ile Met Met Asn Glu Asp Ser Ala Asp
 225 230 235 240
 Ala Trp Ala Asn Pro Glu Phe Phe Arg Asp Asp Leu Arg Ala Gly Ser
 245 250 255
 Pro Pro Asp Gly Glu Asn Pro Gln Gly Gln Asn Trp Gly Phe Pro Ile
 260 265 270
 Tyr Asn Trp Glu Asn Leu Ala Asn Asp Gly Tyr Ser Trp Lys Lys
 275 280 285

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Arg Leu Lys His Ser Ala Arg Tyr Tyr His Ala Tyr Arg Ile Asp His
 290 305 310 315 330 335 340 345 350 355 360 365 370 375 380 385 390 400 405 410 415 420 425 430 435 440 445 450 455 460 465 470 475 480 485 490 495 500 505 510 515 520 525 530 535 540 545 550 555 560 565 570 575 580 585 590 595 600 605 610 615 620 625 630 635 640 645 650 655 660 665 670 675 680

Ile Leu Gly Phe Phe Arg Ile Trp Ala Ile Pro Tyr Gly Glu Tyr Ser
 Gly Tyr Leu Gly Trp Pro Leu Pro His Glu Pro Val Ser Ala Ala Glu
 Leu Ala Glu Arg Gly Phe Ser Lys Asp Arg Leu Arg Trp Leu Thr Glu
 Pro His Leu Pro Thr Arg Ala Ala Glu Glu Ala Asn Asn Trp Asp Tyr
 Leu Gly Thr His Gly Tyr Leu Asn Gln Ile Met Asn Arg Ile Gly Glu
 Glu Glu Leu Trp Leu Phe Lys Pro Glu Ile Thr Cys Glu Ala Asp Ile
 Arg Asn Thr Asn Leu Pro Asp Ala Leu Lys Glu Val Leu Val Arg Gln
 Trp Lys Asn Arg Leu Leu Gln Val Thr Gly Arg Asp Glu Lys Gly Arg
 Thr Ile Tyr Tyr Pro Leu Trp Arg Phe Arg Asp Ser Thr Ala Trp Gln
 Thr Leu Thr Asp Gly Glu Lys His Ser Leu Glu Glu Leu Phe Ala Gln
 Lys Ala Ala His Asn Glu Thr Leu Trp Arg Glu Gln Ala Val Glu Leu
 Leu Gly Glu Leu Thr Arg Ser Thr Asp Met Leu Ala Cys Ala Glu Asp
 Leu Gly Ser Ile Pro His Ser Val Pro Glu Val Leu Ser Asn Leu Ser
 Ile Tyr Ser Leu Arg Val Thr Arg Trp Ala Arg Gln Trp Asp Ala Pro
 Gly Gln Pro Phe His Arg Leu Glu Glu Tyr Pro Leu Met Ser Val Ala
 Thr Pro Ser Val His Asp Ser Ser Thr Leu Arg Gly Trp Trp Glu Thr
 Glu Gly Gly Asp Arg Ala Phe Met Asp Ala Trp Pro Pro Glu Gln Asp
 Ala Tyr Ala Gly Ala Gly Arg His Glu Phe Glu Gly Ala Trp Gly Pro
 Arg Gln Ala Ser Trp Val Leu Arg Lys Leu Cys Glu Ala Arg Ser Ala
 Leu Cys Val Phe Pro Ile Gln Asp Ile Leu Ala Leu Ser Ser Asp Phe
 Tyr Ala Met Thr Ala Asp Glu Glu Arg Ile Asn Ile Pro Gly Ser Val
 Ser Gly Phe Asn Trp Thr Tyr Arg Leu Pro Ala Ala Ile Glu Asp Leu
 Ser Lys Asn Ser Gln Leu Ile Thr Ala Ile Gln Thr Ala Leu Gln Asp
 Arg Arg Ala Arg Lys Ala Gln Gly Ala Gln Gln

<210> 189
 <211> 1596
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 189
 atgcaaacga ttgcaaaaaa aggggatgaa acgatgaaag ggaaaaaatg gacagcttta 60
 gctctaacac tgccgctggc tgctagctta tcaacaggcg ttcacgccga aaccgtacat 120
 aaaggtaaatt ctccagctgc agataaaaac ggtgtctttt atgaggtgta tgtaaactct 180
 ttttacgatg caaataaaga tggacatggg gatttaaaag gtcttacaca aaaactggac 240
 tatttaaatg atggcaattc tcatacaaag aatgatcttc aagtaaaccg gatttggatg 300
 atgccgatca acccttctcc tagctatcat aaatatgatg taacggacta ttataacatt 360
 gattctcagt acggaaatct gcaagatttt cgcaagctaa tgaaagaagc agataaacga 420
 gatgtaaaag ttattatgga cctcgttgtg aatcatacga gcagtgaaca cccttggttt 480

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```

caagctgCGT taaaagataa aaacagcaag tacagagatt actatatTTG ggctgataaa 540
aataccgatt tgaatgaaaa aggatcTTGg ggacaacaag tatggcacaa agctccaaac 600
ggagagTatt tttacggaac gttctgggaa ggaatgcctg acttaaatta cgataaccct 660
gaagtaagaa aagaaatgat taacgtcgga aagttttggc taaagcaagg cgttgacggc 720
ttccgcttag atgctgccct tcatatcttt aaaggTcaaa cacctgaagg cgctaagaaa 780
aatattgtgt ggtggaatga atttagagat gcgatgaaaa aagaaaaccg gaacgtatat 840
ctaacgggCG aagtatggga tcagccggaa gtggtagctc cttattatca gtcgcttgat 900
tccctatTTa actttgattt agcaggaaaa attgtcagct ctgtaaaagc aggaaatgat 960
caaggaatCG ctactgcagc agcggcaaca gatgaactgt tcaaatcata caatccaaat 1020
aaaattgacg gcattttctt aaccaatcat gaccaaattc gcgtcatgag tgagttaagc 1080
ggagatgtca ataaagcaaa gtcagctgcc tctatcttac ttacgcttcc tggaaatccg 1140
tatatttatt acggtgaaga aatcggcatg accggtgaaa agcctgatga attaatccgt 1200
gaaccgttcc gctggtacga aggaaacgga cttggacaaa ctagtTggga aacacctgta 1260
tacaataaaG gcggcaacgg cgtgtctgta gaagcacaaa ccaaacaaaa ggactcTTtg 1320
ttaaatcatt accgtgaaat gattcgcgtg cgtcagcagc acgaagagtt agtaaaagga 1380
acgcttcaat ctatttcagt agacagtaaa gaagtTgttg cttatagccg tacgtataaa 1440
ggcaactcca ttagtgtgta tcataatatt tcaaatcaac ctgtaaaagt atctgtagca 1500
gcgaaaggta aattgatttt tgctagtgaa aaaggTgcta aaaaggTcaa aaatcagctt 1560
gtgattccgg cgaatacaac ggttttagta aaataa 1596

```

<210> 190
 <211> 531
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 190

```

Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
1      5      10      15
Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr
20      25      30
Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Ala Ala Asp
35      40      45
Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala
50      55      60
Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp
65      70      75      80
Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn
85      90      95
Gly Ile Trp Met Met Pro Ile Asn Pro Ser Pro Ser Tyr His Lys Tyr
100     105     110
Asp Val Thr Asp Tyr Tyr Asn Ile Asp Ser Gln Tyr Gly Asn Leu Gln
115     120     125
Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val
130     135     140
Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
145     150     155     160
Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile
165     170     175
Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp Gly Gln
180     185     190
Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe
195     200     205
Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys
210     215     220
Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly
225     230     235     240
Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Pro Glu
245     250     255
Gly Ala Lys Lys Asn Ile Val Trp Trp Asn Glu Phe Arg Asp Ala Met
260     265     270
Lys Lys Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln
275     280     285
Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn
290     295     300
Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp

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```

305      310      315      320
Gln Gly Ile Ala Thr Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser
      325      330      335
Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His Asp Gln
      340      345      350
Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Asn Lys Ala Lys Ser
      355      360      365
Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr
      370      375      380
Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg
385      390      395      400
Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr Ser Trp
      405      410      415
Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala
      420      425      430
Gln Thr Lys Gln Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile
      435      440      445
Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser
      450      455      460
Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys
465      470      475      480
Gly Asn Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys
      485      490      495
Val Ser Val Ala Ala Lys Gly Lys Leu Ile Phe Ala Ser Glu Lys Gly
      500      505      510
Ala Lys Lys Val Lys Asn Gln Leu Val Ile Pro Ala Asn Thr Thr Val
      515      520      525
Leu Val Lys
530

```

<210> 191
 <211> 1596
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

```

<400> 191
atgcaaacga ttgcaaaaaa aggggatgaa acgatgaaag ggaaaaaatg gacagcttta 60
gc tctaacac tgccgctggc tgctagctta tcaacaggcg ttcacgccga aaccgtacat 120
aaaggtaaat ctccaacagc agataaaaac ggtgtctttt atgaagtgtg tgtaaaactct 180
ttttacgatg caaataaaga tggacatggg gacttaaaag gtcttacaca aaagttggac 240
tatttaaatg acggcaattc tcatacaaaa aatgatcttc aagtaaacgg gatttggaag 300
atgccagtca acccttctcc tagctatcat aaatatgatg taacggacta ttataacatt 360
gatccgcagt acggaaatct gcaagatttt cgcaagctga tgaaagaagc agacaaacga 420
gacgtaaaaa tcattatgga ccttgttgtg aatcatatcg gcagtgaaca cccttggttt 480
caagctgcgt taaaagataa aaacagcaag tacagagatt actatatatt ggctgataaa 540
aataccgact tgaatgaaaa aggatcttgg ggacaacaag tatggcataa agctccaaac 600
ggagagtatt ttacggaac gttctgggaa ggaatgcctg acttaaatga cgataaccct 660
gaagtaagaa aagaaatgat taacgtcggg aagttttggc taaagcaagg cgttgacggg 720
ttccgcttag atgctgcgct tcatattttt aaagggtcaa cagctgaagg cgctaagaaa 780
aatatcctgt ggtggaatga gtttagagat gcgatgaaaa aagaaaatcc gaatgtatat 840
ctaacgggtg aagtatggga tcagcctgaa gtggtagctc cttattatca atcgcttgat 900
tctttattta attttgattt agcaggaaaa attgtcagct ctgtaaaagc aggaaatgat 960
caaggaaatc ccactgcagc agcagcaaca gatgaactgt tcaaatcata caatccaaac 1020
aaaattgatg gcataattctt aaccaaccat gaccaaaatc gcgtcatgag tgagctgagc 1080
ggcgaatgta gcaaagcaaa atcagctgct tctatcttac ttacgttcc tggcaaccgc 1140
tatatttatt acggtgaaga aatcggcatg accggtgaaa agcctgatga attaatccgt 1200
gaaccgttcc gctggtacga aggaaacgga cttggacaaa ccagttggga aacacctgta 1260
tacaataaag ccggaacggg tgtgtctgta gaagcacaac ccaaacaaaa ggattctttg 1320
ttaaatcatt accgtgaaat gattcgcgtg cgtcagcagc atgaagagtt agtaaaagga 1380
acgcttcaat ctatttcagt agacagtaaa gaagttgttg cttatagccg tacgtataaa 1440
ggcaactcca ttagtgtgta tcataatatt tcaaatcaac cggtaaaagt atctgtagca 1500
gcgaaaggta aattgatttt tgctagttaa aaaggtgcta agaaagtcaa aaatcagctt 1560
gtggttccgg cgaatacaac ggttttaaat aaataa 1596

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<210> 192

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<211> 531
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 192
Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
1 5 10 15
Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr
20 25 30
Gly Val His Ala Glu Thr Val His Lys Gly Lys Ser Pro Thr Ala Asp
35 40 45
Lys Asn Gly Val Phe Tyr Glu Val Tyr Val Asn Ser Phe Tyr Asp Ala
50 55 60
Asn Lys Asp Gly His Gly Asp Leu Lys Gly Leu Thr Gln Lys Leu Asp
65 70 75 80
Tyr Leu Asn Asp Gly Asn Ser His Thr Lys Asn Asp Leu Gln Val Asn
85 90 95
Gly Ile Trp Met Met Pro Val Asn Pro Ser Pro Ser Tyr His Lys Tyr
100 105 110
Asp Val Thr Asp Tyr Tyr Asn Ile Asp Pro Gln Tyr Gly Asn Leu Gln
115 120 125
Asp Phe Arg Lys Leu Met Lys Glu Ala Asp Lys Arg Asp Val Lys Val
130 135 140
Ile Met Asp Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe
145 150 155 160
Gln Ala Ala Leu Lys Asp Lys Asn Ser Lys Tyr Arg Asp Tyr Tyr Ile
165 170 175
Trp Ala Asp Lys Asn Thr Asp Leu Asn Glu Lys Gly Ser Trp Gly Gln
180 185 190
Gln Val Trp His Lys Ala Pro Asn Gly Glu Tyr Phe Tyr Gly Thr Phe
195 200 205
Trp Glu Gly Met Pro Asp Leu Asn Tyr Asp Asn Pro Glu Val Arg Lys
210 215 220
Glu Met Ile Asn Val Gly Lys Phe Trp Leu Lys Gln Gly Val Asp Gly
225 230 235 240
Phe Arg Leu Asp Ala Ala Leu His Ile Phe Lys Gly Gln Thr Ala Glu
245 250 255
Gly Ala Lys Lys Asn Ile Leu Trp Trp Asn Glu Phe Arg Asp Ala Met
260 265 270
Lys Lys Glu Asn Pro Asn Val Tyr Leu Thr Gly Glu Val Trp Asp Gln
275 280 285
Pro Glu Val Val Ala Pro Tyr Tyr Gln Ser Leu Asp Ser Leu Phe Asn
290 295 300
Phe Asp Leu Ala Gly Lys Ile Val Ser Ser Val Lys Ala Gly Asn Asp
305 310 315 320
Gln Gly Ile Ala Thr Ala Ala Ala Ala Thr Asp Glu Leu Phe Lys Ser
325 330 335
Tyr Asn Pro Asn Lys Ile Asp Gly Ile Phe Leu Thr Asn His Asp Gln
340 345 350
Asn Arg Val Met Ser Glu Leu Ser Gly Asp Val Ser Lys Ala Lys Ser
355 360 365
Ala Ala Ser Ile Leu Leu Thr Leu Pro Gly Asn Pro Tyr Ile Tyr Tyr
370 375 380
Gly Glu Glu Ile Gly Met Thr Gly Glu Lys Pro Asp Glu Leu Ile Arg
385 390 395 400
Glu Pro Phe Arg Trp Tyr Glu Gly Asn Gly Leu Gly Gln Thr Ser Trp
405 410 415
Glu Thr Pro Val Tyr Asn Lys Gly Gly Asn Gly Val Ser Val Glu Ala
420 425 430
Gln Thr Lys Gln Lys Asp Ser Leu Leu Asn His Tyr Arg Glu Met Ile
435 440 445
Arg Val Arg Gln Gln His Glu Glu Leu Val Lys Gly Thr Leu Gln Ser
450 455 460
Ile Ser Val Asp Ser Lys Glu Val Val Ala Tyr Ser Arg Thr Tyr Lys
465 470 475 480

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Gly Asn Ser Ile Ser Val Tyr His Asn Ile Ser Asn Gln Pro Val Lys
 485 490 495
 Val Ser Val Ala Lys Gly Lys Leu Ile Phe Ala Ser Glu Lys Gly
 500 505 510
 Ala Lys Lys Val Lys Asn Gln Leu Val Val Pro Ala Asn Thr Thr Val
 515 520 525
 Leu Met Lys
 530

<210> 193
 <211> 1962
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 193
 atgaaattca aaaagagttt atctgcccgg ctccttttgt tcggagggtc gagcgggtgtg 60
 acaccatccg tcgctgcgga ggtgccacga accgcatttg tccattttatt cgaatggagt 120
 tggcccgata ttgccaccga atgcgaaacc tttcttggtc ctaaggggtt ctctgcggtt 180
 cagggtgtctc cgccgcaaaa aagcgtcagc aatgctgcct ggtggggcgc ctaccaacct 240
 gttagtactt cttttgaagg gcgcagtgga acccgggctc aatttgcgga tatggtccag 300
 cgttgtaaaag cgggtgggggt cgatatttat ctggatgcgg tgatcaacca tatggcagca 360
 caagatcgct attttcacga agtaccttac agcagtaatg attttcacag ttgcacgggc 420
 gatatcgatt attccaaccg ctggtcgatt caaaattgcg atctggttgg gctgaacgat 480
 ctcaaaaaccg agtcagaata cgttcggcag aaaattgcag actatatgaa cgatgcgctc 540
 agtctgggag tggcgggggt tcggattgat gccgccaagc atatcccggc cggcgacatc 600
 gcggcgatca agagcaagct caacggcagc ccgtatatct atcaggaggt tatcggggcg 660
 gcaggggagc cggtaacaac cagcgagtac acgtatatgt gagacgtgac ggaatttaac 720
 ttcgcccgga ccatcgggcc taaatttaag caaggttaata ttaaagacct gcaggggatt 780
 ggttcgtgga gcggctggct gagcagcgac gatcggtga cctttgtgac caaccatgac 840
 gaagaacgcc ataaccctgg ccaggttctc agccatcagg actttggcaa tctgtatttc 900
 ctcggttaacg tgtttactct ggcgatctct tacggctacc caaaagtgat gtcggggtag 960
 tacttcagta attttgatgc cgggccacca tcgacagggg tacattctgg taatgcgtgt 1020
 ggctttgatg gcggtgattg ggtctgcgaa cacaaatggc gtggtgtagc caacatggtg 1080
 gcgtttcgca accacacagc agcccagtgg caggtcactg actggtggga cgatggttac 1140
 aatcaggtgg cgtttggtcg tggcgggctg ggctttgtgg tgatcaatcg agatgacaat 1200
 aaaggcatca atcagagttt ccagacggga atgcccgtg gcgagtattg tgacatcatt 1260
 gccggtgatt tcgacacca gagcggatcat tgcagcgcta cgacgatcac cgtcgacagt 1320
 caggggtatg cacattttac tgtcggtagt catcaggccg ctgcgattca cattggcgcg 1380
 aaactcggct ccgtgtgccg ggactgtggc ggcacggccg cagagacaaa agtctgcttt 1440
 gacaatgcac aaaactttag ccaaccgtat ttgcattact ggaatgtcaa tgcggatcag 1500
 gccgtagcga atgcaacctg gccgggcgtc gcgatgacgg ctgaaaatgg cggttactgc 1560
 tacgattttg gtgtcggtct caattcactt caggttaatt tcagcgataa cggcgccagc 1620
 caaacgcgtg atctgaccgc cagcagtcct acgttgtgtt accagaacgg aacgtggcgt 1680
 gacagtgact tctgtcagag tagcaatgtg ggcaacgaga gttggtattt ccgtggaacc 1740
 tcaaacgggt ggggcgtgag cgcactcact tatgaggctg cgacaggcct gtacactacg 1800
 gtgcagagct ttaacgggga ggagtcgccc gcacgcttta aaattgatga tggcaactgg 1860
 agtgagtcgt atccaagtgc tgattatcaa gtcggtgatt atgccaccta cacgatcacg 1920
 tttgacagcc agacgaaggc catcaccgtg acttcgcagt aa 1962

<210> 194
 <211> 653
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 194
 Met Lys Phe Lys Lys Ser Leu Ser Ala Gly Leu Leu Leu Phe Gly Gly
 1 5 10 15
 Leu Ser Gly Val Thr Pro Ser Val Ala Ala Glu Val Pro Arg Thr Ala
 20 25 30
 Phe Val His Leu Phe Glu Trp Ser Trp Pro Asp Ile Ala Thr Glu Cys
 35 40 45
 Glu Thr Phe Leu Gly Pro Lys Gly Phe Ser Ala Val Gln Val Ser Pro

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Pro 65	50 Gln	Lys	Ser	Val	Ser 70	55 Asn	Ala	Ala	Trp	Trp 75	60 Ala	Arg	Tyr	Gln	Pro 80
Val	Ser	Tyr	Ser	Phe 85	Glu	Gly	Arg	Ser	Gly 90	Thr	Arg	Ala	Gln	Phe 95	Ala
Asp	Met	Val	Gln 100	Arg	Cys	Lys	Ala	Val 105	Gly	Val	Asp	Ile	Tyr 110	Leu	Asp
Ala	Val	Ile 115	Asn	His	Met	Ala	Ala 120	Gln	Asp	Arg	Tyr	Phe 125	Pro	Glu	Val
Pro	Tyr 130	Ser	Ser	Asn	Asp	Phe 135	His	Ser	Cys	Thr	Gly 140	Asp	Ile	Asp	Tyr
Ser 145	Asn	Arg	Trp	Ser	Ile 150	Gln	Asn	Cys	Asp	Leu 155	Val	Gly	Leu	Asn	Asp 160
Leu	Lys	Thr	Glu	Ser 165	Glu	Tyr	Val	Arg	Gln 170	Lys	Ile	Ala	Asp	Tyr 175	Met
Asn	Asp	Ala	Leu 180	Ser	Leu	Gly	Val	Ala 185	Gly	Phe	Arg	Ile	Asp 190	Ala	Ala
Lys	His	Ile 195	Pro	Ala	Gly	Asp	Ile 200	Ala	Ala	Ile	Lys	Ser 205	Lys	Leu	Asn
Gly	Ser 210	Pro	Tyr	Ile	Tyr	Gln 215	Glu	Val	Ile	Gly	Ala 220	Ala	Gly	Glu	Pro
Val 225	Gln	Thr	Ser	Glu	Tyr 230	Thr	Tyr	Ile	Gly	Asp 235	Val	Thr	Glu	Phe	Asn 240
Phe	Ala	Arg	Thr	Ile 245	Gly	Pro	Lys	Phe	Lys 250	Gln	Gly	Asn	Ile	Lys 255	Asp
Leu	Gln	Gly	Ile 260	Gly	Ser	Trp	Ser	Gly 265	Trp	Leu	Ser	Ser	Asp 270	Asp	Ala
Val	Thr	Phe 275	Val	Thr	Asn	His	Asp 280	Glu	Glu	Arg	His	Asn 285	Pro	Gly	Gln
Val	Leu 290	Ser	His	Gln	Asp	Phe 295	Gly	Asn	Leu	Tyr	Phe 300	Leu	Gly	Asn	Val
Phe 305	Thr	Leu	Ala	Tyr	Pro 310	Tyr	Gly	Tyr	Pro	Lys 315	Val	Met	Ser	Gly	Tyr 320
Tyr	Phe	Ser	Asn	Phe 325	Asp	Ala	Gly	Pro	Pro 330	Ser	Thr	Gly	Val	His 335	Ser
Gly	Asn	Ala	Cys 340	Gly	Phe	Asp	Gly	Gly 345	Asp	Trp	Val	Cys	Glu 350	His	Lys
Trp	Arg	Gly 355	Val	Ala	Asn	Met	Val 360	Ala	Phe	Arg	Asn	His 365	Thr	Ala	Ala
Gln	Trp 370	Gln	Val	Thr	Asp	Trp 375	Trp	Asp	Asp	Gly	Tyr 380	Asn	Gln	Val	Ala
Phe 385	Gly	Arg	Gly	Gly	Leu 390	Gly	Phe	Val	Val	Ile 395	Asn	Arg	Asp	Asp	Asn 400
Lys	Gly	Ile	Asn	Gln 405	Ser	Phe	Gln	Thr	Gly 410	Met	Pro	Ala	Gly	Glu 415	Tyr
Cys	Asp	Ile	Ile 420	Ala	Gly	Asp	Phe	Asp 425	Thr	Gln	Ser	Gly	His 430	Cys	Ser
Ala	Thr	Thr 435	Ile	Thr	Val	Asp	Ser 440	Gln	Gly	Tyr	Ala	His 445	Phe	Thr	Val
Gly	Ser 450	His	Gln	Ala	Ala	Ala 455	Ile	His	Ile	Gly	Ala 460	Lys	Leu	Gly	Ser
Val 465	Cys	Gln	Asp	Cys	Gly 470	Gly	Thr	Ala	Ala	Glu 475	Thr	Lys	Val	Cys	Phe 480
Asp	Asn	Ala	Gln	Asn 485	Phe	Ser	Gln	Pro	Tyr 490	Leu	His	Tyr	Trp	Asn 495	Val
Asn	Ala	Asp	Gln 500	Ala	Val	Ala	Asn	Ala 505	Thr	Trp	Pro	Gly	Val 510	Ala	Met
Thr	Ala	Glu 515	Asn	Gly	Gly	Tyr	Cys 520	Tyr	Asp	Phe	Gly	Val 525	Gly	Leu	Asn
Ser	Leu 530	Gln	Val	Ile	Phe	Ser 535	Asp	Asn	Gly	Ala	Ser 540	Gln	Thr	Ala	Asp
Leu 545	Thr	Ala	Ser	Ser	Pro 550	Thr	Leu	Cys	Tyr	Gln 555	Asn	Gly	Thr	Trp	Arg 560
Asp	Ser	Asp	Phe 565	Cys	Gln	Ser	Ser	Asn	Val 570	Gly	Asn	Glu	Ser	Trp 575	Tyr
Phe	Arg	Gly	Thr 580	Ser	Asn	Gly	Trp	Gly 585	Val	Ser	Ala	Leu	Thr	Tyr	Glu
Ala	Ala	Thr	Gly	Leu	Tyr	Thr	Thr	Val	Gln	Ser	Phe	Asn	Gly	Glu	Glu

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      595      600      605
Ser Pro Ala Arg Phe Lys Ile Asp Asp Gly Asn Trp Ser Glu Ser Tyr
      610      615      620
Pro Ser Ala Asp Tyr Gln Val Gly Asp Tyr Ala Thr Tyr Thr Ile Thr
625      630      635      640
Phe Asp Ser Gln Thr Lys Ala Ile Thr Val Thr Ser Gln
      645      650

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<210> 195
 <211> 2790
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample.

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<400> 195
atgctgacag accgttttctt tgatggcgat acatcaaaca acgaccctta caaccagaac      60
tacgatgcta aaaacgaccg gggaacttat cagggcgggc atttttaaagg aatcacgcaa      120
aaattggatt atctcgataa gctaggcggt aacacaatct ggatcagccc gatcgtggaa      180
aatatcaagc atgatgtccg ttatgacaac tctgaagggc attcatacta tgcttaccac      240
ggctactggg caagcaactt cgggtgcgta aacccacact tcggtacaat ggaagatttc      300
catacactga ttgacgctgc ccatgaaaaa ggcatacaaga tcatggttga cgtagtatta      360
aaccacactg gttatggcctt aaaagatatc aacggagaag tttccaatcc tccagccggt      420
taccacaactg acgcagaacg cagcacatat agcagcctgc ttgccaggg ttcaaatgtc      480
ggctctgatg aggttggttg cgaattagct ggcctacctg acttaaaaac agaagacccc      540
gcagtccgcc agacaatcat cgactggcaa acagactgga tcacgaaagc tactacagct      600
aaaggaaaca caattgacta cttccgtgtc gacactgtga agcacgttga agacgcaaca      660
tggatggcat tcaaaaatga cctcactgaa aaaatgccga cacacaaaat gatcggggaa      720
gcttggggag caagtgccaa taaccaactt ggataccttg aaacagggtat gatggactca      780
ctgcttgact tcgacttcaa aggcattgcg cagcatttcg tgaacggcaa gcttaaggca      840
gcaaacgatg cctctgccaa ccgcaacggt aatggaacg acacagctac tttaggttca      900
ttccttggaa gccatgacga agatggtttc ctattttaaag aaggaaatga caaaggcaag      960
cttaagggtg ctgcttccct gcaagcaaca tcaaaaggcc agccggtcat ctattatggt      1020
gaagagcttg gtcaaagtgg agcaaacaac tatccgcaat acgataaccg ttatgacctg      1080
gcatgggaca aagttgaaaa caacgacgtc cttgagcact acactaagggt cctgaacttc      1140
agaagcgctc attcagaagt gttcgctaaa ggtgaacgcg caacaattgg cggttctgac      1200
gtgataaat tcttactttt tgctcgtaaa aatggaacg aagctgctta cgtcggcttg      1260
aacgttgctg acacagcaaa agacgtaaca ctgactgttt ctgcagggtg agtcgtaact      1320
gaccactatg cagataaaac ttatactgct tcagaagctg gagaaatcac attgacgatc      1380
ccggcaaaaag ctgatggcgg tactgtttta ctaacggttg aaggcggaga aatcacagct      1440
gctaaagcgg caagcgaagg cgacggcaca gttgagccag tccctgcgaa ccacatccgc      1500
attcactaca accgtacaga caacaactat gaaaactacg gtgcatggct gtggaacgat      1560
gtagcctccc cttctgccaa ctggccgact ggcgctacaa tgtttgaaaa aacagacagc      1620
tacggtgcat acatcgacgt accacttaaa gagggcgcta agaacatcgg cttcctcggt      1680
atggatgtaa caaaagggtg tcagggtaaa gacggcgggc acaaagggtt tacgatctca      1740
tcacctgaaa tgaacgaaat ttggatcaag caaggttctg acaagggtgta cacttacgag      1800
ccagttgatc ttccggcgaa cactgtccgc gtccactatg tacgtgacaa cgcagactac      1860
gaaaacttcg gtatctggaa ctggggcgat gtaacagcac cttccgaaaa ctggcctaca      1920
ggcgacgcaa aattcgatgg tacagaccgt tacggtgcgt atgtcgacat tacgctaaaa      1980
gaaggcgcaa agaacattgg aatgattgct cttaacactg caaatggaga gaaagacggc      2040
ggagataaat ctttcaacct tctggataaa tataatcgca tttggattaa acaagggtgat      2100
gacaaatgtc acgtttctcc atactgggag aaggtgactg acggcttaac acctgaatct      2220
ttaaaaggag gtcttgtaat taaagattca actggtgctg aagttgccat cgaaagtgc      2280
gaaatcacaa gcgcaacctc tgtaaaagta aaagcaacat tcgatttaga aaagcttcca      2340
ttatccatca catacgaggc cagaacagtt tcagcttcaa ctggctggag aatgcttgat      2400
gaaatgtacg cttatgatgg aaacgacctt ggtgcgactt acaaggacgg agcagcgacg      2460
cttaaattat gggctccgaa agcgaagcaag gtaaccgcta acttctttga taaaaataat      2520
gccgctgaaa aaatcggcag cgtcgagtta acgaagggtg aaaaaggagt ctggctcagct      2580
atggttgctc ctggcgacct gaacgtaacc gatcttgaag gttattttta ccagtatgat      2640
gtaacaaatg acgggtataa tcgccagggt ttagatcctt atgcaaaatc aatggcagcc      2700
tttactgtga atacagaagg caatgctggg cctgacgggg acactgttgg caaggcggca      2760
attcaaaaag cttctcgaga gtacttctag                                     2790

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<210> 196
 <211> 929
 <212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 196

Met 1 Leu Thr Asp Arg 5 Phe Phe Asp Gly Asp 10 Thr Ser Asn Asn 15 Asp Pro
 Tyr Asn Gln Asn 20 Tyr Asp Ala Lys Asn 25 Asp Arg Gly Thr Tyr 30 Gln Gly
 Gly Asp Phe 35 Lys Gly Ile Thr Gln 40 Lys Leu Asp Tyr Leu 45 Asp Lys Leu
 Gly Val 50 Asn Thr Ile Trp Ile 55 Ser Pro Ile Val Glu 60 Asn Ile Lys His
 Asp 65 Val Arg Tyr Asp Asn 70 Ser Glu Gly His Ser 75 Tyr Tyr Ala Tyr His 80
 Gly Tyr Trp Ala Ser 85 Asn Phe Gly Ala Leu 90 Asn Pro His Phe Gly 95 Thr
 Met Glu Asp Phe 100 His Thr Leu Ile Asp 105 Ala Ala His Glu Lys 110 Gly Ile
 Lys Ile Met 115 Val Asp Val Val Leu 120 Asn His Thr Gly Tyr 125 Gly Leu Lys
 Asp Ile Asn Gly Glu Val Ser 135 Asn Pro Pro Ala Gly 140 Tyr Pro Thr Asp
 Ala 145 Glu Arg Ser Thr Tyr 150 Ser Ser Leu Leu Arg 155 Gln Gly Ser Asn Val 160
 Gly Ser Asp Glu Val 165 Val Gly Glu Leu Ala 170 Gly Leu Pro Asp Leu Lys 175
 Thr Glu Asp Pro 180 Ala Val Arg Gln Thr 185 Ile Ile Asp Trp Gln Thr Asp 190
 Trp Ile Thr 195 Lys Ala Thr Thr Ala 200 Lys Gly Asn Thr Ile 205 Asp Tyr Phe
 Arg Val 210 Asp Thr Val Lys His 215 Val Glu Asp Ala Thr 220 Trp Met Ala Phe
 Lys 225 Asn Asp Leu Thr Glu 230 Lys Met Pro Thr His 235 Lys Met Ile Gly Glu 240
 Ala Trp Gly Ala Ser 245 Ala Asn Asn Gln Leu 250 Gly Tyr Leu Glu Thr Gly 255
 Met Met Asp Ser 260 Leu Leu Asp Phe Asp 265 Phe Lys Gly Ile Ala His Asp 270
 Phe Val Asn Gly Lys Leu Lys Ala 280 Ala Asn Asp Ala Leu Thr Ala Arg 285
 Asn Gly 290 Lys Ile Asp Asn Thr 295 Ala Thr Leu Gly Ser Phe Leu Gly Ser 300
 His 305 Asp Glu Asp Gly Phe 310 Leu Phe Lys Glu Gly Asn Asp Lys Gly Lys 320
 Leu Lys Val Ala 325 Ser Leu Gln Ala Thr 330 Ser Lys Gly Gln Pro Val 335
 Ile Tyr Tyr Gly 340 Glu Glu Leu Gly Gln 345 Ser Gly Ala Asn Asn Tyr Pro 350
 Gln Tyr Asp 355 Asn Arg Tyr Asp Leu 360 Ala Trp Asp Lys Val Glu Asn Asn
 Asp Val 370 Leu Glu His Tyr Thr 375 Lys Val Leu Asn Phe Arg Ser Ala His 380
 Ser 385 Glu Val Phe Ala Lys 390 Gly Glu Arg Ala Thr 395 Ile Gly Gly Ser Asp 400
 Ala Asp Lys Phe Leu 405 Leu Phe Ala Arg Lys 410 Asn Gly Asn Glu Ala Ala 415
 Tyr Val Gly Leu 420 Asn Val Ala Asp Thr 425 Ala Lys Asp Val Thr Leu Thr 430
 Val Ser Ala 435 Gly Ala Val Val Thr 440 Asp His Tyr Ala Asp 445 Lys Thr Tyr
 Thr 450 Ala Ser Glu Ala Gly Glu 455 Ile Thr Leu Thr Ile Pro Ala Lys Ala 460
 Asp 465 Gly Gly Thr Val Leu 470 Leu Thr Val Glu Gly 475 Glu Glu Ile Thr Ala 480
 Ala Lys Ala Ala Ser 485 Glu Gly Asp Gly Thr 490 Val Glu Pro Val Pro 495

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Asn	His	Ile	Arg	Ile	His	Tyr	Asn	Arg	Thr	Asp	Asn	Asn	Tyr	Glu	Asn
Tyr	Gly	Ala	Trp	Leu	Trp	Asn	Asp	Val	Ala	Ser	Pro	Ser	Ala	Asn	Trp
Pro	Thr	Gly	Ala	Thr	Met	Phe	Glu	Lys	Thr	Asp	Ser	Tyr	Gly	Ala	Tyr
Ile	Asp	Val	Pro	Leu	Lys	Glu	Gly	Ala	Lys	Asn	Ile	Gly	Phe	Leu	Val
Met	Asp	Val	Thr	Lys	Gly	Asp	Gln	Gly	Lys	Asp	Gly	Gly	Asp	Lys	Gly
Phe	Thr	Ile	Ser	Ser	Pro	Glu	Met	Asn	Glu	Ile	Trp	Ile	Lys	Gln	Gly
Ser	Asp	Lys	Val	Tyr	Thr	Tyr	Glu	Pro	Val	Asp	Leu	Pro	Ala	Asn	Thr
Val	Arg	Val	His	Tyr	Val	Arg	Asp	Asn	Ala	Asp	Tyr	Glu	Asn	Phe	Gly
Ile	Trp	Asn	Trp	Gly	Asp	Val	Thr	Ala	Pro	Ser	Glu	Asn	Trp	Pro	Thr
Gly	Ala	Ala	Lys	Phe	Asp	Gly	Thr	Asp	Arg	Tyr	Gly	Ala	Tyr	Val	Asp
Ile	Thr	Leu	Lys	Glu	Gly	Ala	Lys	Asn	Ile	Gly	Met	Ile	Ala	Leu	Asn
Thr	Ala	Asn	Gly	Glu	Lys	Asp	Gly	Gly	Asp	Lys	Ser	Phe	Asn	Leu	Leu
Asp	Lys	Tyr	Asn	Arg	Ile	Trp	Ile	Lys	Gln	Gly	Asp	Asn	Val	Tyr	
Val	Ser	Pro	Tyr	Trp	Glu	Gln	Ala	Thr	Gly	Ile	Thr	Asn	Ala	Glu	Val
Ile	Ser	Glu	Asp	Thr	Ile	Leu	Leu	Gly	Phe	Thr	Met	Thr	Asp	Gly	Leu
Thr	Pro	Glu	Ser	Leu	Lys	Gly	Gly	Leu	Val	Ile	Lys	Asp	Ser	Thr	Gly
Ala	Glu	Val	Ala	Ile	Glu	Ser	Ala	Glu	Ile	Thr	Ser	Ala	Thr	Ser	Val
Lys	Val	Lys	Ala	Thr	Phe	Asp	Leu	Glu	Lys	Leu	Pro	Leu	Ser	Ile	Thr
Tyr	Ala	Gly	Arg	Thr	Val	Ser	Ala	Ser	Thr	Gly	Trp	Arg	Met	Leu	Asp
Glu	Met	Tyr	Ala	Tyr	Asp	Gly	Asn	Asp	Leu	Gly	Ala	Thr	Tyr	Lys	Asp
Gly	Ala	Ala	Thr	Leu	Lys	Leu	Trp	Ala	Pro	Lys	Ala	Ser	Lys	Val	Thr
Ala	Asn	Phe	Phe	Asp	Lys	Asn	Asn	Ala	Ala	Glu	Lys	Ile	Gly	Ser	Val
Glu	Leu	Thr	Lys	Gly	Glu	Lys	Gly	Val	Trp	Ser	Ala	Met	Val	Ala	Pro
Gly	Asp	Leu	Asn	Val	Thr	Asp	Leu	Glu	Gly	Tyr	Phe	Tyr	Gln	Tyr	Asp
Val	Thr	Asn	Asp	Gly	Ile	Thr	Arg	Gln	Val	Leu	Asp	Pro	Tyr	Ala	Lys
Ser	Met	Ala	Ala	Phe	Thr	Val	Asn	Thr	Glu	Gly	Asn	Ala	Gly	Pro	Asp
Gly	Asp	Thr	Val	Gly	Lys	Ala	Ala	Ile	Gln	Lys	Ala	Ser	Arg	Glu	Tyr
Phe															

<210> 197
 <211> 1401
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 197
 atgaaaccgt caaaattcgt ttttctctct gctgccatcg cttgcagcct ctccagtacc
 gccaatgctg acgccatttt gcatgcattt aactggaagt actccgacgt cacgcaaaac

60
 120

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gcctcgcaaa	tcgcgggcggc	gggttataaa	aaagtgtga	tttcgccagc	actgaaatcg	180
agtggcaatg	aatgggtgggc	acgttatcaa	ccgcaagatc	tgcgcgtgat	cgattcccca	240
cttggcaaca	aaagtgcatt	aaaatccatg	attgatgctc	tgaaggcggg	cggcggttgat	300
gtgtatgccc	atgtggtgct	taaccatatg	gccaatgaaa	catggaagcg	tgaagactta	360
aattaccctg	gcagtgaagt	gctgcaacaa	tacgcagcta	acaccagtta	ttatgctggac	420
caaacgcctt	ttggcaattt	aacggaaaac	ctattctctg	gctttgactt	ccaccagaaa	480
ggctgtatta	gcgattggaa	tgatgccggc	aatgttcagt	actggcgtct	ttgtggcggg	540
gctggtgacc	gagggctgcc	agacttagat	ccgaacaact	gggtgggtgc	acagcaacgt	600
ttgtatttga	atgcgctaaa	aggttttaggt	gtgaaaggct	tccgcattga	tgcgggttaa	660
cacatgagcc	aatatcaaat	cgaccagatt	ttcactgcag	agattaccgc	cggaatgcac	720
gtgtttgggtg	aagtgatcac	cagtgggtggc	aaaggcgact	ccagctatga	gaacttctta	780
gcgccttatc	tcaacgccac	caaccattcg	gcttacgatt	tcccactggt	tgcctctatt	840
cgcaacgcct	tctcctacag	cgggtggcatg	aacatgcttc	atgatccaca	agcctatggc	900
caagggtctg	aaaacgcacg	ttcaattacc	tttaccatca	cgcacgacat	cccaacgaac	960
gacgggtttcc	gttatcaaat	catggatccg	aaagatgaag	agctggctta	cgcttatatc	1020
ctcggtaaag	atggcggcac	acctctgatt	tacagcgaca	acttacctga	taacgaagat	1080
cgtgataatc	gccgttggga	aggtgtttgg	aaccgtgacc	tgatgaagaa	catgttgctc	1140
ttccataacc	aaatgcaagg	gcaagagatg	acgatgctgt	acagcgacca	atgtctactg	1200
atgtttaagc	gcggtaaaca	aggggtgggtc	ggcattaata	aatgctggta	agagcgttct	1260
cataccgttg	acacctatca	gcatgagttc	aactggtatc	agccttacac	agatacactc	1320
actggcgctga	ctgaaaccgt	gagttcgcgt	taccacacct	tccgaattcc	agctcgcagc	1380
gcgcgcgatgt	acatgctcta	a				1401

<210> 198

<211> 466

<212> PRT

<213> Unknown

<220>

<223> obtained from an environmental sample.

<400> 198

Met	Lys	Pro	Ser	Lys	Phe	Val	Phe	Leu	Ser	Ala	Ala	Ile	Ala	Cys	Ser
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Leu	Ser	Ser	Thr	Ala	Asn	Ala	Asp	Ala	Ile	Leu	His	Ala	Phe	Asn	Trp
			20					25					30		
Lys	Tyr	Ser	Asp	Val	Thr	Gln	Asn	Ala	Ser	Gln	Ile	Ala	Ala	Ala	Gly
		35					40					45			
Tyr	Lys	Lys	Val	Leu	Ile	Ser	Pro	Ala	Leu	Lys	Ser	Ser	Gly	Asn	Glu
	50					55					60				
Trp	Trp	Ala	Arg	Tyr	Gln	Pro	Gln	Asp	Leu	Arg	Val	Ile	Asp	Ser	Pro
65					70				75						80
Leu	Gly	Asn	Lys	Ser	Asp	Leu	Lys	Ser	Met	Ile	Asp	Ala	Leu	Lys	Ala
			85					90						95	
Val	Gly	Val	Asp	Val	Tyr	Ala	Asp	Val	Val	Leu	Asn	His	Met	Ala	Asn
			100					105					110		
Glu	Thr	Trp	Lys	Arg	Glu	Asp	Leu	Asn	Tyr	Pro	Gly	Ser	Glu	Val	Leu
		115					120					125			
Gln	Gln	Tyr	Ala	Ala	Asn	Thr	Ser	Tyr	Tyr	Ala	Asp	Gln	Thr	Leu	Phe
	130					135					140				
Gly	Asn	Leu	Thr	Glu	Asn	Leu	Phe	Ser	Gly	Phe	Asp	Phe	His	Pro	Glu
145					150					155					160
Gly	Cys	Ile	Ser	Asp	Trp	Asn	Asp	Ala	Gly	Asn	Val	Gln	Tyr	Trp	Arg
			165						170					175	
Leu	Cys	Gly	Gly	Ala	Gly	Asp	Arg	Gly	Leu	Pro	Asp	Leu	Asp	Pro	Asn
			180					185					190		
Asn	Trp	Val	Val	Ser	Gln	Gln	Arg	Leu	Tyr	Leu	Asn	Ala	Leu	Lys	Gly
		195					200					205			
Leu	Gly	Val	Lys	Gly	Phe	Arg	Ile	Asp	Ala	Val	Lys	His	Met	Ser	Gln
	210					215					220				
Tyr	Gln	Ile	Asp	Gln	Ile	Phe	Thr	Ala	Glu	Ile	Thr	Ala	Gly	Met	His
225					230					235					240
Val	Phe	Gly	Glu	Val	Ile	Thr	Ser	Gly	Gly	Lys	Gly	Asp	Ser	Ser	Tyr
			245						250					255	
Glu	Asn	Phe	Leu	Ala	Pro	Tyr	Leu	Asn	Ala	Thr	Asn	His	Ser	Ala	Tyr
			260					265					270		
Asp	Phe	Pro	Leu	Phe	Ala	Ser	Ile	Arg	Asn	Ala	Phe	Ser	Tyr	Ser	Gly
		275					280					285			

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Gly Met Asn Met Leu His Asp Pro Gln Ala Tyr Gly Gln Gly Leu Glu
 290 300
 Asn Ala Arg Ser Ile Thr Phe Thr Ile Thr His Asp Ile Pro Thr Asn
 305 310 315 320
 Asp Gly Phe Arg Tyr Gln Ile Met Asp Pro Lys Asp Glu Glu Leu Ala
 325 330 335
 Tyr Ala Tyr Ile Leu Gly Lys Asp Gly Gly Thr Pro Leu Ile Tyr Ser
 340 345 350
 Asp Asn Leu Pro Asp Asn Glu Asp Arg Asp Asn Arg Arg Trp Glu Gly
 355 360 365
 Val Trp Asn Arg Asp Leu Met Lys Asn Met Leu Arg Phe His Asn Gln
 370 375 380
 Met Gln Gly Gln Glu Met Thr Met Leu Tyr Ser Asp Gln Cys Leu Leu
 385 390 395 400
 Met Phe Lys Arg Gly Lys Gln Gly Val Val Gly Ile Asn Lys Cys Gly
 405 410 415
 Glu Glu Arg Ser His Thr Val Asp Thr Tyr Gln His Glu Phe Asn Trp
 420 425 430
 Tyr Gln Pro Tyr Thr Asp Thr Leu Thr Gly Val Thr Glu Thr Val Ser
 435 440 445
 Ser Arg Tyr His Thr Phe Arg Ile Pro Ala Arg Ser Ala Arg Met Tyr
 450 455 460
 Met Leu
 465

<210> 199
 <211> 399
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 199
 gtgagtttga ccaaaaaggc tcagtacgaa ccaaatacgg caccaaggct cagtacatct 60
 ctgcaatcaa tgccgcgcac aacaacaata tccaaattta cggcgatggt gtgtttaacc 120
 accgaggtgg tgctgatggg aagtcgtggg tcgataccaa gcgcggtgat tgggacaacc 180
 gcaatattga actgggcgac aaatggattg aagccttgggt tgagtttaat tttcctggcc 240
 gcaacgacaa atactcgaac ttccattgga cttggtatca ctttgacggt gttgactggg 300
 atgacgccgg caaagaaaaa gcgatcttta aattcaaagg cgaaggaaaa gcatgggatt 360
 gggaaagtcag ctctgaaaaa ggcaattacg actacctaa 399

<210> 200
 <211> 132
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 200
 Val Ser Leu Thr Lys Lys Ala Gln Tyr Glu Pro Asn Thr Ala Pro Arg
 1 5 10 15
 Leu Ser Thr Ser Leu Gln Ser Met Pro Arg Thr Thr Thr Ile Ser Lys
 20 25 30
 Phe Thr Ala Met Leu Cys Leu Thr Thr Glu Val Val Leu Met Gly Ser
 35 40 45
 Arg Gly Ser Ile Pro Ser Ala Leu Ile Gly Thr Thr Ala Ile Leu Asn
 50 55 60
 Trp Ala Thr Asn Gly Leu Lys Leu Gly Leu Ser Leu Ile Phe Leu Ala
 65 70 75 80
 Ala Thr Thr Asn Thr Arg Thr Ser Ile Gly Leu Gly Ile Thr Leu Thr
 85 90 95
 Val Leu Thr Gly Met Thr Pro Ala Lys Lys Lys Arg Ser Leu Asn Ser
 100 105 110
 Lys Ala Lys Glu Lys His Gly Ile Gly Lys Ser Ala Leu Lys Lys Ala
 115 120 125
 Ile Thr Thr Thr

130

<210> 201
 <211> 1911
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 201
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 ggcgataaac aggtcggcca tggcaccggc tacggtacca gccatcacaa aggcgatctg 120
 caagggatca ttgactcgct ggattacatt caatcgctgg gcgtcaatgc catttggcta 180
 acgccgattt ttgaatctat tccggtggag ggacaagacc attgggcgga caggcttgat 240
 gctacaggct actttgccag tgactatttc aagatagacc ctgcgtttgg cacgtagaa 300
 caagcccgtg agctgggtga aaaggcacac gcgaaaggct tgtatgtctt ctttgatgga 360
 gtatttggtc accataaagg caatgtgggtg ccatcaccac aaggtagact gcctgtcggg 420
 gaaaataacc cggtcagcta cccagagagc ctggcgtttt acgaagaagt cgccagttac 480
 tgggtgaaa agttaaatg tgatggctgg cgtctggatc aagcctatca agtgccgacc 540
 gatgcatgga aagcgatccg tcagagcgtt gatgaagcgt cacagtccgt aacttatgtg 600
 aataacaaag gggaaaccgt ccatcctttg ggttacatgg tggctgaaat ttggaataac 660
 gaacgtttaca tcacagaaac cggttacggc aaagaaggcg atccggcggt gtgctcggct 720
 tttgattttc cgatgcgttt ccgagtggtc gaaacctttg cggttaacga aagtgggtgtc 780
 agccgaaaag gcggcgaatg gttgaatgac ggcattgtcac tgcacagtca gtatccggat 840
 catgccagc ctaatttaat gttgggcaac catgatgtgg tgcgctttgg ggatctgctg 900
 caacgtggcg gtattgcgtc accagaacaa ccgcaatact ggcagcgcta taaagcggcg 960
 atgtctttct tagcagcgta taccggccca attaccttgt attacggtga agaaattggc 1020
 gatcaggttg acggctttgc taaaaaaatc aaagaagatt gtgccgttat tggtttgtgt 1080
 gatgaccacg tggcgcgcac cagtgcgaag attgatggcg tgacggcgct actgaatgca 1140
 cagcagctct aactcaaagt atatgtctct tcattgatga cattacgtca gcaacatcct 1200
 gcgttatcac aaggggaacg tactaatgtg atggcgacag agacagtata cgtagaccat 1260
 aaacaggcag acaatgaagc cctgtttgtac atgggtagta cgactgataa cgcgagtgca 1320
 gtcacctga agggcaaagc gattggttca caaggtgtgc tgattgattt gttaacgaac 1380
 gagcgtttta tgcccaataa tggggagtat gccattccat taacgggctt tggcgcacga 1440
 ttctcaaga ttgacactcc gacagcggcg ggtgtgatgg cgcaatctgc tgcctcggta 1500
 tcgctagtag gtgaagggat catggcccaa tgtgatacc caaccgttga aggcaccggt 1560
 ccggtagcag aaaccttgta cgtggttggc gattttgccc atgctgggtg gaagcaaaag 1620
 ccgagcgcg cgtatcaata caaagggaag cacaatggca gcaacttgta tcaagtgggt 1680
 gtcgatgaaa aagcgggcgc ctacaagatg caatacgcca cgaaagattg gagccacag 1740
 ttactgcag acggtatggc attgaagccg ggtaccgcaa agtcgctcat agcgggtggc 1800
 tacggtaaag acaccgccgt gacgttgccg gaatccggta agtatgtgtg gagcttaaca 1860
 ttcagtgatc ttggcgagcc ggagcaaatc atggtgtcta agtgtcagta a 1911

<210> 202
 <211> 630
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 202
 Met Thr Ala Lys Ala Asp Asp Leu Arg Ile Tyr Gln Ile Met Val Glu
 1 5 10 15
 Ser Phe Val Asp Gly Asp Lys Gln Val Gly His Gly Thr Gly Tyr Gly
 20 25 30
 Thr Ser His His Lys Gly Asp Leu Gln Gly Ile Ile Asp Ser Leu Asp
 35 40 45
 Tyr Ile Gln Ser Leu Gly Val Asn Ala Ile Trp Leu Thr Pro Ile Phe
 50 55 60
 Glu Ser Ile Pro Val Glu Gly Gln Asp His Trp Ala Asp Arg Leu Asp
 65 70 75 80
 Ala Thr Gly Tyr Phe Ala Ser Asp Tyr Phe Lys Ile Asp Pro Arg Phe
 85 90 95
 Gly Thr Leu Glu Gln Ala Arg Glu Leu Val Glu Lys Ala His Ala Lys
 100 105 110
 Gly Leu Tyr Val Phe Phe Asp Gly Val Phe Gly His His Lys Gly Asn

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Val	Val	115	Ser	Pro	Gln	Gly	120	Leu	Pro	Val	Gly	125	Asn	Asn	Pro
Val	Ser	130	Tyr	Pro	Glu	Ser	135	Ala	Phe	Tyr	Glu	140	Val	Ala	Tyr
Trp	Val	145	Lys	Glu	Leu	Lys	150	Ile	Asp	Gly	Trp	155	Arg	Gln	Ala
Gln	Val	165	Pro	Thr	Asp	Ala	170	Lys	Ala	Ile	Arg	175	Gln	Ser	Val
Ala	Ser	180	Gln	Ser	Val	Thr	185	Val	Asn	Asn	Lys	190	Gly	Glu	Thr
Pro	Leu	195	Gly	Tyr	Met	Val	200	Glu	Ile	Trp	Asn	205	Glu	Arg	Tyr
Thr	Glu	210	Thr	Gly	Tyr	Gly	215	Lys	Glu	Gly	Asp	220	Ala	Leu	Cys
Phe	Asp	225	Phe	Pro	Met	Arg	230	Phe	Arg	Val	Val	235	Glu	Thr	Phe
Glu	Ser	245	Gly	Val	Ser	Arg	250	Lys	Gly	Gly	Trp	255	Leu	Asn	Asp
Ser	Leu	260	His	Ser	Gln	Tyr	265	Pro	Asp	His	Ala	270	Pro	Asn	Leu
Gly	Asn	275	His	Asp	Val	Val	280	Arg	Phe	Gly	Asp	285	Leu	Gln	Arg
Ile	Ala	290	Ser	Pro	Glu	Gln	295	Pro	Gln	Tyr	Trp	300	Arg	His	Lys
Met	Ser	305	Phe	Leu	Ala	Ala	310	Tyr	Thr	Gly	Pro	315	Ile	Thr	Leu
Glu	Glu	325	Ile	Gly	Asp	Gln	330	Val	Asp	Gly	Phe	335	Ala	Lys	Lys
Asp	Cys	340	Ala	Val	Ile	Gly	345	Leu	Cys	Asp	Asp	350	His	Val	Ala
Ala	Lys	355	Ile	Asp	Gly	Val	360	Ala	Ser	Leu	Asn	365	Ala	Gln	Gln
Leu	Lys	370	Val	Tyr	Val	Ser	375	Leu	Met	Thr	Leu	380	Arg	Gln	Gln
Ala	Leu	385	Ser	Gln	Gly	Glu	390	Arg	Thr	Asn	Val	395	Ala	Thr	Glu
Tyr	Val	405	Asp	His	Lys	Gln	410	Ala	Asp	Asn	Glu	415	Ala	Leu	Leu
Ser	Thr	420	Thr	Asp	Asn	Ala	425	Glu	Ser	Val	Thr	430	Leu	Lys	Gly
Gly	Ser	435	Gln	Gly	Val	Leu	440	Ile	Asp	Leu	Leu	445	Thr	Asn	Glu
Pro	Asn	450	Asn	Gly	Glu	Tyr	455	Ala	Ile	Pro	Leu	460	Gly	Phe	Gly
Phe	Leu	465	Lys	Ile	Asp	Thr	470	Pro	Thr	Ala	Ala	475	Val	Met	Ala
Ala	Ala	485	Ser	Val	Ser	Leu	490	Val	Gly	Glu	Gly	495	Ile	Met	Ala
Thr	Pro	500	Thr	Val	Glu	Gly	505	Pro	Val	Ala	Glu	510	Thr	Leu	Tyr
Val	Gly	515	Asp	Phe	Ala	Asp	520	Gly	Trp	Lys	Gln	525	Lys	Pro	Gln
Tyr	Gln	530	Tyr	Lys	Gly	Lys	535	His	Asn	Gly	Ser	540	Leu	Tyr	Gln
Val	Asp	545	Glu	Lys	Ala	Gly	550	Ala	Tyr	Lys	Met	555	Gln	Tyr	Ala
Trp	Ser	565	Pro	Gln	Phe	Thr	570	Ala	Asp	Gly	Met	575	Ala	Leu	Lys
Ala	Lys	580	Ser	Leu	Ile	Ala	585	Gly	Tyr	Gly	Lys	590	Thr	Ala	Val
Leu	Pro	595	Glu	Ser	Gly	Lys	600	Val	Trp	Ser	Leu	605	Phe	Ser	Asp
Gly	Glu	610	Pro	Glu	Gln	Ile	615					620			
625						630									

<210> 203
<211> 2601

<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 203
atgaagatga agtccccgggc gtggttggtta ggtagtgagc tggccatggc gttggcctct 60
tcggcagcca atgccggtgt catggttcac ctgttccagt ggaagtacaa tgacatcgcc 120
aacgagtgcg aaaagggtgct cgggtccaaa gggatgaag cagtgagat cagccgcct 180
gctgaacacc tgcaaggctc ctcttggtgg gtggtctatc agcccgtag ctacaagaac 240
ttcacttctc tggcggttaa cgaggccgaa ctcaaaagca tgatcgccg ttgcaaggcc 300
gccggggtca agatttacgc cgatgcggtt ttcaaccagc tggctggtgg atcaggcgct 360
ggtacaggtg gtagcagcta caatgccggc agcttcagct atccccatt tggctacaac 420
gatttccatc acgctgggag cctcaccaac tatgccgacc gcaacaatgt gcaaaacggc 480
gccctgctgg ggctgccgga tctggatacc ggctctgcct atgtgcagga tcagctggct 540
acatatatga agaccctgag tggctggggg gtggcaggtt ttcgtcttga tgcagcaaag 600
catatgagcg ttgccgatct ctccggccatc gtcagcaagg cgggcaatcc ttttgtctac 660
tccgaggtga ttggtgccac ggggtgaacca atccagccgg gcgaatatac cggcattggt 720
gccgtgacgg aatttaata cggcaccgat tccatgacca acttcaaggg gcagatcaag 780
aatctcaaga gcatggcgga gactggtggg ctgcttgcgt cgaacaaggc tgaagtcttt 840
gtggtcaacc atgaccgtga gcggggacat ggcggtggcg gtatgctgac ctacaaggat 900
ggtgccctct acaatctggc caacatcttc atgtggcct ggccctatgg cgcctatccc 960
caggtgatgt ccggctatga tttcggcacc aataccgata ttggtggggc gagcgctacc 1020
ccttggtctt ccggctctag ctggaactgc gaacaccgct ggagcaacat cgccaacatg 1080
gtctcgttcc ccaatgccg ccaaggcacg tccatgacca actggtggga taatggtaat 1140
aaccagatcg cctttggtcg cggcgccaag gcctttgtgg tgatcaacaa tgaatcttcc 1200
actctgagca agagcctgca gacgggtctg ccagccgggg agtactgcaa cattctggcc 1260
ggtgatgccc tgtgcagcgg cagcaccatc aagggtgatg ccagcggtat ggccaccttc 1320
aacgtggcag ggatgaaggc ggcagcgatc catatcaatg ccaagcccga tagcaccagc 1380
agtggcagct caggctcttc ctctggctct tcttctctg ccaccagtaa caagtttgcc 1440
agcatgaatc tgcggggcac caacaatggc tggggcagca cgcctatgac agtggatgac 1500
aaccgtgtct ggtcggcgga tgtcaccttt accggggccg cggatgccaa tggtgcccag 1560
cgcttcaagt ttgatgtcta tggcaactgg acagagagct atggcgatac acaagccgat 1620
ggcattgccc acaaggggag cgccaaggac atctatttca atggtgtggg caagtatcgt 1680
gtctcgctca aggagagcga catgagctac accctgacct tccgtcaagc tgggtgactc agtgggtgtc 1800
ccgggtggcg ccatcaccac caagacactc tccgtcaggc acagctggc taccggtggc 1860
gatgctccg gctccaccga tgatgtgggt gctctgggt gctctgggt ccaagaccat taccgtgaca 1920
agtggcaaga ccgaaactgt gctgtttgat gctctgggt gccagtgcc cgtcaccgt caccgatggc 1980
gtggccgatg ccgatggctt gacctccaag gccagtgcc cgtcaccgt caacagttgg 2040
agcgtggctt ataacagcaa ctttgccagc ctgaacttcc gtggcactcc caacttcgat 2100
ggcgcgagcag ccatgacgct ggtggcagac aacacctggg aggcaacggc caacttcgat 2160
ggtcaggcca atcagcgctt caagttcgat atcaaggggt actggagcca gaactatggt 2220
gatatcaaca aggatggggg ggccgaacgt accggtgccg atatttacac cactgtgacc 2280
ggtcaatata aggtgcaatt taacgactcc actttgaagt acaccctgac caagctggcc 2340
gatatgagcg ccaccagcta tagcgcaaac tttgccagcc tctacctgag tggcaccgcc 2400
aacagctggg gcaccaccgc catgaagctg gtggccaata acagctggca ggccgaggtg 2460
accttcaccg gcaagggcga tgccactggg gcccaacgct tcaagttcga cgtcaagggg 2520
gactggagcc agaactacgg tgacagcaac atggacggga ctgccgaacg gactgggtggc 2580
gatatcacca gtgccgtggg gggcacctat ctggtgacct ttaatgacag cacactgaaa 2601
tacaccctga ccgccaata a

<210> 204
<211> 866
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 204
Met Lys Met Lys Ser Arg Ala Trp Leu Leu Gly Ser Ala Val Ala Met
1 5 10 15
Ala Leu Ala Ser Ser Ala Ala Asn Ala Gly Val Met Val His Leu Phe
20 25 30
Gln Trp Lys Tyr Asn Asp Ile Ala Asn Glu Cys Glu Lys Val Leu Gly
35 40 45
Pro Lys Gly Tyr Glu Ala Val Gln Ile Thr Pro Pro Ala Glu His Leu

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Gln 65	50 Gly	Ser	Ser	Trp	Trp 70	55 Val	Val	Tyr	Gln	Pro 75	60 Val	Ser	Tyr	Lys	Asn 80
Phe	Thr	Ser	Leu	Gly 85	Gly	Asn	Glu	Ala	Glu 90	Leu	Lys	Ser	Met	Ile 95	Ala
Arg	Cys	Lys	Ala 100	Ala	Gly	Val	Lys	Ile 105	Tyr	Ala	Asp	Ala	Val 110	Phe	Asn
Gln	Leu	Ala 115	Gly	Gly	Ser	Gly	Val 120	Gly	Thr	Gly	Gly	Ser 125	Ser	Tyr	Asn
Ala 130	Gly	Ser	Phe	Ser	Tyr	Pro 135	Gln	Phe	Gly	Tyr	Asn 140	Asp	Phe	His	His
Ala 145	Gly	Ser	Leu	Thr	Asn 150	Tyr	Ala	Asp	Arg	Asn 155	Asn	Val	Gln	Asn	Gly 160
Ala	Leu	Leu	Gly	Leu 165	Pro	Asp	Leu	Asp	Thr 170	Gly	Ser	Ala	Tyr	Val 175	Gln
Asp	Gln	Leu	Ala 180	Thr	Tyr	Met	Lys	Thr 185	Leu	Ser	Gly	Trp	Gly 190	Val	Ala
Gly	Phe	Arg 195	Leu	Asp	Ala	Ala	Lys 200	His	Met	Ser	Val	Ala 205	Asp	Leu	Ser
Ala	Ile 210	Val	Ser	Lys	Ala	Gly 215	Asn	Pro	Phe	Val	Tyr 220	Ser	Glu	Val	Ile
Gly 225	Ala	Thr	Gly	Glu	Pro 230	Ile	Gln	Pro	Gly	Glu 235	Tyr	Thr	Gly	Ile	Gly 240
Ala	Val	Thr	Glu	Phe 245	Lys	Tyr	Gly	Thr	Asp 250	Leu	Ala	Ser	Asn	Phe 255	Lys
Gly	Gln	Ile	Lys 260	Asn	Leu	Lys	Ser	Met 265	Gly	Glu	Ser	Trp	Gly 270	Leu	Leu
Ala	Ser	Asn 275	Lys	Ala	Glu	Val	Phe 280	Val	Val	Asn	His	Asp 285	Arg	Glu	Arg
Gly	His 290	Gly	Gly	Gly	Gly	Met 295	Leu	Thr	Tyr	Lys	Asp 300	Gly	Ala	Leu	Tyr
Asn 305	Leu	Ala	Asn	Ile	Phe 310	Met	Leu	Ala	Trp	Pro 315	Tyr	Gly	Ala	Tyr	Pro 320
Gln	Val	Met	Ser	Gly 325	Tyr	Asp	Phe	Gly	Thr 330	Asn	Thr	Asp	Ile	Gly 335	Gly
Pro	Ser	Ala	Thr 340	Pro	Cys	Ser	Ser	Gly 345	Ser	Ser	Trp	Asn	Cys 350	Glu	His
Arg	Trp	Ser 355	Asn	Ile	Ala	Asn	Met 360	Val	Ser	Phe	His	Asn 365	Ala	Ala	Gln
Gly	Thr 370	Ser	Met	Thr	Asn	Trp 375	Trp	Asp	Asn	Gly	Asn 380	Asn	Gln	Ile	Ala
Phe 385	Gly	Arg	Gly	Ala	Lys 390	Ala	Phe	Val	Val	Ile 395	Asn	Asn	Glu	Ser	Ser 400
Thr	Leu	Ser	Lys	Ser 405	Leu	Gln	Thr	Gly	Leu 410	Pro	Ala	Gly	Glu	Tyr 415	Cys
Asn	Ile	Leu	Ala 420	Gly	Asp	Ala	Leu	Cys 425	Ser	Gly	Ser	Thr	Ile 430	Lys	Val
Asp	Ala	Ser 435	Gly	Met	Ala	Thr	Phe 440	Asn	Val	Ala	Gly	Met 445	Lys	Ala	Ala
Ala	Ile 450	His	Ile	Asn	Ala	Lys 455	Pro	Asp	Ser	Thr	Ser 460	Ser	Gly	Ser	Ser
Gly 465	Ser	Ser	Ser	Gly	Ser 470	Ser	Ser	Ser	Ala	Thr 475	Ser	Asn	Lys	Phe	Ala 480
Ser	Met	Asn	Leu	Arg 485	Gly	Thr	Asn	Asn	Gly 490	Trp	Ala	Ser	Thr	Ala 495	Met
Thr	Val	Asp	Ala 500	Asn	Arg	Val	Trp	Ser 505	Ala	Asp	Val	Thr	Phe 510	Thr	Gly
Ala	Ala	Asp 515	Ala	Asn	Gly	Ala	Gln 520	Arg	Phe	Lys	Phe	Asp 525	Val	Tyr	Gly
Asn	Trp 530	Thr	Glu	Ser	Tyr	Gly 535	Asp	Thr	Gln	Ala	Asp 540	Gly	Ile	Ala	Asp
Lys 545	Gly	Ser	Ala	Lys	Asp 550	Ile	Tyr	Phe	Asn	Gly 555	Val	Gly	Lys	Tyr	Arg 560
Val	Ser	Leu	Lys	Glu 565	Ser	Asp	Met	Ser	Tyr 570	Thr	Leu	Thr	Gln	Leu 575	Ser
Ser	Asn	Gln 580	Ala	Pro	Val	Ala	Ala	Ile 585	Thr	Pro	Lys	Thr	Leu 590	Ser	Val
Lys	Leu	Gly	Asp	Ser	Val	Val	Phe	Asp	Ala	Ser	Gly	Ser	Thr	Asp	Asp

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Val Gly 595 Thr Gly Tyr Ser 600 Trp Ser Thr Gly Gly 605 Ser Ala Lys Thr
 610 Val Thr Val Leu Phe Asp 615 Ala Leu Gly Thr Lys 620 Thr Ile Thr Val Thr
 625 Val Ala Asp Ala Asp 630 Gly Leu Thr Ser Lys 635 Ala Ser Ala Thr Val Thr
 Val Thr Asp Gly 645 Ser Val Ala Tyr Asn 650 Ser Asn Phe Ala Ser 655 Leu Asn
 Phe Arg Gly Thr Pro Asn Ser Trp 665 Gly Ala Ala Ala Met 670 Thr Leu Val
 675 Ala Asp Asn Thr Trp Glu Ala 680 Thr Val Asn Phe Asp 685 Gly Gln Ala Asn
 690 Gln Arg Phe Lys Phe Asp 695 Ile Lys Gly Asp Trp 700 Ser Gln Asn Tyr Gly
 705 Asp Ser Asn Lys Asp 710 Gly Val Ala Glu Arg Thr Gly Ala Asp Ile Tyr
 Thr Thr Val Thr 725 Gly Gln Tyr Lys Val 730 Phe Asn Asp Ser 735 Thr Leu
 Lys Tyr Thr 740 Leu Thr Lys Leu Ala 745 Asp Ser Ser Ala Thr 750 Ser Tyr Ser
 755 Ala Asn Phe Ala Ser Leu Tyr Leu Arg Gly Thr Pro Asn Ser Trp Gly
 770 Thr Thr Ala Met Lys Leu Val Ala Asn Asn Ser Trp 780 Gln Ala Glu Val
 785 Thr Phe Thr Gly Lys 805 Gly Asp Ala Thr Gly 810 Ala Gln Arg Phe Lys 815 Phe
 Asp Val Lys Gly Asp Trp Ser Gln Asn Tyr Gly Asp Ser Asn Met Asp
 820 Gly Thr Ala Glu Arg Thr Gly Gly Asp Ile Thr Ser Ala Val Val Gly
 835 Thr Tyr Leu Val Thr Phe Asn 840 Asp Ser Thr Leu Lys Tyr Thr Leu Thr
 850 Ala Lys
 865

<210> 205

<211> 1674

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 205

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gaggctgcca	atgtatat	tttgttaact	gaccg	tttta	acaacggtaa	cccaaacaat	180	
gacatcaatt	ttaataggac	taaagaatca	ggaaaactcc	gcaattttat	gggaggcgat	240		
atcaagggca	tcacccaaaa	aataaatgag	gggtatttta	gtaaactagg	cgttaatgcc	300		
atctggccta	ccccggttgt	tgaacaaata	catggcagtg	ttgatgaagg	taccggcaat	360		
acctatgcct	ttcatggcta	ttgggcaaaa	gattggacaa	acttagaccc	aaattttggc	420		
acaaaagaag	accttgccga	actggtggca	actgcccatg	caaaaggcat	caggatactt	480		
ttagatgtgg	taataaacca	caccggcccc	gtaaccgacc	aagaccgggt	ttggggagaa	540		
gattgggtac	gtacagcccc	gcagtgtacc	tatgataatt	acaccaatac	caccagttgc	600		
acgctggtag	ccaatttacc	tgatatactt	acagaaagta	atgaaaatgt	ggccttacca	660		
acctttttgt	tagataaatg	gaaagccgaa	ggcagattag	agcaagaact	aaaagaactt	720		
gacgattttt	tttcccgcac	aggccaccca	cgcgcacccc	gctttttacat	tattaaatgg	780		
cttaccgatt	acatccgaga	at	ttggggta	gatgggttta	gggttgatac	840		
accgaagaaa	cggtttgggc	cgagttgtat	gatgaagccg	taattgcttt	tgccgaatat	900		
aaaaagcca	acccagacaa	ggtattggac	gataatgaat	tttatatggt	aggcgaagt	960		
tacaactacg	gtattttccg	cggaagggtc	tatgatttcg	gcgataaaaa	ggtggactat	1020		
tttgaccacg	gattttaaag	cctcatcaat	tttgaaatga	aatatgatgc	caattttacc	1080		
tacgatacac	tttttaggaa	gtacgatacc	cttttgcata	ccaaacttaa	aggcagaagt	1140		
gtgctcaact	acctctcatc	tcacgacgat	ggaagtccat	ttgataaaaat	gcggcaaaaa	1200		
ccatacgagt	cggtacaaa	attactgctc	actccggg	cg	catcccaaat	ttattacggt	1260	
gacgaaaccg	ccagaagcct	taacatagaa	ggcgacacag	gagatgctac	gcttcgttcg	1320		
ttttgaatt	gggaagagct	cgcagaagac	cctgccaa	gc	aaaaatact	tcagcatttg	1380	

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caaaaactgg	gcagtttcag	gaacaaccac	cccgcagttg	gtgccggaag	gcacaaaacc	1440
cttggcaaaa	agccgtttta	caccttttagc	agggttttatc	aaaaaaatgg	ttttattgac	1500
aaagtgtgg	tagcattaga	tgccccctaaa	ggccaaaaaac	aaattaccgt	taatgggtgtt	1560
tttgatgacg	gtacaaaact	tgtagatgcc	tattcaggca	aagaaacctc	agttaaaaat	1620
ggtatcgttt	cactttcttc	tgaatttgat	attgttttgt	tagaacaaaa	ataa	1674

<210> 206
 <211> 557
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 206

Met	Tyr	Arg	Val	Ile	Pro	Ile	Ile	Leu	Ile	Met	Ser	Met	Ile	Val	Ala
1				5				10					15		
Cys	Glu	Ser	Pro	Lys	Lys	Lys	Thr	Thr	Glu	Thr	Ala	Gln	Pro	Ser	Thr
			20					25					30		
Asn	Ala	Glu	Lys	Pro	Phe	Val	Trp	Glu	Ala	Ala	Asn	Val	Tyr	Phe	Leu
		35					40					45			
Leu	Thr	Asp	Arg	Phe	Asn	Asn	Gly	Asn	Pro	Asn	Asn	Asp	Ile	Asn	Phe
		50				55					60				
Asn	Arg	Thr	Lys	Glu	Ser	Gly	Lys	Leu	Arg	Asn	Phe	Met	Gly	Gly	Asp
65					70					75					80
Ile	Lys	Gly	Ile	Thr	Gln	Lys	Ile	Asn	Glu	Gly	Tyr	Phe	Ser	Lys	Leu
				85					90					95	
Gly	Val	Asn	Ala	Ile	Trp	Leu	Thr	Pro	Val	Val	Glu	Gln	Ile	His	Gly
			100					105					110		
Ser	Val	Asp	Glu	Gly	Thr	Gly	Asn	Thr	Tyr	Ala	Phe	His	Gly	Tyr	Trp
		115					120					125			
Ala	Lys	Asp	Trp	Thr	Asn	Leu	Asp	Pro	Asn	Phe	Gly	Thr	Lys	Glu	Asp
		130				135					140				
Leu	Ala	Glu	Leu	Val	Ala	Thr	Ala	His	Ala	Lys	Gly	Ile	Arg	Ile	Leu
145					150					155					160
Leu	Asp	Val	Val	Ile	Asn	His	Thr	Gly	Pro	Val	Thr	Asp	Gln	Asp	Pro
				165					170					175	
Val	Trp	Gly	Glu	Asp	Trp	Val	Arg	Thr	Gly	Pro	Gln	Cys	Thr	Tyr	Asp
			180					185					190		
Asn	Tyr	Thr	Asn	Thr	Thr	Ser	Cys	Thr	Leu	Val	Ala	Asn	Leu	Pro	Asp
		195					200					205			
Ile	Leu	Thr	Glu	Ser	Asn	Glu	Asn	Val	Ala	Leu	Pro	Thr	Phe	Leu	Leu
	210				215						220				
Asp	Lys	Trp	Lys	Ala	Glu	Gly	Arg	Leu	Glu	Gln	Glu	Leu	Lys	Glu	Leu
225					230					235					240
Asp	Asp	Phe	Phe	Ser	Arg	Thr	Gly	His	Pro	Arg	Ala	Pro	Arg	Phe	Tyr
				245					250					255	
Ile	Ile	Lys	Trp	Leu	Thr	Asp	Tyr	Ile	Arg	Glu	Phe	Gly	Val	Asp	Gly
			260					265					270		
Phe	Arg	Val	Asp	Thr	Val	Lys	His	Thr	Glu	Glu	Thr	Val	Trp	Ala	Glu
		275					280					285			
Leu	Tyr	Asp	Glu	Ala	Val	Ile	Ala	Phe	Ala	Glu	Tyr	Lys	Lys	Ala	Asn
	290					295					300				
Pro	Asp	Lys	Val	Leu	Asp	Asn	Glu	Phe	Tyr	Met	Val	Gly	Glu	Val	
305					310				315					320	
Tyr	Asn	Tyr	Gly	Ile	Ser	Gly	Gly	Arg	Phe	Tyr	Asp	Phe	Gly	Asp	Lys
				325					330					335	
Lys	Val	Asp	Tyr	Phe	Asp	His	Gly	Phe	Lys	Ser	Leu	Ile	Asn	Phe	Glu
			340					345					350		
Met	Lys	Tyr	Asp	Ala	Asn	Phe	Thr	Tyr	Asp	Thr	Leu	Phe	Arg	Lys	Tyr
		355					360					365			
Asp	Thr	Leu	Leu	His	Thr	Lys	Leu	Lys	Gly	Arg	Ser	Val	Leu	Asn	Tyr
	370					375					380				
Leu	Ser	Ser	His	Asp	Asp	Gly	Ser	Pro	Phe	Asp	Lys	Met	Arg	Gln	Lys
385					390					395					400
Pro	Tyr	Glu	Ser	Ala	Thr	Lys	Leu	Leu	Leu	Thr	Pro	Gly	Ala	Ser	Gln
				405					410					415	
Ile	Tyr	Tyr	Gly	Asp	Glu	Thr	Ala	Arg	Ser	Leu	Asn	Ile	Glu	Gly	Ala

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Gln	Gly	Asp	420	Ala	Thr	Leu	Arg	Ser	425	Phe	Met	Asn	Trp	Glu	430	Glu	Leu	Ala
		435						440						445				
Glu	Asp	Pro	Ala	Lys	Gln	Lys	Ile	Leu	Gln	His	Trp	Gln	Lys	Leu	Gly			
	450					455					460							
Ser	Phe	Arg	Asn	Asn	His	Pro	Ala	Val	Gly	Ala	Gly	Arg	His	Lys	Thr			
465					470					475					480			
Leu	Gly	Lys	Lys	Pro	Phe	Tyr	Thr	Phe	Ser	Arg	Val	Tyr	Gln	Lys	Asn			
				485					490					495				
Gly	Phe	Ile	Asp	Lys	Val	Val	Val	Ala	Leu	Asp	Ala	Pro	Lys	Gly	Gln			
			500					505					510					
Lys	Gln	Ile	Thr	Val	Asn	Gly	Val	Phe	Asp	Asp	Gly	Thr	Lys	Leu	Val			
		515					520					525						
Asp	Ala	Tyr	Ser	Gly	Lys	Glu	Thr	Ser	Val	Lys	Asn	Gly	Ile	Val	Ser			
	530					535					540							
Leu	Ser	Ser	Glu	Phe	Asp	Ile	Val	Leu	Leu	Glu	Gln	Lys						
545					550					555								

<210> 207
 <211> 1378
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 207
 ctgtcgactg agccttttcgt tttgggctcg agactgactc tcagcccacc ccgcagtagc 60
 tccagacgga gtagccgtaa tagccgttgg ccgggtcgtg ggcaggggccc tcgaggtaca 120
 cccacccgct tgagtccacc cacttggtcca cccagccgcc gaggttgccg gtgtactcgt 180
 ggatgcacgc tcccgcgaac ttcggaacgt agaccacct tccggctttg cttgaggcga 240
 ggttgatgta tgttatcagt cccggcttgc ttcctgtagcc gtttctcacg aatatcagct 300
 cgctcgttgc gtagtaaacg acgtcagtgc ttcctccggc cagggttgta tgtatccaga 360
 tgaggttctt gagcttatcc ttgttgagcc actcctcgtg gtcgcggtag aatatgtcgc 420
 gctggccctc gtaggtgagg atgaacgcgt aggttgata cttgttccag attatatcgc 480
 tgctcgtggtt tgcaacgaag gttacggcct taaacgggtc gcggctgacg actgtgcccc 540
 cgttcttgag ggctcgcagc agtgcgaggaa tgttcttggc gtcaaaggcc gcgtccatct 600
 ttagtagag cggaagtcg aagaccttgg cgccgctcga gtaggccagc ttgaggagtg 660
 catcaacggt ggtgtccag tactcgccaa cggcccagcc gccccaccag ttgagccagt 720
 ccttgacgac ccacgctccg tggcccttca cgtagtcaaa gcgccaggca tcaacgccga 780
 tgctccttag gtaggcggcg tagctctcat cgctcgccca gagccagtgc tgggtcccagc 840
 tcttctcgtg ggctatgtct gggaagcctc caaatgtgcc ctctgtcacag cacttgacct 900
 cgttggggtg gaagtcgagg tagttggcag tatatttgcc cgaggccacc tttgagaagt 960
 ccgtccagggt gtagtcccca acgaacgggt tccactcgag gtctccgcct gcgcggtggt 1020
 ttatgacgat gtccgctatg acctttatgc cgtaggcatg ggccgtgttt atcatgttca 1080
 cgagctcctg cttggagcca aagcgcgtct ctaccgttcc cttctgggtc tactcaccga 1140
 ggtcaaagaa gtcgtagggg tcgtagccca tcgaataggc gccgcccag ccttgcctgc 1200
 ccgggggaat ccaaattggcg gatattcccc cctcgtacca ctccggtatc ttgctcctga 1260
 tgggtgcccc ccagattcct ccacctggga cgtccagata gaaggcctgc attataacgc 1320
 cgccctcttc cagctcggag tacttgccca taagttacct cctactagta gattaaaa 1378

<210> 208
 <211> 439
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 208
 Leu Ser Thr Glu Pro Phe Val Leu Gly Ser Arg Leu Thr Leu Ser Pro
 1 5 10 15
 Pro Arg Ser Ser Ser Arg Arg Ser Ser Arg Asn Ser Arg Trp Pro Gly
 20 25 30
 Arg Gly Gln Gly Pro Arg Gly Thr Pro Thr Arg Leu Ser Pro Pro Thr
 35 40 45
 Cys Pro Pro Ser Arg Arg Gly Cys Arg Cys Thr Arg Gly Cys Thr Leu
 50 55 60

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Pro  Arg  Thr  Ser  Glu  Arg  Arg  Pro  Thr  Phe  Arg  Leu  Cys  Leu  Arg  Arg
65      70      75      80
Gly  Cys  Met  Leu  Ser  Val  Pro  Ala  Cys  Phe  Arg  Ser  Arg  Phe  Ser  Arg
      85      90      95
Ile  Ser  Ala  Arg  Arg  Cys  Arg  Ser  Lys  Arg  Arg  Gln  Cys  Phe  Leu  Arg
      100      105      110
Pro  Gly  Cys  His  Val  Ser  Arg  Gly  Ser  Ala  Tyr  Pro  Cys  Ala  Thr  Pro
      115      120      125
Arg  Ser  Arg  Gly  Arg  Ile  Leu  Ser  Ala  Gly  Pro  Arg  Arg  Gly  Thr  Arg
      130      135      140
Arg  Leu  Asp  Thr  Cys  Ser  Arg  Leu  Tyr  Arg  Cys  Arg  Gly  Leu  Gln  Arg
      145      150      155
Arg  Leu  Arg  Pro  Thr  Gly  Arg  Gly  Arg  Leu  Cys  Pro  Arg  Ser  Gly  Pro
      160      165      170      175
Arg  Arg  Val  Arg  Glu  Cys  Ser  Cys  Cys  Gln  Arg  Pro  Arg  Pro  Ser  Cys
      180      185      190
Ser  Arg  Ala  Gly  Ser  Arg  Arg  Pro  Trp  Arg  Arg  Ser  Ser  Arg  Pro  Ser
      195      200      205
Gly  Val  His  Gln  Arg  Trp  Cys  Pro  Ser  Thr  Arg  Gln  Arg  Pro  Ser  Arg
      210      215      220
Pro  Thr  Ser  Ala  Ser  Pro  Arg  Pro  Thr  Leu  Arg  Gly  Pro  Ser  Arg  Ser
      225      230      235      240
Gln  Ser  Ala  Arg  His  Gln  Arg  Arg  Cys  Ser  Leu  Gly  Arg  Arg  Arg  Ser
      245      250      255
Ser  His  Arg  Ser  Pro  Arg  Ala  Ser  Ala  Gly  Pro  Ser  Ser  Ser  Arg  Gly
      260      265      270
Leu  Cys  Leu  Gly  Ser  Leu  Gln  Met  Cys  Pro  Arg  His  Ser  Thr  Pro  Arg
      275      280      285
Trp  Gly  Gly  Ser  Arg  Gly  Ser  Trp  Gln  Tyr  Ile  Cys  Pro  Arg  Pro  Pro
      290      295      300
Leu  Arg  Ser  Pro  Ser  Arg  Cys  Ser  Pro  Gln  Arg  Thr  Gly  Ser  Thr  Arg
      305      310      315      320
Gly  Leu  Arg  Leu  Arg  Gly  Gly  Leu  Arg  Cys  Pro  Leu  Pro  Leu  Cys  Arg
      325      330      335
Arg  His  Gly  Pro  Cys  Leu  Ser  Cys  Ser  Arg  Ala  Pro  Ala  Trp  Ser  Gln
      340      345      350
Ser  Ala  Ser  Leu  Pro  Phe  Pro  Ser  Gly  Arg  Thr  His  Arg  Gly  Gln  Arg
      355      360      365
Ser  Arg  Arg  Gly  Arg  Ser  Pro  Ser  Asn  Arg  Arg  Arg  Pro  Cys  Pro  Cys
      370      375      380
Ser  Pro  Gly  Glu  Ser  Lys  Trp  Arg  Ile  Phe  Pro  Pro  Arg  Thr  Thr  Pro
      385      390      395      400
Val  Ser  Cys  Ser  Trp  Cys  Pro  Thr  Arg  Phe  Leu  His  Leu  Gly  Arg  Pro
      405      410      415
Ser  Arg  Arg  Pro  Ala  Leu  Arg  Arg  Pro  Leu  Pro  Ala  Arg  Ser  Thr  Trp
      420      425      430
Pro  Val  Thr  Ser  Tyr  Ile  Lys
      435

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<210> 209

<211> 1416

<212> DNA

<213> Unknown

<220>

<223> obtained from an environmental sample.

<400> 209

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atgattcagc ccattgcactc tcgggaacag gcctgccgtc tcattccggc actgatcatg      60
acatttgcac tggcactgcc gttgcaaatt cgtgccgatg tcaccctgca tgctttcaac      120
tggagctatg ccgatgtcgc tgatcgggcc gttgacatcg ctgcagcagg gtacagtgcc      180
gtgctggttg ccccgccact tcgatccgaa ggcacggcct ggtgggcgcg ataccagccc      240
caggatctcc gccttatcga ccattccgtg ggcaatacac atgacttcgt caacatgatc      300
gatgctctcg atgatgtggg tgtgggcgtg tacgccgaca tcgtgctcaa ccacatggcc      360
aatgaggctg cacaaaggcc tgacctgaac taccctggtc aggcagtgct tgacgaatat      420
gcttccgatc ccggtcattt cgagggcttg aggctgttcg gtaatctgag cttcaatttc      480
ctgtcggaac atgatttcgg acccgcccag tgcattcagg attacagcga tgtgtttcag      540
gtccagaact ggcggctgtg cggaccgccg ccggaccggg gcctgccgca cctggtcgcc      600

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aatgactggg tgatctctca acagcgccag tatctggaag ccatcaaggc gctgggtgtg 660
gctggcatgc gcatcgacgc ggtcaagcat atgccatga gccatatcaa tgccgttctc 720
acccccgaga tccggtcggg cttgcatgtg tttggcgaag tcatcacctc cggtggggct 780
ggtgatacat cctacgaccg ttttctggcc cttacctgg cacaaagcga ccatggtgcc 840
tatgactttc cattgtttga aaccattcgc cgtgctttcg gcttcggtgg cagcatgagt 900
gaactggtcg atcctgctgc ctacggtcag gccctgccac cggaccgcgc catcaccttc 960
gtcatcacgc acgatattcc gaacaatgac ggatttcgct accagatact cgaccccgctc 1020
gatgaatcac tggcctacgc ctacattctg ggccgcgatg gcggtgtccc gcttctgtat 1080
tccgacaaca atgaaagcgg cgatggccgc tggatcgatg cctggcaacg tccggatctg 1140
gttgcaatgg tcggcttcca caatgcagtc cacggtcagg acatggccgt gctttcacat 1200
gacgactgcc acctgctgtt tcggcgcggc agcctcgga ttgtcggcat caacaagtgc 1260
ggccatgcac tcagctcctg ggtcaacatg aaccagagcg tactgtggtg gtacgcggac 1320
tacacagacg tgctcgacag caacagcgtt gtcaacatcc agtcatcctg gcacgagttc 1380
atccttcccc cccgccaggc acgcctgtgg ttgcga 1416

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<210> 210
 <211> 472
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 210

Met	Ile	Gln	Pro	Met	His	Ser	Arg	Glu	Gln	Ala	Cys	Arg	Leu	Ile	Pro
1				5				10					15		
Ala	Leu	Ile	Met	Thr	Phe	Ala	Leu	Ala	Leu	Pro	Leu	Gln	Ile	Arg	Ala
			20					25					30		
Asp	Val	Thr	Leu	His	Ala	Phe	Asn	Trp	Ser	Tyr	Ala	Asp	Val	Ala	Asp
		35					40					45			
Arg	Ala	Val	Asp	Ile	Ala	Ala	Gly	Tyr	Ser	Ala	Val	Leu	Val	Ala	
		50				55					60				
Pro	Pro	Leu	Arg	Ser	Glu	Gly	Thr	Ala	Trp	Trp	Ala	Arg	Tyr	Gln	Pro
65					70					75					80
Gln	Asp	Leu	Arg	Leu	Ile	Asp	His	Pro	Leu	Gly	Asn	Thr	His	Asp	Phe
				85					90					95	
Val	Asn	Met	Ile	Asp	Ala	Leu	Asp	Asp	Val	Gly	Val	Gly	Val	Tyr	Ala
			100					105					110		
Asp	Ile	Val	Leu	Asn	His	Met	Ala	Asn	Glu	Ala	Ala	Gln	Arg	Pro	Asp
		115					120					125			
Leu	Asn	Tyr	Pro	Gly	Gln	Ala	Val	Leu	Asp	Glu	Tyr	Ala	Ser	Asp	Pro
		130				135					140				
Gly	His	Phe	Glu	Gly	Leu	Arg	Leu	Phe	Gly	Asn	Leu	Ser	Phe	Asn	Phe
145					150					155					160
Leu	Ser	Glu	His	Asp	Phe	Gly	Pro	Ala	Gln	Cys	Ile	Gln	Asp	Tyr	Ser
				165					170					175	
Asp	Val	Phe	Gln	Val	Gln	Asn	Trp	Arg	Leu	Cys	Gly	Pro	Pro	Pro	Asp
		180						185					190		
Pro	Gly	Leu	Pro	Asp	Leu	Val	Ala	Asn	Asp	Trp	Val	Ile	Ser	Gln	Gln
		195					200					205			
Arg	Gln	Tyr	Leu	Glu	Ala	Ile	Lys	Ala	Leu	Gly	Val	Ala	Gly	Met	Arg
		210				215					220				
Ile	Asp	Ala	Val	Lys	His	Met	Pro	Met	Ser	His	Ile	Asn	Ala	Val	Leu
225					230					235					240
Thr	Pro	Glu	Ile	Arg	Ser	Gly	Leu	His	Val	Phe	Gly	Glu	Val	Ile	Thr
				245					250					255	
Ser	Gly	Gly	Ala	Gly	Asp	Thr	Ser	Tyr	Asp	Arg	Phe	Leu	Ala	Pro	Tyr
			260					265					270		
Leu	Ala	Gln	Ser	Asp	His	Gly	Ala	Tyr	Asp	Phe	Pro	Leu	Phe	Glu	Thr
		275					280					285			
Ile	Arg	Arg	Ala	Phe	Gly	Phe	Gly	Gly	Ser	Met	Ser	Glu	Leu	Val	Asp
		290				295					300				
Pro	Ala	Ala	Tyr	Gly	Gln	Ala	Leu	Pro	Pro	Asp	Arg	Ala	Ile	Thr	Phe
305					310					315					320
Val	Ile	Thr	His	Asp	Ile	Pro	Asn	Asn	Asp	Gly	Phe	Arg	Tyr	Gln	Ile
				325					330					335	
Leu	Asp	Pro	Val	Asp	Glu	Ser	Leu	Ala	Tyr	Ala	Tyr	Ile	Leu	Gly	Arg
			340					345					350		

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Asp Gly Gly Val Pro Leu Leu Tyr Ser Asp Asn Asn Glu Ser Gly Asp
 355 360 365
 Gly Arg Trp Ile Asp Ala Trp Gln Arg Pro Asp Leu Val Ala Met Val
 370 375 380
 Gly Phe His Asn Ala Val His Gly Gln Asp Met Ala Val Leu Ser His
 385 390 395 400
 Asp Asp Cys His Leu Phe Arg Arg Gly Ser Leu Gly Ile Val Gly
 405 410 415
 Ile Asn Lys Cys Gly His Ala Leu Ser Trp Val Asn Met Asn Gln
 420 425 430
 Ser Val Leu Trp Trp Tyr Ala Asp Tyr Thr Asp Val Leu Asp Ser Asn
 435 440 445
 Ser Val Val Asn Ile Gln Ser Trp His Glu Phe Ile Leu Pro Ala
 450 455 460
 Arg Gln Ala Arg Leu Trp Leu Arg
 465 470

<210> 211
 <211> 1491
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 211
 gtgttttcgtt ctgacacagt ttcgcgtacc tgcattgtatg gtgcgctgcg taatgcctac 60
 caaccgatc ggggtgtttac tggagtcacg gtgcggacat gcaacttaaa aaagcatgct 120
 catcgccagg cgctgttggt catcgtgacg cgggtgcctgt gcctgaaatc caggcagacc 180
 cataaaaaaca acaacaaacc gataacaaac gacccaagcc ttctaagagg agaaaacggg 240
 atggctttta aactacgcaa aaaggcgctc gttggcctgt tcacggccgg cgcaatggta 300
 tatgccggtg cagcggcgag tggtgaaatc attctgcagg gcttccactg gcactccaag 360
 tggggcggca acaatcaggg ttggtggcag gtgatggaag gtcaggccaa caccatcgcc 420
 aacgccggct ttacgcacgt gtggttccc cgggtccata actcggccga tgccgaggg 480
 tacctacccc gcgagctgaa caacctcaac tccagctatg gctccgaagc acagctgcgc 540
 agcgccatcc aggcactgaa caatcgcggc gtgcattgca ttgccgatgt ggtcatgaac 600
 caccgggtgg gctgctctgg ctgggaggat ttctgtaacc cggactggcc gacctggtac 660
 atcgtcgcca atgattcctg gcccggtggc ccgaaaagcc agaactggga cacgggtgag 720
 acgtaccacg ccgcccgtga cctcgatcac gccaatccgc aggtgcgcaa cgatatctcg 780
 cactaccta acagccgcct caaggacgtc ggcttctccg gctggcgctg ggactatgcc 840
 aagggtttct ggcccggcta tgtcggcgag tacaactgga acaccaacc gaacttctgt 900
 gtgggtgagg tgtgggacga tctcgacccc aacaatccca acccgaccg ccagcaactg 960
 gtggactggg ttgatgctac cgggtggcag tgtcacgtct tcgacttcac caccaagggg 1020
 ctgacgaact atgcgtgca gcatggccag tactggcgcc tgcagggtga taatgggtgg 1080
 ccggctggcg gcatcggtg gtggccgcaa cgcattggtga ccttcgtcga caaccatgac 1140
 acgggcccga gcaatcactg tggtagcggc cagaacctct ggcccgtgcc ctgtgacaag 1200
 gtcatggagg cgtatgccta catcctgacc catccggcg tgccgctcgg gtactggacg 1260
 cacttcttca actggaatct tggtagcagg atcagccagt tgatgcagat ccgcaagaac 1320
 cagggcgtgc actccggttc cgacgtctgg atcgccagg cccgtcacgg cctgtacgcc 1380
 gcctatatca acggtaatgt ggcgatgaag atgggctggg ataactggag cccgggctgg 1440
 ggctggtcgc tggcggcctc cggttaacaac tgggcccgtct ggacacgctg a 1491

<210> 212
 <211> 496
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 212
 Val Phe Arg Ser Asp Thr Val Ser Arg Thr Cys Met Tyr Gly Ala Leu
 1 5 10 15
 Arg Asn Ala Tyr Gln Pro Asp Arg Val Phe Thr Gly Val Thr Val Arg
 20 25 30
 Thr Cys Asn Leu Lys Lys His Ala His Arg Gln Ala Leu Phe Ile
 35 40 45
 Val Thr Arg Cys Leu Cys Leu Lys Ser Arg Gln Thr His Lys Asn Asn

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50	Asn	Lys	Pro	Ile	Thr	55	Asn	Pro	Ser	Leu	60	Leu	Arg	Gly	Glu	Asn	Gly
65	Met	Ala	Phe	Lys	Leu	70	Arg	Lys	Lys	Ala	75	Leu	Val	Gly	Leu	Phe	Thr
				85	Tyr	Ala	Gly	Ala	Ala	Ala	90	Ala	Ser	Gly	Glu	Ile	95
	Gln	Gly	Phe	100	His	Trp	His	Ser	Lys	105	Trp	Gly	Gly	Asn	Asn	110	Gln
	Trp	Gln	Val	115	Met	Glu	Gly	Gln	Ala	120	Asn	Thr	Ile	Ala	Asn	Ala	Gly
	Thr	His	Val	130	Trp	Phe	Pro	Pro	Val	135	His	Asn	Ser	Ala	Asp	Ala	Glu
	Tyr	Leu	Pro	145	Arg	Glu	Leu	Asn	Asn	150	Leu	Asn	Ser	Ser	Tyr	Gly	Ser
	Ala	Gln	Leu	165	Ser	Ala	Ile	Gln	Ala	170	Leu	Asn	Asn	Arg	Gly	Val	His
	Ala	Ile	Ala	180	Asp	Val	Val	Met	Asn	185	His	Arg	Val	Gly	Cys	Ser	Gly
	Ala	Asp	Phe	195	Cys	Asn	Pro	Asp	Trp	200	Pro	Thr	Trp	Tyr	Ile	Val	Ala
	Asp	Ser	Trp	210	Pro	Gly	Gly	Pro	Lys	215	Ser	Gln	Asn	Trp	Asp	Thr	Gly
	Thr	Tyr	His	225	Ala	Ala	Arg	Asp	Leu	230	Asp	His	Ala	Asn	Pro	Gln	Val
	Asn	Asp	Ile	245	His	Tyr	Leu	Asn	Ser	250	Arg	Leu	Lys	Asp	Val	Gly	Phe
	Ser	Gly	Trp	260	Arg	Trp	Asp	Tyr	Ala	265	Lys	Gly	Phe	Trp	Pro	Gly	Tyr
	Gly	Glu	Tyr	275	Asn	Trp	Asn	Thr	Asn	280	Pro	Asn	Phe	Cys	Val	Gly	Glu
	Trp	Asp	Asp	290	Leu	Asp	Pro	Asn	Asn	295	Pro	Asn	Pro	His	Arg	Gln	Gln
	Val	Asp	Trp	305	Val	Asp	Ala	Thr	Gly	310	Gly	Ser	Cys	His	Val	Phe	Asp
	Thr	Thr	Lys	325	Gly	Leu	Thr	Asn	Tyr	330	Ala	Leu	Gln	His	Gly	Gln	Tyr
	Arg	Leu	Gln	340	Gly	Asp	Asn	Gly	Gly	345	Pro	Ala	Gly	Gly	Ile	Gly	Trp
	Pro	Gln	Arg	355	Met	Val	Thr	Phe	Val	360	Asp	Asn	His	Asp	Thr	Gly	Pro
	Asn	His	Cys	370	Gly	Asp	Gly	Gln	Asn	375	Leu	Trp	Pro	Val	Pro	Cys	Asp
	Val	Met	Glu	385	Ala	Tyr	Ala	Tyr	Ile	390	Leu	Thr	His	Pro	Gly	Val	Pro
	Val	Tyr	Trp	405	His	Phe	Phe	Asn	Trp	410	Asn	Leu	Gly	Ser	Glu	Ile	Ser
	Gln	Leu	Met	420	Gln	Ile	Arg	Lys	Asn	425	Gln	Gly	Val	His	Ser	Gly	Ser
	Val	Trp	Ile	435	Ala	Glu	Ala	Arg	His	440	Gly	Leu	Tyr	Ala	Ala	Tyr	Ile
	Gly	Asn	Val	450	Ala	Met	Lys	Met	Gly	455	Trp	Asp	Asn	Trp	Ser	Pro	Gly
	Gly	Trp	Ser	465	Ala	Ala	Ser	Gly	Asn	470	Asn	Trp	Ala	Val	Trp	Thr	Arg
				485						490							

<210> 213
 <211> 23
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 213
 Met Phe Leu Leu Ala Phe Leu Leu Thr Ala Ser Leu Phe Cys Pro Thr
 1 5 10 15
 Gly Gln Pro Ala Lys Ala Ala

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20

<210> 214
<211> 23
<212> PRT
<213> Bacterial

<400> 214
Val Leu Thr Phe His Arg Ile Ile Arg Lys Gly Trp Met Phe Leu Leu
1 5 10 15
Ala Phe Leu Leu Thr Ala Ser
20

<210> 215
<211> 33
<212> PRT
<213> Bacterial

<400> 215
Met Lys Ser Phe Ala Phe Met Pro Ile Leu Phe Tyr Ala Asn Asp Phe
1 5 10 15
Ile Ser Glu Arg Glu Gly Gly Gly Lys Met Gly Lys Asn Met Arg Arg
20 25 30
Arg

<210> 216
<211> 31
<212> PRT
<213> Bacterial

<400> 216
Met Arg Lys Lys Met Ser His Ser Arg Phe Thr Phe Leu Leu Ile Leu
1 5 10 15
Ala Leu Phe Ile Phe Phe Ser Gly Cys Ile Ser Glu Val Lys Ser
20 25 30

<210> 217
<211> 30
<212> PRT
<213> Bacterial

<400> 217
Met Tyr Thr Leu Phe Ile Arg Ser Phe Tyr Asp Thr Asn Asn Asp Gly
1 5 10 15
Val Gly Asp Tyr Asn Gly Val Ala Glu Lys Val Asp Tyr Leu
20 25 30

<210> 218
<211> 22
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 218
Val Leu Thr Phe His Arg Ile Ile Arg Lys Gly Trp Met Phe Leu Leu
1 5 10 15
Ala Phe Leu Leu Thr Ala
20

<210> 219
<211> 33
<212> PRT
<213> Unknown

<220>

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<223> Obtained from an environmental sample.

<400> 219

Met	Ser	Leu	Phe	Lys	Lys	Ile	Phe	Pro	Trp	Ile	Val	Ser	Leu	Leu	Leu
1				5					10				15		
Leu	Phe	Ser	Phe	Ile	Ala	Pro	Phe	Ser	Ile	Gln	Thr	Glu	Lys	Val	Arg
			20					25					30		

Ala

<210> 220

<211> 25

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 220

Met	Ala	Arg	Lys	Thr	Leu	Ala	Ile	Phe	Phe	Val	Leu	Leu	Val	Leu	Leu
1				5					10					15	
Ser	Leu	Ser	Ala	Val	Pro	Ala	Lys	Ala							
			20					25							

<210> 221

<211> 35

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 221

Met	Pro	Ala	Phe	Lys	Ser	Lys	Val	Met	His	Met	Lys	Leu	Lys	Tyr	Leu
1				5					10					15	
Ala	Leu	Val	Leu	Leu	Ala	Val	Ala	Ser	Ile	Gly	Leu	Leu	Ser	Thr	Pro
			20					25					30		

Val Gly Ala
35

<210> 222

<211> 28

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 222

Met	Lys	Gln	Gln	Lys	Arg	Leu	Tyr	Ala	Arg	Leu	Leu	Thr	Leu	Leu	Phe
1				5					10					15	
Ala	Leu	Ile	Phe	Leu	Leu	Pro	His	Ser	Ala	Ala	Ala				
			20					25							

<210> 223

<211> 21

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 223

Met	Arg	Val	Phe	Leu	Val	Val	Pro	Lys	Leu	Ser	Arg	Pro	Phe	Gln	Ala
1				5					10					15	
Glu	Ser	Gln	Gln	Gln											
			20												

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<210> 224
<211> 30
<212> PRT
<213> Bacterial

<400> 224
Met Gln Thr Phe Ala Phe Leu Phe Tyr Ser Lys Lys Gly Trp Val Cys
1 5 10 15
Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val
20 25 30

<210> 225
<211> 35
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 225
Met Pro Gln Ala Ile Arg Thr Phe Ser Arg Trp Thr Leu Phe Gly Leu
1 5 10 15
Ile Gly Val Phe Leu Leu Gly Leu Val Phe Ser Val Pro Pro Arg Ala
20 25 30
Ile Gln Ala
35

<210> 226
<211> 28
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 226
Val Val His Met Lys Leu Lys Tyr Leu Ala Leu Val Leu Leu Ala Val
1 5 10 15
Ala Ser Ile Gly Leu Leu Ser Thr Pro Val Gly Ala
20 25

<210> 227
<211> 30
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 227
Val Cys Met Asn Tyr Leu Lys Lys Val Trp Leu Tyr Tyr Ala Ile Val
1 5 10 15
Ala Thr Leu Ile Ile Tyr Phe Leu Thr Pro Phe Ser Thr Ala
20 25 30

<210> 228
<211> 30
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 228
Met Pro Gln Leu Tyr Pro Leu Pro Pro Arg Trp Arg Arg Ala Ala Arg
1 5 10 15
Gln Gly Leu Ala Ala Leu Thr Leu Ala Thr Thr Ala Leu Gly
20 25 30

<210> 229
 <211> 30
 <212> PRT
 <213> Unknown

<220>
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<400> 229
 Met Asn Asn Val Lys Lys Val Trp Leu Tyr Tyr Ser Ile Ile Ala Thr
 1 5 10 15
 Leu Val Ile Ser Phe Phe Thr Pro Phe Ser Thr Ala Gln Ala
 20 25 30

<210> 230
 <211> 28
 <212> PRT
 <213> Unknown

<220>
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<400> 230
 Val Gly Arg Ala Gly Leu Ala His His Ser Asn Thr Ser Ala Lys Gly
 1 5 10 15
 Thr Tyr Gly Ser Pro Leu Glu Leu Arg Pro Asp Arg
 20 25

<210> 231
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 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 231
 Met Lys Thr Phe Asn Leu Lys Pro Thr Leu Leu Pro Leu Thr Leu Leu
 1 5 10 15
 Leu Ser Ser Pro Val Leu Ala
 20

<210> 232
 <211> 23
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 232
 Met Lys Pro Ile Asn Thr Leu Leu Ile Ser Ala Leu Ala Val Cys Ser
 1 5 10 15
 Phe Ser Ser Ala Thr Tyr Ala
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<210> 233
 <211> 30
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 233
 Met Pro Lys Ser Thr Phe Thr Lys Ser Ile Thr Lys Ser Leu Leu Ala
 1 5 10 15
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Thr Ser Val Val Val Ser Leu Leu Pro Ala Tyr Ala Gln Ala
20 25 30

<210> 234
<211> 27
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 234
Met Leu Lys Arg Ile Thr Val Val Cys Leu Leu Phe Ile Leu Leu Phe
1 5 10 15
Pro Asn Ile Tyr Gly Arg Asn Lys Ala Glu Ala
20 25

<210> 235
<211> 29
<212> PRT
<213> Unknown

<220>
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<400> 235
Met Ser Leu Asn Asn Phe Lys Val Lys Leu Leu Ser Phe Ala Val Ser
1 5 10 15
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20 25

<210> 236
<211> 28
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 236
Met Ile Leu Ser Asn Phe Lys Val Lys Leu Leu Ser Phe Ala Val Ser
1 5 10 15
Ser Ala Val Leu Thr Leu Ala Ala Asn Val Ala Asn
20 25

<210> 237
<211> 27
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 237
Met Leu Lys Arg Ile Thr Val Val Cys Leu Leu Phe Ile Leu Leu Phe
1 5 10 15
Pro Asn Ile Tyr Glu Gly Asn Lys Ala Glu Ala
20 25

<210> 238
<211> 26
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 238

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Met Pro Ser Ile Asn Ala Ser Asp Cys Lys Lys Lys Gly Asp Arg Ser
 1 5 10 15
 Met Lys Arg Lys Lys Trp Thr Ala Leu Ala
 20 25

<210> 239
 <211> 33
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 239
 Val Ser Arg Met Phe Ala Lys Arg Phe Lys Thr Ser Leu Leu Pro Leu
 1 5 10 15
 Phe Ala Gly Phe Leu Leu Leu Phe His Leu Val Leu Ala Gly Pro Thr
 20 25 30
 Ala

<210> 240
 <211> 25
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 240
 Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
 1 5 10 15
 Trp Thr Ala Leu Ala Leu Thr Leu Pro
 20 25

<210> 241
 <211> 25
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

<400> 241
 Met Gln Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala
 1 5 10 15
 Leu Ala Leu Thr Leu Pro Leu Ala Ala
 20 25

<210> 242
 <211> 36
 <212> PRT
 <213> Bacterial

<400> 242
 Val Asp Pro Lys Asn Cys Ser Gln Phe Met Gln Thr Ile Ala Lys Lys
 1 5 10 15
 Gly Asp Glu Thr Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr
 20 25 30
 Leu Pro Leu Ala
 35

<210> 243
 <211> 36
 <212> PRT
 <213> Unknown

<220>

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<223> Obtained from an environmental sample.

<400> 243

Met	Gln	Thr	Ile	Ala	Lys	Lys	Gly	Asp	Glu	Thr	Met	Lys	Gly	Lys	Lys
1				5					10				15		
Trp	Thr	Ala	Leu	Ala	Leu	Thr	Leu	Pro	Leu	Ala	Ala	Ser	Leu	Ser	Thr
			20					25					30		
Gly	Val	His	Ala												
		35													

<210> 244

<211> 23

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 244

Met	Lys	Thr	Phe	Lys	Leu	Lys	Arg	Thr	Phe	Leu	Pro	Leu	Thr	Leu	Leu
1				5					10					15	
Leu	Ser	Ala	Pro	Ala	Phe	Ala									
			20												

<210> 245

<211> 25

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 245

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1				5					10				15		
Trp	Thr	Ala	Leu	Ala	Leu	Thr	Leu	Pro							
			20					25							

<210> 246

<211> 22

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 246

Met	Lys	Asn	Ile	Ile	Arg	Leu	Cys	Ala	Ala	Ser	Ala	Ile	Leu	Thr	Val
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Ser	His	Ala	Ser	Tyr	Ala										
			20												

<210> 247

<211> 23

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 247

Met	Lys	Thr	Phe	Lys	Leu	Lys	Arg	Thr	Phe	Leu	Pro	Leu	Thr	Leu	Leu
1				5					10					15	
Leu	Ser	Ala	Pro	Ala	Phe	Ala									
			20												

<210> 248

<211> 23

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<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 248
Met Lys Thr Phe Lys Leu Lys Arg Thr Phe Leu Pro Leu Thr Leu Leu
1 5 10 15
Leu Ser Ala Pro Ala Phe Ala
20

<210> 249
<211> 25
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 249
Met Lys Leu Met Lys Gly Lys Lys Trp Thr Ala Leu Ala Leu Thr Leu
1 5 10 15
Pro Leu Ala Ala Ser Leu Ser Thr Gly
20 25

<210> 250
<211> 36
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 250
Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
1 5 10 15
Trp Thr Ala Leu Ala Leu Thr Leu Pro Leu Ala Ala Ser Leu Ser Thr
20 25 30
Gly Val His Ala
35

<210> 251
<211> 25
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 251
Met Gln Thr Ile Ala Lys Lys Gly Asp Glu Thr Met Lys Gly Lys Lys
1 5 10 15
Trp Thr Ala Leu Ala Leu Thr Leu Pro
20 25

<210> 252
<211> 25
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 252
Met Lys Phe Lys Lys Ser Leu Ser Ala Gly Leu Leu Leu Phe Gly Gly
1 5 10 15
Leu Ser Gly Val Thr Pro Ser Val Ala

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20

25

<210> 253
<211> 23
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 253
Met Lys Pro Ser Lys Phe Val Phe Leu Ser Ala Ala Ile Ala Cys Ser
1 5 10 15
Leu Ser Ser Thr Ala Asn Ala
20

<210> 254
<211> 23
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 254
Val Ser Leu Thr Lys Lys Ala Gln Tyr Glu Pro Asn Thr Ala Pro Arg
1 5 10 15
Leu Ser Thr Ser Leu Gln Ser
20

<210> 255
<211> 30
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 255
Met Thr Ala Lys Ala Asp Asp Leu Arg Ile Tyr Gln Ile Met Val Glu
1 5 10 15
Ser Phe Val Asp Gly Asp Lys Gln Val Gly His Gly Thr Gly
20 25 30

<210> 256
<211> 25
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 256
Met Lys Met Lys Ser Arg Ala Trp Leu Leu Gly Ser Ala Val Ala Met
1 5 10 15
Ala Leu Ala Ser Ser Ala Ala Asn Ala
20 25

<210> 257
<211> 16
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample.

<400> 257
Met Tyr Arg Val Ile Pro Ile Ile Leu Ile Met Ser Met Ile Val Ala

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<211> 33			
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<211> 33			
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gaacgtctca ggcgctttga ctacgtgaag ggc			33
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<220>
<223> Primer

<400> 264
gaaccgtctc acgatataat ctggaacaag taccttgc 38

<210> 265
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<212> DNA
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<220>
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<400> 265
gaaccgtctc agaagcacga gcatagttta ctacg 35

<210> 266
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<220>
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<400> 266
gaaccgtctc aaaggtgggt ttatgtgccg 30

<210> 267
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<400> 267
gaacgtctca ggaatccaaa tggcggatat tcccgc 36

<210> 268
<211> 33
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<220>
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<400> 268
gaacggtctc agtttatcat attgatgagc tcc 33

<210> 269
<211> 33
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<220>
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<400> 269
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<210> 270
<211> 31

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<212> DNA
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<400> 270
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gaacggctctc aatcgggtgc gtggtttgct ac 32

<210> 273
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<220>
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<400> 273
gaaccgtctc acttcacact gcgaggtggt c 31

<210> 274
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<220>
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<400> 274
gaaccgtctc accttccaac cttgctcgag c 31

<210> 275
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<400> 275
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<210> 276
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<400> 276
gaacactagt aggaggtaac ttatggccaa gtacctggag ctcgaagagg 50

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 <210> 302
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<400> 307
gaacgtctca ttgtagtaga gggcgaagtc aaag 34

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<400> 308
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<400> 310
gaaccgtctc accttccggc ttgcttgag gc 32

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<400> 311
tcgagactga ctctcaccca acaccgcagt agctcc 36

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<400> 312
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<210> 313
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<400> 313

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 Met Ala Ala Val Ala Gln Pro Ala Ser Ala Ala Lys Tyr Ser Glu Leu
 20 25 30
 Glu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Ala
 35 40 45
 Gly Gly Ile Trp Trp Asp Thr Ile Arg Ser Lys Ile Pro Glu Trp Tyr
 50 55 60
 Glu Ala Gly Ile Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys Gly Met
 65 70 75 80
 Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu
 85 90 95
 Gly Glu Tyr Asn Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser Lys
 100 105 110
 Gln Glu Leu Ile Asn Met Ile Asn Thr Ala His Ala Tyr Gly Ile Lys
 115 120 125
 Val Ile Ala Asp Ile Val Ile Asn His Arg Ala Gly Asp Leu Glu
 130 135 140
 Trp Asn Pro Phe Val Gly Asp Tyr Thr Trp Thr Asp Phe Ser Lys Val
 145 150 155 160
 Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu
 165 170 175
 Val Lys Cys Cys Asp Glu Gly Thr Phe Gly Gly Phe Pro Asp Ile Ala
 180 185 190
 His Glu Lys Glu Trp Asp Gln His Trp Leu Trp Ala Ser Asp Glu Ser
 195 200 205
 Tyr Ala Ala Tyr Leu Arg Ser Ile Gly Val Asp Ala Trp Arg Phe Asp
 210 215 220
 Tyr Val Lys Gly Tyr Gly Ala Trp Val Val Lys Asp Trp Leu Asn Trp
 225 230 235 240
 Trp Gly Gly Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala
 245 250 255
 Leu Leu Asn Trp Ala Tyr Ser Ser Gly Ala Lys Val Phe Asp Phe Pro
 260 265 270
 Leu Tyr Tyr Lys Met Asp Glu Ala Phe Asp Asn Thr Asn Ile Pro Ala
 275 280 285
 Leu Val Asp Ala Leu Gln Asn Gly Gly Thr Val Val Ser Arg Asp Pro
 290 295 300
 Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Asp Ile Ile Trp
 305 310 315 320
 Asn Lys Tyr Pro Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro
 325 330 335
 Val Ile Phe Tyr Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Lys Leu
 340 345 350
 Asn Asn Leu Ile Trp Ile His Asp His Leu Ala Gly Gly Ser Thr Ser
 355 360 365
 Ile Val Tyr Tyr Asp Ser Asp Glu Leu Ile Phe Val Arg Asn Gly Asp
 370 375 380
 Ser Lys Arg Pro Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser Ser Lys
 385 390 395 400
 Val Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Ala Cys Ile His
 405 410 415
 Glu Tyr Thr Gly Asn Leu Gly Gly Trp Val Asp Lys Tyr Val Glu Ser
 420 425 430
 Ser Gly Trp Val Tyr Leu Glu Ala Pro Ala Tyr Asp Pro Ala Ser Gly
 435 440 445
 Gln Tyr Gly Tyr Thr Val Trp Ser Tyr Cys Gly Val Gly
 450 455 460

<210> 314

<211> 460

<212> PRT

<213> Pyrococcus furiosus

<400> 314

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Val 1	Asn	Ile	Lys	Lys 5	Leu	Thr	Pro	Leu	Leu 10	Thr	Leu	Leu	Leu	Phe 15	Phe
Ile	Val	Leu	Ala 20	Ser	Pro	Val	Ser	Ala 25	Ala	Lys	Tyr	Leu	Glu 30	Leu	Glu
Glu	Gly	Gly 35	Val	Ile	Met	Gln	Ala 40	Phe	Tyr	Trp	Asp	Val 45	Pro	Gly	Gly
Gly	Ile 50	Trp	Trp	Asp	His	Ile 55	Arg	Ser	Lys	Ile	Pro 60	Glu	Trp	Tyr	Glu
Ala 65	Gly	Ile	Ser	Ala	Ile 70	Trp	Leu	Pro	Pro	Pro 75	Ser	Lys	Gly	Met	Ser 80
Gly	Gly	Tyr	Ser	Met 85	Gly	Tyr	Asp	Pro	Tyr 90	Asp	Tyr	Phe	Asp	Leu 95	Gly
Glu	Tyr	Tyr	Gln 100	Lys	Gly	Thr	Val	Glu 105	Thr	Arg	Phe	Gly	Ser 110	Lys	Glu
Glu	Leu	Val 115	Arg	Leu	Ile	Gln	Thr 120	Ala	His	Ala	Tyr	Gly 125	Ile	Lys	Val
Ile 130	Ala	Asp	Val	Val	Ile	Asn 135	His	Arg	Ala	Gly	Gly 140	Asp	Leu	Glu	Trp
Asn 145	Pro	Phe	Val	Gly	Asp 150	Tyr	Thr	Trp	Thr	Asp 155	Phe	Ser	Lys	Val	Ala 160
Ser	Gly	Lys	Tyr	Thr 165	Ala	Asn	Tyr	Leu	Asp 170	Phe	His	Pro	Asn	Glu 175	Leu
His	Cys	Cys	Asp 180	Glu	Gly	Thr	Phe	Gly 185	Gly	Phe	Pro	Asp	Ile 190	Cys	His
His	Lys	Glu 195	Trp	Asp	Gln	Tyr	Trp 200	Leu	Trp	Lys	Ser	Asn 205	Glu	Ser	Tyr
Ala 210	Ala	Tyr	Leu	Arg	Ser	Ile 215	Gly	Phe	Asp	Gly	Trp 220	Arg	Phe	Asp	Tyr
Val 225	Lys	Gly	Tyr	Gly	Ala 230	Trp	Val	Val	Arg	Asp 235	Trp	Leu	Asn	Trp	Trp 240
Gly	Gly	Trp	Ala	Val 245	Gly	Glu	Tyr	Trp	Asp 250	Thr	Asn	Val	Asp	Ala 255	Leu
Leu	Ser	Trp	Ala 260	Tyr	Glu	Ser	Gly	Ala 265	Lys	Val	Phe	Asp	Phe 270	Pro	Leu
Tyr	Tyr	Lys 275	Met	Asp	Glu	Ala	Phe 280	Asp	Asn	Asn	Asn	Ile 285	Pro	Ala	Leu
Val 290	Tyr	Ala	Leu	Gln	Asn	Gly 295	Gln	Thr	Val	Val	Ser 300	Arg	Asp	Pro	Phe
Lys 305	Ala	Val	Thr	Phe	Val 310	Ala	Asn	His	Asp	Thr 315	Asp	Ile	Ile	Trp	Asn 320
Lys	Tyr	Pro	Ala	Tyr 325	Ala	Phe	Ile	Leu	Thr 330	Tyr	Glu	Gly	Gln	Pro 335	Val
Ile	Phe	Tyr	Arg 340	Asp	Phe	Glu	Glu	Trp 345	Leu	Asn	Lys	Asp	Lys 350	Leu	Ile
Asn	Leu	Ile 355	Trp	Ile	His	Asp	His 360	Leu	Ala	Gly	Gly	Ser 365	Thr	Thr	Ile
Val 370	Tyr	Tyr	Asp	Asn	Asp	Glu 375	Leu	Ile	Phe	Val	Arg 380	Asn	Gly	Asp	Ser
Arg 385	Arg	Pro	Gly	Leu	Ile 390	Thr	Tyr	Ile	Asn	Leu 395	Ser	Pro	Asn	Trp	Val 400
Gly	Arg	Trp	Val	Tyr 405	Val	Pro	Lys	Phe	Ala 410	Gly	Ala	Cys	Ile	His 415	Glu
Tyr	Thr	Gly	Asn 420	Leu	Gly	Gly	Trp	Val 425	Asp	Lys	Arg	Val	Asp 430	Ser	Ser
Gly	Trp	Val 435	Tyr	Leu	Glu	Ala	Pro 440	Pro	His	Asp	Pro	Ala 445	Asn	Gly	Tyr
Tyr	Gly 450	Tyr	Ser	Val	Trp	Ser 455	Tyr	Cys	Gly	Val	Gly 460				

<210> 315

<211> 484

<212> PRT

<213> Thermococcus sp.

<400> 315

Ser 1	Glu	Ser	Gln	Cys 5	Thr	Ala	Thr	Cys	Thr 10	Trp	Arg	Val	Val	Tyr 15	Met
Ser	Ala	Lys	Lys	Leu	Leu	Ala	Leu	Leu	Phe	Val	Leu	Ala	Val	Leu	Val

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Gly	Val	Ala	20	Val	Ile	Pro	Ala	Arg	25	Val	Gly	Ile	Ala	Pro	30	Val	Ser	Ala
Gly	Ala	Thr	35	Ser	Arg	Pro	Ser	Leu	40	Glu	Glu	Gly	Gly	Val	Ile	Met	Gln	
Ala	Phe	Tyr	50	Trp	Asp	Val	Pro	Ala	55	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	
Arg	Ser	Lys	65	Ile	Pro	Asp	Trp	Ala	70	Ser	Ala	Gly	Ile	Ser	Ala	Ile	Trp	
Ile	Pro	Pro	85	Ala	Ser	Lys	Gly	Met	90	Gly	Ala	Tyr	Ser	Met	Gly	Tyr		
Asp	Pro	Tyr	100	Asp	Phe	Phe	Asp	Leu	105	Gly	Glu	Tyr	Tyr	Gln	Lys	Gly	Thr	
Val	Glu	Thr	115	Arg	Phe	Gly	Ser	Lys	120	Gln	Glu	Leu	Ile	Asn	Met	Ile	Asn	
Thr	Ala	His	130	Ser	Tyr	Gly	Ile	Lys	135	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	
His	Arg	Ala	145	Gly	Gly	Asp	Leu	Glu	150	Trp	Asn	Pro	Phe	Thr	Asn	Ser	Tyr	
Thr	Trp	Thr	165	Asp	Phe	Ser	Lys	Val	170	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	
Tyr	Leu	Asp	180	Phe	His	Pro	Asn	Glu	185	Val	Lys	Cys	Cys	Asp	Glu	Gly	Thr	
Phe	Gly	Gly	195	Phe	Pro	Asp	Ile	Ala	200	His	Glu	Lys	Ser	Trp	Asp	Gln	Tyr	
Trp	Leu	Trp	210	Ala	Ser	Gln	Lys	Ser	215	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	
Gly	Ile	Asp	225	Ala	Trp	Arg	Phe	Asp	230	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	
Val	Val	Lys	245	Asp	Trp	Leu	Lys	Trp	250	Ala	Leu	Ala	Val	Gly	Glu	Tyr		
Trp	Asp	Thr	260	Asn	Val	Asp	Ala	Leu	265	Asn	Trp	Ala	Tyr	Ser	Ser	Gly		
Ala	Lys	Val	275	Phe	Asp	Phe	Pro	Leu	280	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	
Asp	Asn	Lys	290	Asn	Ile	Pro	Ala	Leu	295	Val	Ser	Ala	Leu	Gln	Asn	Gly	Gln	
Thr	Val	Val	305	Ser	Arg	Asp	Pro	Phe	310	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn	
His	Asp	Thr	325	Asp	Ile	Ile	Trp	Asn	330	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile	
Leu	Thr	Tyr	340	Glu	Gly	Gln	Pro	Val	345	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu	
Trp	Leu	Asn	355	Lys	Asp	Arg	Leu	Lys	360	Asn	Leu	Ile	Trp	Ile	His	Asn	Asn	
Leu	Ala	Gly	370	Gly	Ser	Thr	Ser	Ile	375	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu	
Ile	Phe	Val	385	Arg	Asn	Gly	Tyr	Gly	390	Asn	Lys	Pro	Gly	Leu	Ile	Thr	Tyr	
Ile	Asn	Leu	405	Gly	Ser	Ser	Lys	Val	410	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys	
Phe	Ala	Gly	420	Ser	Cys	Ile	His	Glu	425	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp	
Val	Asp	Lys	435	Tyr	Val	Gly	Ser	Asn	440	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro	
Ala	His	Asp	450	Pro	Ala	Lys	Gly	Gln	455	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr	
Cys	Gly	Val	465	Gly					470									

<210> 316
 <211> 457
 <212> PRT
 <213> Thermococcus hydrothermalis

<400> 316
 Met Ala Arg Lys Val Leu Val Ala Leu Leu Val Phe Leu Val Val Leu
 1 5 10 15

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Ser Val Ser Ala Val Pro Ala Lys Ala Glu Thr Leu Glu Asn Gly Gly
20 25 30
Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro Gly Gly Ile Trp
35 40 45
Trp Asp Thr Ile Ala Gln Lys Ile Pro Asp Trp Ala Ser Ala Gly Ile
50 55 60
Ser Ala Ile Trp Ile Pro Pro Ala Ser Lys Gly Met Ser Gly Gly Tyr
65 70 75 80
Ser Met Gly Tyr Asp Pro Tyr Asp Phe Phe Asp Leu Gly Glu Tyr Tyr
85 90 95
Gln Lys Gly Ser Val Glu Thr Arg Phe Gly Ser Lys Glu Glu Leu Val
100 105 110
Asn Met Ile Asn Thr Ala His Ala His Asn Met Lys Val Ile Ala Asp
115 120 125
Ile Val Ile Asn His Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe
130 135 140
Thr Asn Ser Tyr Thr Trp Thr Asp Phe Ser Lys Val Ala Ser Gly Lys
145 150 155 160
Tyr Thr Ala Asn Tyr Leu Asp Phe His Pro Asn Glu Leu His Ala Gly
165 170 175
Asp Ser Gly Thr Phe Gly Gly Tyr Pro Asp Ile Cys His Asp Lys Ser
180 185 190
Trp Asp Gln His Trp Leu Trp Ala Ser Asn Glu Ser Tyr Ala Ala Tyr
195 200 205
Leu Arg Ser Ile Gly Ile Asp Ala Trp Arg Phe Asp Tyr Val Lys Gly
210 215 220
Tyr Ala Pro Trp Val Val Lys Asn Trp Leu Asn Arg Trp Gly Gly Trp
225 230 235 240
Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu Ser Trp
245 250 255
Ala Tyr Asp Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys
260 265 270
Met Asp Glu Ala Phe Asp Asn Asn Asn Ile Pro Ala Leu Val Asp Ala
275 280 285
Leu Lys Asn Gly Gly Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val
290 295 300
Thr Phe Val Ala Asn His Asp Thr Asn Ile Ile Trp Asn Lys Tyr Pro
305 310 315 320
Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro Ala Ile Phe Tyr
325 330 335
Arg Asp Tyr Glu Glu Trp Leu Asn Lys Asp Arg Leu Arg Asn Leu Ile
340 345 350
Trp Ile His Asp His Leu Ala Gly Ser Thr Asp Ile Ile Tyr Tyr
355 360 365
Asp Ser Asp Glu Leu Ile Phe Val Arg Asn Gly Tyr Gly Asp Lys Pro
370 375 380
Gly Leu Ile Thr Tyr Ile Asn Leu Gly Ser Ser Lys Ala Gly Arg Trp
385 390 395 400
Val Tyr Val Pro Lys Phe Ala Gly Ser Cys Ile His Glu Tyr Thr Gly
405 410 415
Asn Leu Gly Gly Trp Ile Asp Lys Trp Val Asp Ser Ser Gly Arg Val
420 425 430
Tyr Leu Glu Ala Pro Ala His Asp Pro Ala Asn Gly Gln Tyr Gly Tyr
435 440 445
Ser Val Trp Ser Tyr Cys Gly Val Gly
450 455

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<210> 317

<211> 340

<212> PRT

<213> Artificial Sequence

<220>

<223> Consensus sequence

<400> 317

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Ser Ala Leu Glu Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val
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Pro Gly Gly Ile Trp Trp Asp Ile Lys Ile Pro Trp Ala Gly Ile Ser
 20 25 30
 Ala Ile Trp Pro Pro Ser Lys Gly Met Gly Tyr Ser Met Gly Tyr Asp
 35 40 45
 Pro Tyr Asp Phe Asp Leu Gly Glu Tyr Gln Lys Gly Glu Thr Arg Phe
 50 55 60
 Gly Ser Lys Glu Leu Ile Thr Ala His Lys Val Ile Ala Asp Val Ile
 65 70 75 80
 Asn His Arg Ala Gly Gly Asp Leu Glu Trp Asn Pro Phe Tyr Thr Trp
 85 90 95
 Thr Asp Phe Ser Lys Val Ala Ser Gly Lys Tyr Thr Ala Asn Tyr Leu
 100 105 110
 Asp Phe His Pro Asn Glu Asp Gly Thr Phe Gly Gly Pro Asp Ile His
 115 120 125
 Lys Trp Asp Gln Trp Leu Trp Ser Ser Tyr Ala Ala Tyr Leu Arg Ser
 130 135 140
 Ile Gly Asp Trp Arg Phe Asp Tyr Val Lys Gly Tyr Trp Val Val Trp
 145 150 155 160
 Leu Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Leu
 165 170 175
 Trp Ala Tyr Ser Gly Ala Lys Val Phe Asp Phe Pro Leu Tyr Tyr Lys
 180 185 190
 Met Asp Glu Ala Phe Asp Asn Asn Ile Pro Ala Leu Val Ala Leu Asn
 195 200 205
 Gly Thr Val Val Ser Arg Asp Pro Phe Lys Ala Val Thr Phe Val Ala
 210 215 220
 Asn His Asp Thr Ile Ile Trp Asn Lys Tyr Ala Tyr Ala Phe Ile Leu
 225 230 235 240
 Thr Tyr Glu Gly Gln Pro Ile Phe Tyr Arg Asp Glu Glu Trp Leu Asn
 245 250 255
 Lys Asp Leu Asn Leu Ile Trp Ile His Leu Ala Gly Gly Ser Thr Ile
 260 265 270
 Tyr Tyr Asp Asp Glu Ile Phe Val Arg Asn Gly Pro Gly Leu Ile Thr
 275 280 285
 Tyr Ile Asn Leu Gly Arg Trp Val Tyr Val Pro Lys Phe Ala Gly Cys
 290 295 300
 Ile His Glu Tyr Thr Gly Asn Leu Gly Gly Trp Asp Lys Val Ser Gly
 305 310 315 320
 Val Tyr Glu Ala Pro Asp Pro Ala Gly Tyr Gly Tyr Val Trp Ser Tyr
 325 330 335
 Cys Gly Val Gly
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<210> 318
 <211> 463
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> clone

<400> 318
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 1 5 10 15
 Leu Ala Gly Ile Tyr Tyr Pro Ser Thr Ala Ala Lys Tyr Ser Glu
 20 25 30
 Leu Glu Gln Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro
 35 40 45
 Glu Gly Gly Ile Trp Trp Asp Thr Ile Arg Gln Lys Ile Pro Glu Trp
 50 55 60
 Tyr Asp Ala Gly Ile Ser Ala Ile Trp Ile Pro Ala Ser Lys Gly
 65 70 75 80
 Met Gly Gly Ala Tyr Ser Met Gly Tyr Asp Pro Tyr Asp Tyr Phe Asp
 85 90 95
 Leu Gly Glu Phe Tyr Gln Lys Gly Thr Val Glu Thr Arg Phe Gly Ser
 100 105 110
 Lys Glu Glu Leu Val Asn Met Ile Ser Thr Ala His Gln Tyr Gly Ile
 115 120 125

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Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His	Arg	Ala	Gly	Gly	Asp	Leu
130	130					135					140				
Glu	Trp	Asn	Pro	Tyr	Val	Gly	Asp	Tyr	Thr	Trp	Thr	Asp	Phe	Ser	Lys
145					150					155					160
Val	Ala	Ser	Gly	Lys	Tyr	Lys	Ala	His	Tyr	Met	Asp	Phe	His	Pro	Asn
				165					170					175	
Asn	Tyr	Ser	Thr	Ser	Asp	Glu	Gly	Thr	Phe	Gly	Gly	Phe	Pro	Asp	Ile
			180					185					190		
Asp	His	Leu	Val	Pro	Phe	Asn	Gln	Tyr	Trp	Leu	Trp	Ala	Ser	Asn	Glu
		195					200					205			
Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly	Ile	Asp	Ala	Trp	Arg	Phe
	210					215					220				
Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val	Val	Lys	Asp	Trp	Leu	Ser
225					230					235					240
Gln	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	Trp	Asp	Thr	Asn	Val	Asp
				245					250					255	
Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Gly	Ala	Lys	Val	Phe	Asp	Phe
			260					265					270		
Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	Asp	Asn	Lys	Asn	Ile	Pro
		275					280					285			
Ala	Leu	Val	Tyr	Ala	Ile	Gln	Asn	Gly	Glu	Thr	Val	Val	Ser	Arg	Asp
	290					295					300				
Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn	His	Asp	Thr	Asn	Ile	Ile
305					310					315					320
Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile	Leu	Thr	Tyr	Glu	Gly	Gln
				325					330					335	
Pro	Val	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu	Trp	Leu	Asn	Lys	Asp	Lys
			340					345					350		
Leu	Asn	Asn	Leu	Ile	Trp	Ile	His	Glu	His	Leu	Ala	Gly	Gly	Ser	Thr
		355					360					365			
Lys	Ile	Leu	Tyr	Tyr	Asp	Asp	Glu	Leu	Ile	Phe	Met	Arg	Glu	Gly	
	370					375				380					
Tyr	Gly	Asp	Arg	Pro	Gly	Leu	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	Ser	Asp
385					390					395					400
Trp	Ala	Glu	Arg	Trp	Val	Asn	Val	Gly	Ser	Lys	Phe	Ala	Gly	Tyr	Thr
				405					410					415	
Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Trp	Val	Asp	Arg	Tyr	Val	
			420					425					430		
Gln	Tyr	Asp	Gly	Trp	Val	Lys	Leu	Thr	Ala	Pro	Pro	His	Asp	Pro	Ala
		435					440					445			
Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr	Ala	Gly	Val	Gly	
	450					455					460				

<210> 319
 <211> 306
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Consensus sequence

<400> 319

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Trp	Asp	Ile	Lys	Ile	Pro	Trp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Pro	Pro
			20					25					30		
Ser	Lys	Gly	Met	Gly	Tyr	Ser	Met	Gly	Tyr	Asp	Pro	Tyr	Asp	Phe	Asp
		35					40					45			
Leu	Gly	Glu	Gln	Lys	Gly	Glu	Thr	Arg	Phe	Gly	Ser	Lys	Glu	Leu	Ile
	50					55					60				
Thr	Ala	His	Lys	Val	Ile	Ala	Asp	Val	Ile	Asn	His	Arg	Ala	Gly	Gly
65					70					75					80
Leu	Glu	Trp	Asn	Pro	Tyr	Thr	Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser
			85						90					95	
Gly	Lys	Tyr	Ala	Tyr	Asp	Phe	His	Pro	Asn	Asp	Gly	Thr	Phe	Gly	Gly
			100					105					110		
Pro	Asp	Ile	His	Gln	Trp	Leu	Trp	Ser	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg
		115					120					125			

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Ser Ile Gly Asp Trp Phe Asp Tyr Val Lys Gly Tyr Trp Val Val Trp
 130 135 140
 Leu Trp Ala Val Gly Glu Tyr Trp Asp Thr Asn Val Asp Ala Leu Trp
 145 150 155 160
 Ala Tyr Ser Ala Lys Val Phe Asp Phe Leu Tyr Tyr Lys Met Asp Ala
 165 170 175
 Phe Asp Asn Asn Ile Pro Ala Leu Val Ala Gly Thr Val Val Ser Arg
 180 185 190
 Asp Pro Phe Lys Ala Val Thr Phe Val Ala Asn His Asp Thr Ile Ile
 195 200 205
 Trp Asn Lys Tyr Ala Tyr Ala Phe Ile Leu Thr Tyr Glu Gly Gln Pro
 210 215 220
 Ile Phe Tyr Arg Asp Glu Glu Trp Leu Asn Lys Asp Leu Asn Leu Ile
 225 230 235 240
 Trp Ile His Leu Ala Gly Gly Ser Thr Ile Tyr Tyr Asp Asp Glu Ile
 245 250 255
 Phe Arg Gly Pro Gly Leu Ile Thr Tyr Ile Asn Leu Arg Trp Val Val
 260 265 270
 Lys Phe Ala Gly Ile His Glu Tyr Thr Gly Leu Gly Gly Trp Asp Val
 275 280 285
 Gly Val Ala Pro Asp Pro Ala Gly Tyr Gly Tyr Val Trp Ser Tyr Gly
 290 295 300
 Val Gly
 305

<210> 320
 <211> 1392
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Clone

<400> 320
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 caggccttct actgggacgt tccggagggg ggaatctggg gggacacaa acggcagaag 180
 atccctgaat ggtacgatgc aggcataatcc gccatctgga taccctcggc gagcaagggc 240
 atgggcgggg cctactcgat gggctacgac ccctacgatt acttcgatct gggcgagttt 300
 taccagaagg gaaccgttga gacccgcttc ggctccaagg aagagctcgt caacatgatc 360
 tccacggccc accagtacgg catcaagggtt atagcggaca tagtgataaa ccaccgcgca 420
 ggtggagacc tcgaatggaa cccatacgtc ggcgactata cctggacgga cttttctaag 480
 gtcgcctccg ggaaatacaa ggcccactac atggacttcc atccaaacaa ctacagcacc 540
 tcagacgagg gaaccttcgg tggcttccca gacattgatc acctcgtgcc cttcaaccag 600
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 gcgtggcgct ttgactacgt taagggttac ggcgcgtggg tcgtcaagga ctggctgagt 720
 cagtggggcg gctgggccgt cggcgagtac tgggacacca acgtcgatgc gtcctcaac 780
 tgggcttaca gcagcggcgc caaggtcttc gacttcccgc tctactacaa gatggacgag 840
 gcctttgaca acaagaacat tcccgccttc gtttacgcca tccagaacgg tgaaaccgtc 900
 gtcagcaggg atcccttcaa ggccgttacc ttcgtggcta accacgatac gaacataatc 960
 tgaacaagt accctgccta tgccttcata ctgacctacg aaggtcagcc cgtcatcttc 1020
 taccgcgact acgaggagt gctcaacaag gacaaactca acaacctcat atggattcac 1080
 gagcacctgg cagggggaag caccaagatc ctctactacg acgacgatga gctcatcttc 1140
 atgaggggaag gctacggcga caggccccgg cttataacct acatcaacct cggtagcgac 1200
 tgggcggaga gatgggtgaa cgttggtcca aagttcgcgg gctatacaat ccacgaatac 1260
 accggaaacc tcggcggctg ggtcgacagg tacgtccagt acgacggctg ggtcaagctt 1320
 accgctccgc cacacgatcc ggcaaacggc tattacggct actcggctct gagctacgcc 1380
 ggagttggat ga 1392

<210> 321
 <211> 846
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Consensus sequence

<400> 321

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atgggcggaat tcgcattgga tccccgcaga agggatggcg ggtatcatgg gctacgaccc      120
tacgatttga ctgggagtag agaaggatg aacgttggtc aaagacttaa atgatacgc      180
caacataagg tatgcgaatg tataaccacc gcgcgggggc tgatggaacc tcctaacctg      240
gacgatttca agtgctcggg ataagccact actgacttcc accaacagag gacttggggg      300
ccgaatcacc taccagactg gcttgggcag cgaagctacg cgctactgga gcatgggtgag      360
ctgggttgac tacgtaaggg ctagctgggt gtaactggct gtggggggct gggcgtggga      420
gtactgggac acaacgtgag ctctactggg ctacagcggc aagtcttgac ttccctctac      480
tacaagatgg aggccttgaa aaaacatccg ctgtgctagg acgtgtccgg accttaagcg      540
tacttgtgca accacgaaca ataattggaa caagtacgct agcttcatcc tacctagagg      600
cagccatttt accgcgacta cgaggagtgg ctcaacaagg aataaacctc attggatcag      660
aacctgcggg gagacattta ctacgacgag atattttaga ggctaggcac cggctataac      720
taatcaacct gcggagtggg tagtcaagtt cgcgatcac gataacggac tcgggggtgg      780
gacagtgttc gggggtatgc cccacgaccg caaggatagg ctactcgttg gagctacggg      840
tggatga                                           846

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<210> 322
 <211> 1392
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 322
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tactaccctt ccacgagtgc cgcgaagtac tccgagctgg agcagggcgg agtcataatg      120
caggccttct actgggacgt tccggagggg ggaatctggg gggacacaaat acggcagaag      180
atccctgaat ggtacgatgc aggcataatcc gccatctgga taccctcggc gagcaagggc      240
atgggcgggg cctactcgat gggctacgac ccctacgatt acttcgatct gggcggagttt      300
taccagaagg gaaccgttga gaccgccttc ggctccaagg aagagctcgt caacatgac      360
tccacggccc accagtacgg catcaagggt atagcggaca tagtgataaa ccaccgcgca      420
ggtgggagacc tcgaatggaa cccatacgtc ggcgactata cctggacgga cttttctaag      480
gtcgccctcc ggaaatacaa ggcccactac atggacttcc atccaaacaa ctacagcacc      540
tcagacgagg gaaccttcgg tggcttccca gacattgatc acctcgtgcc ctccaaccag      600
tactggctgt gggcgagcaa cgagagctac gccgcctacc tcaggagcat agggatcgat      660
gcgtggcgct ttgactacgt taagggtac ggcgctgggg tcgtcaagga ctggctgagt      720
cagtggggcg gctgggccgt cggcgagtac tgggacacca acgtcgtatg gctcctcaac      780
tgggcctaca gcagcggcgc caaggtcttc gacttccgcg tctactacaa gatggacgag      840
gcctttgaca acaagaacat tcccgccttc gtttacgcca tccagaacgg tgaaaccgct      900
gtcagcaggg atcccttcaa ggccgttacc ttcgtggcta accacgatac gaacataatc      960
tggaacaagt accctgccta tgccttcata ctgacctacg aaggtcagcc cgtcatcttc      1020
taccgcgact acgaggagtg gctcaacaag gacaaactca acaacctcat atggattcac      1080
gagcacctgg cagggggaag caccaagatc ctctactacg acgacgatga gctcatcttc      1140
atgaggggaa gctacggcga caggcccggg cttataacct acatcaacct cggtagcgac      1200
tgggcggaga gatgggtgaa cgttggctca aagtctcgcg gctatacaat ccacgaatac      1260
accggaaacc tcggcggctg ggtcgacagg tacgtccagt acgacggctg ggtcaagctt      1320
accgctccgc cacacgatcc ggcaaacggc tattacggct actcggctct gagctacgcc      1380
ggagtggat ga                                           1392

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<210> 323
 <211> 463
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(27)

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<400> 323
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Leu Ala Gly Ile Tyr Tyr Pro Ser Thr Ser Ala Ala Lys Tyr Ser Glu
      20           25           30
Leu Glu Gln Gly Gly Val Ile Met Gln Ala Phe Tyr Trp Asp Val Pro
 35           40           45

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Glu	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg	Gln	Lys	Ile	Pro	Glu	Trp
50	50	50	50	50	50	55	55	55	55	55	60	60	60	60	60
Tyr	Asp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile	Pro	Pro	Ala	Ser	Lys	Gly
65	65	65	65	65	70	70	70	70	70	75	75	75	75	75	80
Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr	Asp	Pro	Tyr	Asp	Tyr	Phe	Asp
				85					90					95	
Leu	Gly	Glu	Phe	Tyr	Gln	Lys	Gly	Thr	Val	Glu	Thr	Arg	Phe	Gly	Ser
			100					105					110		
Lys	Glu	Glu	Leu	Val	Asn	Met	Ile	Ser	Thr	Ala	His	Gln	Tyr	Gly	Ile
		115					120					125			
Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His	Arg	Ala	Gly	Gly	Asp	Leu
	130					135					140				
Glu	Trp	Asn	Pro	Tyr	Val	Gly	Asp	Tyr	Thr	Trp	Thr	Asp	Phe	Ser	Lys
145					150					155					160
Val	Ala	Ser	Gly	Lys	Tyr	Lys	Ala	His	Tyr	Met	Asp	Phe	His	Pro	Asn
				165					170					175	
Asn	Tyr	Ser	Thr	Ser	Asp	Glu	Gly	Thr	Phe	Gly	Gly	Phe	Pro	Asp	Ile
			180					185					190		
Asp	His	Leu	Val	Pro	Phe	Asn	Gln	Tyr	Trp	Leu	Trp	Ala	Ser	Asn	Glu
		195					200					205			
Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly	Ile	Asp	Ala	Trp	Arg	Phe
	210					215					220				
Asp	Tyr	Val	Lys	Gly	Tyr	Gly	Ala	Trp	Val	Val	Lys	Asp	Trp	Leu	Ser
225					230					235					240
Gln	Trp	Gly	Gly	Trp	Ala	Val	Gly	Glu	Tyr	Trp	Asp	Thr	Asn	Val	Asp
				245					250					255	
Ala	Leu	Leu	Asn	Trp	Ala	Tyr	Ser	Ser	Gly	Ala	Lys	Val	Phe	Asp	Phe
			260					265					270		
Pro	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe	Asp	Asn	Lys	Asn	Ile	Pro
		275					280					285			
Ala	Leu	Val	Tyr	Ala	Ile	Gln	Asn	Gly	Glu	Thr	Val	Val	Ser	Arg	Asp
	290					295					300				
Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn	His	Asp	Thr	Asn	Ile	Ile
305					310					315					320
Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile	Leu	Thr	Tyr	Glu	Gly	Gln
				325					330					335	
Pro	Val	Ile	Phe	Tyr	Arg	Asp	Tyr	Glu	Glu	Trp	Leu	Asn	Lys	Asp	Lys
			340					345					350		
Leu	Asn	Asn	Leu	Ile	Trp	Ile	His	Glu	His	Leu	Ala	Gly	Gly	Ser	Thr
		355					360					365			
Lys	Ile	Leu	Tyr	Tyr	Asp	Asp	Asp	Glu	Leu	Ile	Phe	Met	Arg	Glu	Gly
	370					375					380				
Tyr	Gly	Asp	Arg	Pro	Gly	Leu	Ile	Thr	Tyr	Ile	Asn	Leu	Gly	Ser	Asp
385					390					395					400
Trp	Ala	Glu	Arg	Trp	Val	Asn	Val	Gly	Ser	Lys	Phe	Ala	Gly	Tyr	Thr
				405					410					415	
Ile	His	Glu	Tyr	Thr	Gly	Asn	Leu	Gly	Gly	Trp	Val	Asp	Arg	Tyr	Val
			420					425					430		
Gln	Tyr	Asp	Gly	Trp	Val	Lys	Leu	Thr	Ala	Pro	Pro	His	Asp	Pro	Ala
		435					440					445			
Asn	Gly	Tyr	Tyr	Gly	Tyr	Ser	Val	Trp	Ser	Tyr	Ala	Gly	Val	Gly	
	450					455					460				

<210> 324
 <211> 1296
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 324	
atgaccagca gcctgaaaac caccagcata cccttcatct acaacatttt cccgtcgctc	60
ctcggcgagg tcgacaagtg gattccgcat gtcgagcgtg cggtggaaat gggcttcaac	120
tggatattca tcaatcccat tcatttgacc ggcgccagcg gatcgctgta cgccatccgc	180
gactatttcc agttcaaccc ggcgtttttc ccctacgact cgttcgacgc gcagaaaaac	240
cgtctgtcgg cgtttgtctc gcagtgccgc gagatggggg cggaggtgat gatcgatctg	300
gtcatcaacc ataccgccta cgacaacccc ctgaccgcgc atcaccgcaa ctggtacaag	360

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ctgaacgccg	acggttcgat	aaaaaatccc	ggcgccaggg	acgacagcgc	gcccggcggg	420
tatgtgggtt	ggggcgactt	gaacgagata	gacaacgagc	attccccgga	ccggaacaat	480
ttgtgggaat	attggtgggtc	ggtggtggac	atgtatctgt	cctgcggcgt	gcgcggattc	540
cgttgcgacg	cggcttatca	gattccgaac	gacttgtggc	ggctgcttat	agaacgcgcc	600
aagaacaagc	attccgaatg	ccgtttcttc	gcggagtctc	tcggctgtcc	catggaacag	660
accgtcaacc	tggcgaaagc	gggccacgac	ttcgttttca	actccggcaa	gtggtgggat	720
tatcgccagg	actggtttat	cgatcagatg	cgcattggtt	ccggccaggc	gtcttcgatt	780
tgtttccccg	agtcaccacg	caccgaacgt	ttggcggcgg	agtggggctc	ggatttggag	840
cggatcaagc	agcattatct	tttcagcgcg	ttgatatacgt	ccggcgtcct	catccttctc	900
ggcttcgagt	acggcttcat	gaaaaagcca	catgtcgtgc	actcgcattc	gcacgattac	960
gagggcgcg	attacgattt	gaccgactac	atccgggaaa	tcaattccct	caagcgcgcc	1020
caccgggtat	tcgccgagga	caataaattc	gaacggcttg	attccggaga	cgacggcgtg	1080
ctggcgctgc	ggaagacgac	gctggacggc	aaggagagcg	gcctgcttct	cttcaaccgt	1140
accggcgatc	gcaaaaccct	tcacctgggc	gacttgatca	accgcatgca	tgccaaccgc	1200
taccgtttca	ccgcggccgc	caacgtcaag	ctccccgcca	aggccgagcg	tatcgagatc	1260
aaccgggtatg	gcgcgggtggc	gttcgggttg	tattga			1296

<210> 325

<211> 431

<212> PRT

<213> Unknown

<220>

<223> obtained from an environmental sample

<400> 325

Met	Thr	Ser	Ser	Leu	Lys	Thr	Thr	Ser	Ile	Pro	Phe	Ile	Tyr	Asn	Ile
1				5				10						15	
Phe	Pro	Ser	Leu	Leu	Gly	Glu	Val	Asp	Lys	Trp	Ile	Pro	His	Val	Glu
			20					25					30		
Arg	Ala	Val	Glu	Met	Gly	Phe	Asn	Trp	Ile	Phe	Ile	Asn	Pro	Ile	His
		35					40					45			
Leu	Thr	Gly	Ala	Ser	Gly	Ser	Leu	Tyr	Ala	Ile	Arg	Asp	Tyr	Phe	Gln
	50					55					60				
Phe	Asn	Pro	Ala	Phe	Phe	Pro	Tyr	Asp	Ser	Phe	Asp	Ala	Gln	Lys	Asn
65				70						75				80	
Arg	Leu	Ser	Ala	Phe	Val	Ser	Gln	Cys	Arg	Glu	Met	Gly	Ala	Glu	Val
			85					90						95	
Met	Ile	Asp	Leu	Val	Ile	Asn	His	Thr	Ala	Tyr	Asp	Asn	Pro	Leu	Thr
			100					105					110		
Arg	Asp	His	Arg	Asn	Trp	Tyr	Lys	Leu	Asn	Ala	Asp	Gly	Ser	Ile	Lys
		115					120					125			
Asn	Pro	Gly	Ala	Arg	Asp	Asp	Ser	Ala	Pro	Gly	Gly	Tyr	Val	Val	Trp
	130					135					140				
Gly	Asp	Leu	Asn	Glu	Ile	Asp	Asn	Glu	His	Ser	Pro	Asp	Arg	Asn	Asn
145				150						155				160	
Leu	Trp	Glu	Tyr	Trp	Trp	Ser	Val	Val	Asp	Met	Tyr	Leu	Ser	Cys	Gly
			165					170						175	
Val	Arg	Gly	Phe	Arg	Cys	Asp	Ala	Ala	Tyr	Gln	Ile	Pro	Asn	Asp	Leu
			180					185					190		
Trp	Arg	Leu	Leu	Ile	Glu	Arg	Ala	Lys	Asn	Lys	His	Ser	Glu	Cys	Arg
		195					200					205			
Phe	Phe	Ala	Glu	Ser	Leu	Gly	Cys	Pro	Met	Glu	Gln	Thr	Val	Asn	Leu
	210					215					220				
Ala	Lys	Ala	Gly	His	Asp	Phe	Val	Phe	Asn	Ser	Gly	Lys	Trp	Trp	Asp
225				230						235				240	
Tyr	Arg	Gln	Asp	Trp	Phe	Ile	Asp	Gln	Met	Arg	Met	Val	Ser	Gly	Gln
			245					250						255	
Ala	Ser	Ser	Ile	Cys	Phe	Pro	Glu	Ser	His	Asp	Thr	Glu	Arg	Leu	Ala
			260					265					270		
Ala	Glu	Trp	Gly	Arg	Asp	Leu	Glu	Arg	Ile	Lys	Gln	His	Tyr	Leu	Phe
		275					280					285			
Ser	Ala	Leu	Ile	Ser	Ser	Gly	Val	Leu	Ile	Leu	Leu	Gly	Phe	Glu	Tyr
	290					295					300				
Gly	Phe	Met	Lys	Lys	Pro	His	Val	Val	His	Ser	His	Pro	His	Asp	Tyr
305					310					315				320	
Glu	Gly	Ala	Asn	Tyr	Asp	Leu	Thr	Asp	Tyr	Ile	Arg	Glu	Ile	Asn	Ser
			325						330					335	

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Leu Lys Arg Ala His Arg Val Phe Ala Glu Asp Asn Lys Phe Glu Arg
 340 345 350
 Leu Asp Ser Gly Asp Asp Gly Val Leu Ala Leu Arg Lys Thr Thr Leu
 355 360 365
 Asp Gly Lys Glu Ser Gly Leu Leu Leu Phe Asn Arg Thr Gly Asp Arg
 370 375 380
 Lys Thr Leu His Leu Gly Asp Leu Ile Asn Arg Met His Ala Asn Pro
 385 390 395 400
 Tyr Arg Phe Thr Ala Ala Asn Val Lys Leu Pro Ala Lys Ala Glu
 405 410 415
 Arg Ile Glu Ile Asn Arg Tyr Gly Ala Val Ala Phe Arg Leu Tyr
 420 425 430

<210> 326
 <211> 1359
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 326
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 acgaattata tggagatcga tccggaatac gggaccaaag atgatttcag gagattgctg 120
 gacgaagccc acaagcgtgg cctcaagggtc atcatcgatt ttgtgatcaa ccacaccagc 180
 agccagcacc cctggttcct ggaggccagg aaaggcaaag acaatcctta ccgcaattat 240
 tacctgtgga tggagcccg cgaatcgaa aggatgggca tcgcggtgcg ggaaaaaacc 300
 gccgactcct gggaaacgca gccctggcac tgggccagcc gcggcgacaa ggaaaagtat 360
 tacgggatgt tctggaacgg gatgcctgat cttaatatgg acaaccgtga ggtgagagaa 420
 cagatttatg ccatcggcaa atactggctc gaattcgggg ctgacggctt caggatggat 480
 gctgccaaac acatttatcc cagttgggaa gccgaaaagt cgcacgactt ctgggtggag 540
 ttcagagagc ggatggaggc tgtcaagcct ggcttttacc tgggtgggga ggtgtggaca 600
 tcggcagaaa aggtagcccc gtttttccgg ggactgaaag ccaacttcca ttttgacctg 660
 agcctggccc tgcagaaaat agcggcggga caggacagcg gatcaggact ggtgaacatg 720
 ctgctgggca attacgagat attcggaaag gagaaccgg cgtttattga cgccacgatg 780
 ctgaccaacc acgaccagga gaggctggga agtgcggtag gcaacgataa aaataagatg 840
 aagctggctg ctaacctgct gctgaccctg cccggtaatc cctatcttta ttacggggaa 900
 gaactgggga tgctcgcaa aaagcctgat gaaaacatca gggaggcggt tctgtgggat 960
 acgcgggcga atgaccgcga ccggaccaac tggagaacaa ccaattttta taccgatcc 1020
 cgggtgacac ccctgaagtt gcaacagcag gacccggact cagtttatca tcattacaag 1080
 aaactgatcc gattaagaaa ggatcacccg gcattgtggc aggtatcccc gccgaatctt 1140
 caggcggcac aaacggctca ggaaggtttc gtgacgttca ttcgtccgca cgaagcggg 1200
 agtttgctgg tcatacagaa cctgcagaac aacaccgcc tgggtggaagt aaaagaggac 1260
 atcactgaga cggatttcag cactgcaggt tttcggaata ccgaaggga ggtcatggaa 1320
 ataccggggt ttgggatggt ggttttttagg ttggggtag 1359

<210> 327
 <211> 452
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 327
 Met Glu Gly Leu Trp Leu Thr Pro Phe Phe Gln Ser Pro Ser Tyr His
 1 5 10 15
 Lys Tyr Asp Val Thr Asn Tyr Met Glu Ile Asp Pro Glu Tyr Gly Thr
 20 25 30
 Lys Asp Asp Phe Arg Arg Leu Leu Asp Glu Ala His Lys Arg Gly Leu
 35 40 45
 Lys Val Ile Ile Asp Phe Val Ile Asn His Thr Ser Ser Gln His Pro
 50 55 60
 Trp Phe Leu Glu Ala Arg Lys Gly Lys Asp Asn Pro Tyr Arg Asn Tyr
 65 70 75 80
 Tyr Leu Trp Met Glu Pro Ala Glu Ile Glu Arg Met Gly Ile Ala Val
 85 90 95
 Arg Glu Lys Thr Ala Asp Ser Trp Glu Thr Gln Pro Trp His Trp Ala

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Ser	Arg	Gly	100	Asp	Lys	Glu	Lys	Tyr	105	Tyr	Gly	Met	Phe	Trp	110	Asn	Gly	Met
Pro	Asp	Leu	115	Asn	Met	Asp	Asn	Arg	120	Glu	Val	Arg	Glu	Gln	125	Ile	Tyr	Ala
Ile	Gly	Lys	130	Tyr	Trp	Leu	Glu	Phe	135	Gly	Ala	Asp	Gly	Phe	140	Arg	Met	Asp
Ala	Ala	Lys	145	His	Ile	Tyr	Pro	Ser	150	Trp	Glu	Ala	Glu	Lys	155	Ser	His	Asp
Phe	Trp	Val	160	Glu	Phe	Arg	Glu	Arg	165	Met	Glu	Ala	Val	Lys	170	Pro	Gly	Phe
Tyr	Leu	Val	180	Gly	Glu	Val	Trp	Thr	185	Ser	Ala	Glu	Lys	Val	190	Ala	Pro	Phe
Phe	Arg	Gly	195	Leu	Lys	Ala	Asn	Phe	200	His	Phe	Asp	Leu	Ser	205	Leu	Ala	Leu
Gln	Lys	Ile	210	Ala	Ala	Gly	Gln	Asp	215	Ser	Gly	Ser	Gly	Leu	220	Val	Asn	Met
Leu	Leu	Gly	225	Asn	Tyr	Glu	Ile	Phe	230	Gly	Arg	Glu	Asn	Pro	235	Ala	Phe	Ile
Asp	Ala	Thr	240	Met	Leu	Thr	Asn	His	245	Asp	Gln	Glu	Arg	Leu	250	Gly	Ser	Ala
Val	Gly	Asn	255	Asp	Lys	Asn	Lys	Met	260	Lys	Leu	Ala	Ala	Asn	265	Leu	Leu	Leu
Thr	Leu	Pro	270	Gly	Asn	Pro	Tyr	Leu	275	Tyr	Tyr	Gly	Glu	Glu	280	Leu	Gly	Met
Leu	Gly	Lys	285	Lys	Pro	Asp	Glu	Asn	290	Ile	Arg	Glu	Ala	Phe	295	Leu	Trp	Asp
Thr	Arg	Ala	300	Asn	Asp	Arg	Asp	Arg	305	Asn	Trp	Lys	Lys	Pro	310	Asn	Phe	
Asn	Thr	Asp	315	Ser	Arg	Val	Thr	Pro	320	Leu	Lys	Leu	Gln	Gln	325	Gln	Asp	Pro
Asp	Ser	Val	330	Tyr	His	His	Tyr	Lys	335	Lys	Leu	Ile	Arg	Leu	340	Arg	Lys	Asp
His	Pro	Ala	345	Leu	Trp	Gln	Val	Ser	350	Pro	Pro	Asn	Leu	Gln	355	Ala	Ala	Gln
Thr	Ala	Gln	360	Glu	Gly	Phe	Val	Thr	365	Phe	Ile	Arg	Pro	His	370	Glu	Ser	Gly
Ser	Leu	Leu	375	Val	Ile	Gln	Asn	Leu	380	Asn	Asn	Thr	Ala	Leu	385	Val	Glu	
Val	Lys	Glu	390	Asp	Ile	Thr	Glu	Thr	395	Val	Phe	Ser	Thr	Ala	400	Gly	Phe	Arg
Asn	Thr	Glu	405	Gly	Lys	Val	Met	Glu	410	Ile	Pro	Gly	Phe	Gly	415	Met	Val	Val
Phe	Arg	Leu	420	Gly					425						430			
			435						440						445			
			450															

<210> 328

<211> 1677

<212> DNA

<213> Unknown

<220>

<223> obtained from an environmental sample

<400> 328

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cgcgccgagc	tcgccagcat	ggtgagccgg	tgcgccgccg	cgggcggtga	gatctacgcc	180
gacgcggtga	tcaaccacat	gacgggcggc	tcgggcacga	cgtcgagcgc	cggaaccggg	240
ccgtggggcg	tgaagagcta	cccgaacgtc	ccgtacggcg	tgaacgactt	ccacgcgacg	300
tgcgcgatca	ccagctacgg	tgacgcaa	caggtccaga	actgcgagct	cgccgggctc	360
caggacctga	acaccggctc	ctcctacgtg	cgcggaaga	tcgtcgacta	cctgggtcgat	420
ctctacaacc	tcggcgctcc	cggtctccgg	atcgacgcgg	ccaagcacat	cagcccgacc	480
gacgtgacgg	cgatcgctgc	cgccgtcgag	gcccgcgtcc	cgacgaagcc	gttctggttc	540
ctcgagggtga	tcggcgcttc	cggcgagggc	gtgcagccga	atcagtactt	caatgtcgcc	600
gaggggcgcg	cgacggtgac	ggagttcgcg	tacggccgcg	agctgtacgg	caagttcgcg	660
ggcggcgggc	ggctggccga	actccgcacg	ttcggcgaga	cgtggaacct	cttgccgggc	720
gaccgcgcga	tcgcgttcac	cgacaaccac	gacaagcagc	gcggccacgg	cggcggcggc	780

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acgtacctca	cgtatcacaa	cggggcgacc	tacgacctcg	gcaacgtctt	catgctggcg	840
tggccgtacg	gctaccgggc	gctcatgtcg	agctacgcgt	tcaacaagac	caccgattac	900
gacacgagct	acggggccgc	gcacgactcg	gcgaccggcg	cgacgcgcgg	cccgtgggac	960
ggcggcgctct	ccgccccgc	gtgcttcaac	cagaccgcgc	gcggctgggt	ctgcgagcac	1020
cgcttccgcc	cgatcggcaa	catggtggcg	ttccgcaagg	cgacgatggg	caactggacc	1080
gtcaccgact	ggtgggacaa	cggcaataac	cagatcgcgt	ttggccgcgg	cggctcggga	1140
ttcgtgggtca	tcaacaagga	ggcggccgcg	ctgactcgca	cgttcaagac	gagcctcgcc	1200
gccggcaagt	actgcgacgt	catctcgggc	gactggaacg	cgggcgcggg	cacctgcgcc	1260
ggcaacgtcg	tgaccgtcga	cgcgaccggc	aacgcgagca	tcaccgtcgg	ggcggttcggc	1320
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tacaagtaca	tcaagaagga	cggcgcgggc	accgtcatct	gggagtcggg	cgccaaccgc	1620
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<210> 329
 <211> 558
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 329

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Gly	Ala	Pro	Tyr	Pro	Trp	Trp	Met	Arg	Tyr	Gln	Pro	Val	Ser	Tyr	Ser
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Leu	Asp	Arg	Ser	Arg	Ser	Gly	Thr	Arg	Ala	Glu	Leu	Ala	Ser	Met	Val
		35					40					45			
Ser	Arg	Cys	Ala	Ala	Ala	Gly	Val	Gln	Ile	Tyr	Ala	Asp	Ala	Val	Ile
	50					55					60				
Asn	His	Met	Thr	Gly	Gly	Ser	Gly	Thr	Thr	Ser	Ser	Ala	Gly	Asn	Arg
65				70					75					80	
Pro	Trp	Gly	Val	Lys	Ser	Tyr	Pro	Asn	Val	Pro	Tyr	Gly	Val	Asn	Asp
				85					90					95	
Phe	His	Ala	Thr	Cys	Ala	Ile	Thr	Ser	Tyr	Gly	Asp	Ala	Asn	Gln	Val
			100					105					110		
Gln	Asn	Cys	Glu	Leu	Ala	Gly	Leu	Gln	Asp	Leu	Asn	Thr	Gly	Ser	Ser
		115					120					125			
Tyr	Val	Arg	Gly	Lys	Ile	Val	Asp	Tyr	Leu	Val	Asp	Leu	Tyr	Asn	Leu
	130					135					140				
Gly	Val	Arg	Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His	Ile	Ser	Pro	Thr
145					150					155					160
Asp	Val	Thr	Ala	Ile	Val	Ala	Ala	Val	Glu	Ala	Arg	Val	Pro	Thr	Lys
				165					170					175	
Pro	Phe	Trp	Phe	Leu	Glu	Val	Ile	Gly	Ala	Phe	Gly	Glu	Ala	Val	Gln
			180					185					190		
Pro	Asn	Gln	Tyr	Phe	Asn	Val	Ala	Glu	Gly	Arg	Ala	Thr	Val	Thr	Glu
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Phe	Ala	Tyr	Gly	Arg	Glu	Leu	Tyr	Gly	Lys	Phe	Ala	Gly	Gly	Gly	Arg
	210					215					220				
Leu	Ala	Glu	Leu	Arg	Thr	Phe	Gly	Glu	Thr	Trp	Asn	Leu	Leu	Pro	Gly
225					230					235					240
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Gly	Gly	Val	Ser	Ala	Pro	Ala	Cys	Phe	Asn	Gln	Thr	Arg	Gly	Gly	Trp
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Lys	Ala	Thr	340	Met	Gly	Asn	Trp	Thr	345	Val	Thr	Asp	Trp	Trp	350	Asp	Asn	Gly
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Asn	Lys	Glu	370	Ala	Ala	Ala	Leu	Thr	375	Arg	Thr	Phe	Lys	Thr	Ser	Leu	Ala	
Ala	Gly	Lys	385	Tyr	Cys	Asp	Val	Ile	390	Ser	Gly	Asp	Trp	Asn	Ala	Gly	Ala	
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Ser	Ile	Thr	420	Val	Gly	Ala	Phe	Gly	425	Ala	Ala	Ala	Ile	His	Ala	Gly	Ala	
Lys	Leu	Gly	435	Gly	Gly	Gly	Gly	Gly	440	Gly	Thr	Ala	Ser	Val	Thr	Phe	Asn	
Glu	Ala	Ala	450	Asp	Thr	Val	Phe	Gly	455	Gln	Asn	Ile	Tyr	Val	Val	Gly	Ser	
Val	Pro	Ala	465	Leu	Ala	Ser	Trp	Asn	470	Pro	Ala	Ser	Ala	Val	Pro	Met	Thr	
Trp	Ile	Ser	485	Gly	Ser	Gly	Thr	Arg	490	Gly	Asn	Trp	Arg	Ala	Thr	Val	Thr	
Leu	Pro	Ala	500	Ser	Thr	Ala	Ile	Glu	505	Tyr	Lys	Tyr	Ile	Lys	Lys	Asp	Gly	
Ala	Gly	Thr	515	Val	Ile	Trp	Glu	Ser	520	Gly	Ala	Asn	Arg	Thr	Leu	Thr	Thr	
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<210> 330
 <211> 3129
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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caggccgagg	ccctgggtccc ttacttcgac gagctgggta ttagcgattg ttacgcctcg 180
ccccgtgtgg	cggcctgtcc tggcagcgag cacggctatg atgtctgtga tcccagccgg 240
ctcaatccgg	ccctgggcga tgagcaagcg tttgagtctt ttacctcggc cctccacgcc 300
cacaatatgg	gcctcatttt agatacgggt cccaaccata tgggcattag cctgcctgag 360
aatggctggg	ggttggatgt cctggaaaat ggcccggcgt caccctacgc cgcctatttt 420
gatattgact	ggcatccaat caagtccgaa ctgaacaata aagtcctgct gcctattttg 480
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caagagtttc	gcagtatcct cacggccttg gacgacttgc cgccccgcac cgccgcgcag 720
acagagcccc	tggtcgaacg gagccgggaa aaagaggtgg tcaaacgccg gctggccaat 780
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<210> 331
 <211> 1042
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<400> 331

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Phe Asn Ala Asn Phe Thr Leu Ala Gln Ala Glu Ala Leu Val Pro Tyr
 35           40           45
Phe Asp Glu Leu Gly Ile Ser Asp Cys Tyr Ala Ser Pro Leu Leu Ala
 50           55           60
Ala Cys Pro Gly Ser Glu His Gly Tyr Asp Val Cys Asp Pro Ser Arg
 65           70           75           80
Leu Asn Pro Ala Leu Gly Asp Glu Gln Ala Phe Glu Ser Phe Thr Ser
 85           90           95
Ala Leu His Ala His Asn Met Gly Leu Ile Leu Asp Thr Val Pro Asn
100           105           110
His Met Gly Ile Ser Leu Pro Glu Asn Gly Trp Trp Leu Asp Val Leu
115           120           125
Glu Asn Gly Pro Ala Ser Pro Tyr Ala Ala Tyr Phe Asp Ile Asp Trp
130           135           140
His Pro Ile Lys Ser Glu Leu Asn Asn Lys Val Leu Leu Pro Ile Leu
145           150           155           160
Glu Asn Gln Tyr Gly Arg Val Leu Glu Ser Gly Lys Leu Glu Leu Val
165           170           175
Tyr Glu Asn Gly Ala Phe Phe Ile Tyr Tyr Tyr Asp Thr Lys Leu Pro
180           185           190
Val Ser Pro Arg Ser Tyr Arg Leu Ile Leu Thr Glu Cys Leu Ala Arg
195           200           205
Leu Asp Glu Glu Leu Ala Glu Ala Asp Glu Asp Leu Gln Glu Phe Arg
210           215           220
Ser Ile Leu Thr Ala Leu Glu His Leu Pro Pro Arg Thr Ala Ala Glu
225           230           235           240
Thr Glu Ala Leu Val Glu Arg Ser Arg Glu Lys Glu Val Val Lys Arg
245           250           255
Arg Leu Ala Asn Leu Thr Gln Ser Asn Ala Thr Val Gln Ala Ala Ile
260           265           270
Glu Ala Ala Val Thr Thr Leu Asn Gly Arg Pro Gly Glu Pro Glu Ser
275           280           285

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305					310					315					320
Ile	Asn	Ser	Leu	Ala	Ala	Ile	Arg	Val	Glu	Leu	Pro	Gln	Val	Phe	Ala
				325					330					335	
Asp	Thr	His	Ala	Leu	Ile	Phe	Arg	Trp	Leu	Ala	Glu	Gly	Lys	Val	Thr
			340					345					350		
Gly	Leu	Arg	Ile	Asp	His	Pro	Asp	Gly	Leu	Trp	Ser	Pro	Gly	Gly	Tyr
		355					360					365			
Phe	Lys	Gln	Leu	His	Glu	Asn	Tyr	Leu	Leu	Asp	Gln	Leu	Ala	Pro	Ala
	370					375					380				
Ala	Thr	Gly	Val	Glu	Gln	Glu	Glu	Leu	His	Gln	Ala	Ile	Val	Ala	Trp
385					390					395					400
Leu	Glu	Thr	Val	Glu	Ala	Ser	Trp	Thr	Glu	Ala	Asp	Gln	Ala	Glu	Pro
				405					410					415	
Ala	Glu	Ala	Ala	Leu	Pro	Trp	Pro	Leu	Tyr	Ile	Val	Ala	Glu	Lys	Ile
			420					425					430		
Leu	Ser	Glu	Thr	Glu	Pro	Leu	Pro	Pro	Asp	Trp	Ala	Val	Asp	Gly	Thr
		435					440					445			
Thr	Gly	Tyr	Asp	Phe	Leu	Asn	Gln	Val	Asn	Gly	Val	Phe	Ile	Asn	Arg
	450					455					460				
Gln	Ser	Glu	Arg	Ala	Phe	Asp	Arg	Ile	Tyr	Arg	Arg	Phe	Thr	Gly	Ile
465					470					475					480
Ser	Ala	Asn	Phe	Ala	Glu	Leu	Ala	Ile	Gln	Thr	Lys	Asp	Met	Ile	Met
				485					490					495	
Arg	Leu	Ala	Leu	Ala	Ser	Glu	Val	Asn	Leu	Leu	Ala	His	Gln	Leu	Glu
			500					505					510		
Gln	Ile	Asn	Glu	Lys	Asn	Arg	Arg	Tyr	Arg	Asp	Phe	Thr	Leu	Ser	Gly
		515					520					525			
Val	Thr	Ala	Ala	Leu	Arg	Glu	Val	Leu	Ala	Cys	Leu	Pro	Ile	Tyr	Arg
	530					535					540				
Thr	Tyr	Ile	Thr	Pro	Pro	His	Pro	Val	Thr	Ala	Arg	Asp	Glu	Ser	Tyr
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Ile	Glu	Ala	Ala	Val	Ala	Glu	Ala	Arg	Arg	Arg	Asn	Pro	Arg	Leu	Ser
				565					570					575	
Thr	Ser	Leu	Leu	Asn	Phe	Leu	Arg	Asp	Thr	Leu	Leu	Leu	Arg	Asn	Leu
			580					585					590		
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		595					600					605			
Lys	Phe	Gln	Gln	Val	Ser	Gly	Pro	Leu	Thr	Ala	Lys	Gly	Leu	Glu	Asp
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Thr	Ala	Phe	Tyr	Val	Tyr	Asn	Arg	Phe	Val	Ser	Leu	Asn	Glu	Val	Gly
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Gly	His	Pro	Gly	Gln	Phe	Gly	Leu	Arg	Val	Glu	Asp	Phe	His	Gln	Gln
				645					650					655	
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Thr	His	Asp	Thr	Lys	Arg	Ser	Glu	Asp	Val	Arg	Ala	Arg	Leu	Asn	Val
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Phe	Ser	Glu	Met	Pro	Glu	Glu	Trp	Arg	Met	Ala	Leu	Ser	Arg	Trp	Arg
	690					695					700				
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Asp	Arg	Asn	Asp	Glu	Tyr	Leu	Phe	Tyr	Gln	Thr	Val	Val	Gly	Ala	Trp
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Pro	Phe	Ala	Val	Asn	Glu	Gly	Met	Asp	Ser	Ala	Ala	Gln	Ser	Gly	Ile
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Ala	Arg	Pro	Val	Val	Ser	Arg	Leu	Met	Gln	Pro	Gly	Arg	Gly	Glu	Leu
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	770					775					780				
Glu	Ala	Lys	Val	His	Thr	Ser	Trp	Ile	Asn	Pro	Asn	Glu	Ala	Tyr	Asp
785					790					795					800
Lys	Ala	Leu	Glu	Lys	Phe	Val	Cys	Asp	Ser	Leu	Asp	Asn	Gln	Arg	Ala
				805					810					815	
Asn	Arg	Phe	Leu	Gly	Glu	Leu	Leu	Ala	Phe	Thr	Glu	Arg	Val	Ala	Tyr
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Leu Val Asp Pro Asp Asn Arg Arg Pro Val Asp Tyr Asp Gln Arg Gln
865 870 875 880
Arg Leu Leu Ala Glu Leu Lys Gln Ser Ile Ala Gly Gly Gln Pro Glu
885 890 895
Gln Gly Gln Ile Leu Ala Gln Asn Leu Leu Glu Thr Ser His Asp Gly
900 905 910
Arg Ile Lys Phe Tyr Ile Thr Ala Arg Thr Leu Asn Phe Arg Arg Glu
915 920 925
Gln Pro Glu Leu Phe Asp Gln Gly Asp Tyr Gln Pro Leu Ala Ala Ser
930 935 940
Gly Asp Gln Ser Asp His Ile Ile Ala Phe Ser Arg Ser Trp Gly Asp
945 950 955 960
Glu Asn Phe Ile Val Val Val Pro Arg Leu Val Phe Ser Leu Met Ala
965 970 975
Gly Val Glu Arg Pro Pro Leu Ala Asn Val Trp Gly Asp Thr Trp Leu
980 985 990
His Leu Pro Asp Pro Ala Ala Asn Arg Ser Tyr Arg His Leu Phe Thr
995 1000 1005
Gly Gln Thr Leu Ser Pro Gln Ser Ile Ser Asp Gly Pro Gly Leu Pro
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<210> 332
<211> 1707
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

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<210> 333
<211> 569
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(22)

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35 40 45
Lys Ala Gly Glu Leu Gly Ala Ala Gly Ile Asn His Val Trp Leu Pro
50 55 60
Pro Pro Ser Lys Ser Gly Ala Pro Gln Gly Tyr Leu Pro Thr Glu Leu
65 70 75 80
Asn Val Leu Asp Ser Ala Tyr Gly Thr Glu Leu Gln Leu Lys Asp Ala
85 90 95
Ile Ser Ala Leu Asn Ala Ala Gly Val Ser Ala Val Ala Asp Ile Val
100 105 110
Val Asn His Arg Asn Gly Ser Thr Ala Trp His Asp Phe Arg Asn Pro
115 120 125
Asp Trp Gly Thr Asp Ser Ile Val Ala Asn Asp Glu Cys Trp Ser Thr
130 135 140
Ala Gly Ser Thr Cys Thr Ser Ser Met Val Arg Gly Ala Asn Asp Ser
145 150 155 160
Gly Glu Pro Tyr Ala Ala Ala Arg Asp Val Asp His Gly Lys Gln Tyr
165 170 175
Val Arg Asp Ser Leu Val Gly Trp Met Gly Thr Arg Leu Ala Asn Val
180 185 190
Gly Phe Asp Gly Trp Arg Phe Asp Phe Val Lys Gly Phe Ser Gly Ala
195 200 205
Tyr Val Gly Glu Tyr Val Ala Arg Thr Ala Pro Thr Phe Cys Val Gly
210 215 220
Glu Phe Trp Pro Thr Asn Tyr Phe Asp Leu Asn Asp Pro Ala Asn Trp
225 230 235 240
Arg Asn Gln Ile Ile Gly Trp Val Asp Ala Thr Gly Gly Arg Cys Ser
245 250 255
Ala Phe Asp Phe Val Thr Lys Gly Leu Leu Asn Gln Val Leu Ala Asn
260 265 270
Gly Asp Tyr Gly Arg Leu Lys Thr Ala Asp Gly Lys Pro Thr Gly Thr
275 280 285
Ile Gly Val Arg Pro Ala Arg Ser Val Thr Phe Val Asp Asn His Asp
290 295 300
Thr Gly Pro Ser Glu Ala Cys Gly Asp Gly Gln Asn His Trp Pro Val
305 310 315 320
Pro Cys Asp Lys Val Met Ala Gly Tyr Ala Tyr Ile Leu Thr His Pro
325 330 335
Gly Val Pro Thr Val Tyr Trp Thr His Tyr Tyr Asn Phe Gly Leu Lys
340 345 350
Ala Gln Leu Asp Lys Leu Ile Ser Ile Arg Lys Thr Asn Gly Leu Gln
355 360 365
Ser Glu Ser Val Val Asn Ile Val Arg Ala Glu Gln Asn Leu Tyr Ala
370 375 380
Ala Ile Ile Asp Ala Lys Val Ala Met Lys Ile Gly Ser Gly Ser Trp
385 390 395 400
Ser Pro Gly Thr Asp Trp Thr Leu Ala Ala Ser Gly Thr Asp Tyr Ala
405 410 415
Val Trp Thr Lys Gly Thr Thr Pro Pro Pro Thr Gly Cys Thr Gln
420 425 430
Pro Ser Met Asn Leu Arg Gly Thr Phe Asn Asn Phe Gly Ser Ala Ala
435 440 445

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Met Thr Cys Lys Gly Gly Asn Thr Trp Glu Ile Thr Asn Val Thr Phe
  450 455 460
Gly Gly Ala Ala Thr Asp Arg Phe Lys Phe Asp Val Phe Gly Asp Trp
  465 470 475 480
Thr Arg Asn Tyr Gly Asp Thr Asn Asn Asp Gly Thr Ala Glu Leu Ser
  485 490 495
Gly Gly Asp Ile Ser Leu Thr Thr Ala Gly Thr Tyr Thr Leu Gln Phe
  500 505 510
Asn Asp Ser Thr Leu Arg Tyr Thr Val Thr Arg Thr Ser Thr Pro Pro
  515 520 525
Pro Pro Thr Gly Thr Ser Val Arg Phe Val Cys Gln Asn Gly Thr Thr
  530 535 540
Phe Val Gly Gln Asn Val His Val Val Gly Ser Ile Pro Glu Leu Gly
  545 550 555 560
Ser Trp Asp Ala Thr Lys Ser Ile Lys
  565

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<210> 334
 <211> 2061
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

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<400> 334
atggtagcag gctttgggct ctacggagct gccttgctca ccccaatggc ggcacaggcc 60
gaaacgggca gagaagtcac gctgcagggc tttaactgga attccacctc gcaagcgggt 120
gggcactata atgagctggc aaatcgtgct ggtgaaatcg ctggggctgg tatcgacatt 180
gtttggttcc cacctccatc cagagccgca gaccgggttg gatacctgcc caacgagtgg 240
tataatatga actccaatta tggaaatcgg acgactctac aggcggctat cgggaatcct 300
cgcaacaacg gtgtcaaaac cgtcgcggat attgtcggtt accaccgtgt gggcacaacg 360
aattgggicgg acttcacgaa tccttctttc ggtgataaca accgtgccat cacacgcgat 420
gatgaatggc atcagtcgtc gggttaactgg gataccgggtg aggcgtatag tgccgcccg 480
gacttgacc acacctacgg ccccgtgcaa aatgagatta aaaactggct gaattgggtg 540
aagagtgcac tcggctttga tggttggcgc tatgatattg tcaagggatt cagcgggtac 600
tatgtgggcg agtacaacac agcaactagc ccctacattt cgttgggtga gttcttcgac 660
tatgatcgcc agaaggtggg cggctggatc aacgccacca atgctcgctc tcgcgcttc 720
gattttccaa cagcaacct cctctatgtg gctgtcacc aaacaacta cggagtctt 780
cgtgatggg aaggttaagg caatggactc atcggttggg ggccacaacg ggctatcaca 840
ttcattgaga accatgatac ggaagaagcc cgcaacgggt agtacacccc cgccttcccc 900
caatgggcta ccatgcaggg ttatgcctac attttgacgc acccggcat tccatgcgtt 960
ttctggaatg attggcgttg ggacttccgt tcggaaatca atcaactgat cgccatccgc 1020
agggcccaag gcatcaacga tggcagcagc ctacagcatc aggttgccga tggcagccgc 1080
tcggggcgca tcatcaatgg aaacacggcg gtcaagattg gtcctggaaa ctggagccca 1140
agtggtagtt ggacgctggc tgccgccgga accaattacg ccgtctggac aaacgggtggc 1200
ggaacaccaa cgccaacacc cacaccaacc ccaactcagg gccccacaac ggttacttgg 1260
aatccatcca ctccgacagc cggacaaaat gtgacgatca cttatccttc cggccgctcc 1320
cttgccagtt cctccaacgt caacctctac tggggtgtca atggatggac caatgtccaa 1380
accaaagcga tgaccaagaa cagttccaat gactggacaa ccacgatcac acttccctcc 1440
aacacaacac gtttgaactt tgtgttcaat aacggctcca gttgggacaa caacagcagt 1500
caggattgga atgtcaacgt gaccgctgta acgcccacac caacaccgac accgactccc 1560
acacctacgg caactccgac accgacaccc accccgacgc ccactgccac acccacccca 1620
actccgactc cttcgatcat ttggtatcgc atcgaagccc gtcacagtgg taaagttcta 1680
gacgttgcca gtgcatcaac ttcaaacggg ggcaatgtgc atcagtgggt ttatgctgg 1740
ggacagaaac aacaatggcg tgtgtcgcac gctgggaacg ggtttgttta catcctaaac 1800
cggaacagtg gaaaagcact tgaagtgggc aacttctcca ccagtaacgg cgggaacgtc 1860
cagcaatggg attatgccgg tggttccagt caacagtgga agctcattga gacgaccaac 1920
ggatatgtgc aaatccagaa tcggaacagc ggcaaagcca ttgatgtttc cgctgcttcg 1980
actaccaatg gtgccaacat ccatcagtgg acctacggcg gcggcaacaa tcagcagtgg 2040
aagctgatcc caatcaatta a
  2061

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<210> 335
 <211> 686
 <212> PRT
 <213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(20)

<400> 335

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Met Val Ala Gly Phe Gly Leu Tyr Gly Ala Ala Leu Leu Thr Pro Met
1 10 15
Ala Ala Gln Ala Glu Thr Gly Arg Glu Val Met Leu Gln Gly Phe Asn
20 25 30
Trp Asn Ser Thr Ser Gln Ala Gly Gly His Tyr Asn Glu Leu Ala Asn
35 40 45
Arg Ala Gly Glu Ile Ala Gly Ala Gly Ile Asp Ile Val Trp Phe Pro
50 55 60
Pro Pro Ser Arg Ala Ala Asp Arg Val Gly Tyr Leu Pro Asn Glu Trp
65 70 75 80
Tyr Asn Met Asn Ser Asn Tyr Gly Asn Arg Thr Thr Leu Gln Ala Ala
85 90 95
Ile Gly Asn Leu Arg Asn Asn Gly Val Lys Thr Val Ala Asp Ile Val
100 105 110
Val Asn His Arg Val Gly Thr Thr Asn Trp Ala Asp Phe Thr Asn Pro
115 120 125
Ser Phe Gly Asp Asn Asn Arg Ala Ile Thr Arg Asp Asp Glu Trp His
130 135 140
Gln Ser Ser Gly Asn Trp Asp Thr Gly Glu Ala Tyr Ser Ala Ala Arg
145 150 155 160
Asp Leu Asp His Thr Tyr Gly Pro Val Gln Asn Glu Ile Lys Asn Trp
165 170 175
Leu Asn Trp Leu Lys Ser Asp Ile Gly Phe Asp Gly Trp Arg Tyr Asp
180 185 190
Met Val Lys Gly Phe Ser Gly Tyr Tyr Val Gly Glu Tyr Asn Thr Ala
195 200 205
Thr Ser Pro Tyr Ile Ser Val Gly Glu Phe Phe Asp Tyr Asp Arg Gln
210 215 220
Lys Val Val Gly Trp Ile Asn Ala Thr Asn Ala Arg Ser Arg Ala Phe
225 230 235 240
Asp Phe Pro Thr Arg Asn Leu Leu Tyr Val Ala Val Thr Gln Asn Asn
245 250 255
Tyr Gly Val Leu Arg Asp Gly Glu Gly Lys Ala Asn Gly Leu Ile Gly
260 265 270
Trp Trp Pro Gln Arg Ala Ile Thr Phe Ile Glu Asn His Asp Thr Glu
275 280 285
Glu Ala Arg Asn Gly Glu Tyr Thr Pro Ala Phe Pro Gln Trp Ala Thr
290 295 300
Met Gln Gly Tyr Ala Tyr Ile Leu Thr His Pro Gly Ile Pro Cys Val
305 310 315 320
Phe Trp Asn Asp Trp Arg Trp Asp Phe Arg Ser Glu Ile Asn Gln Leu
325 330 335
Ile Ala Ile Arg Arg Ala Gln Gly Ile Asn Asp Gly Ser Ser Leu Ser
340 345 350
Ile Gln Val Ala Asp Gly Ser Arg Tyr Gly Ala Ile Ile Asn Gly Asn
355 360 365
Thr Ala Val Lys Ile Gly Pro Gly Asn Trp Ser Pro Ser Gly Ser Trp
370 375 380
Thr Leu Ala Ala Ala Gly Thr Asn Tyr Ala Val Trp Thr Asn Gly Gly
385 390 395 400
Gly Thr Pro Thr Pro Thr Pro Thr Pro Thr Pro Thr Gln Gly Pro Thr
405 410 415
Thr Val Thr Trp Asn Pro Ser Thr Pro Thr Ala Gly Gln Asn Val Thr
420 425 430
Ile Thr Tyr Pro Ser Gly Arg Ser Leu Ala Ser Ser Ser Asn Val Asn
435 440 445
Leu Tyr Trp Gly Val Asn Gly Trp Thr Asn Val Gln Thr Lys Ala Met
450 455 460
Thr Lys Asn Ser Ser Asn Asp Trp Thr Thr Thr Ile Thr Leu Pro Ser
465 470 475 480
Asn Thr Thr Arg Leu Asn Phe Val Phe Asn Asn Gly Ser Ser Trp Asp
485 490 495

```


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Asn	Asn	Ser	Ser	Gln	Asp	Trp	Asn	Val	Asn	Val	Thr	Ala	Val	Thr	Pro
Thr	Pro	Thr	Pro	Thr	Pro	Thr	Pro	Thr	Pro	Thr	Ala	Thr	Pro	Thr	Pro
Thr	Pro	Thr	Pro	Thr	Pro	Thr	Ala	Thr	Pro	Thr	Pro	Thr	Pro	Thr	Pro
Ser	Ile	Ile	Trp	Tyr	Arg	Ile	Glu	Ala	Arg	His	Ser	Gly	Lys	Val	Leu
Asp	Val	Ala	Ser	Ala	Ser	Thr	Ser	Asn	Gly	Gly	Asn	Val	His	Gln	Trp
Ser	Tyr	Ala	Gly	Gly	Gln	Asn	Gln	Gln	Trp	Arg	Val	Val	Asp	Ala	Gly
Asn	Gly	Phe	Val	Tyr	Ile	Leu	Asn	Arg	Asn	Ser	Gly	Lys	Ala	Leu	Glu
Val	Gly	Asn	Phe	Ser	Thr	Ser	Asn	Gly	Gly	Asn	Val	Gln	Gln	Trp	Asp
Tyr	Ala	Gly	Gly	Ser	Ser	Gln	Gln	Trp	Lys	Leu	Ile	Glu	Thr	Thr	Asn
Gly	Tyr	Val	Gln	Ile	Gln	Asn	Arg	Asn	Ser	Gly	Lys	Ala	Ile	Asp	Val
Ser	Ala	Ala	Ser	Thr	Thr	Asn	Gly	Ala	Asn	Ile	His	Gln	Trp	Thr	Tyr
Gly	Gly	Gly	Asn	Asn	Gln	Gln	Trp	Lys	Leu	Ile	Pro	Ile	Asn		

<210> 336
 <211> 1731
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 336

atgaccgaac	gacacctcag	tcgcagggcg	aagggggcg	tggcgggcct	cgccgctgcc	60
gccgtcgccg	ccgtcaccct	cgcggtcccc	cccgcagcgc	aggcatcgcc	ccccgggaac	120
aaggacgtca	ccgctgtcct	gttctcgtgg	aacttcgact	ccatagcccg	ggagtgcacc	180
gaccgcttgg	gccccgccgg	ctacggcttc	gtccagggtc	caccgcccc	ggagcacatc	240
cagggctccg	cgtggtggac	ccagtaccag	cccgtcagct	accggctgga	gagcaggctc	300
ggcaaccgcg	cccagttccg	gaacatggtc	gacacctgca	acagcgccgg	tgtgggcgtg	360
gtcgtcgaca	ccgtcatcaa	ccacatgacc	gcgggggtccg	gcaccggcac	cggcggttcg	420
cagtacagca	agtaccacta	ccccgacgcc	ccttacagcg	acggggactt	caacgactgc	480
cgccgcgaca	tcagcaacta	ccgcgaccgg	tacgacgtcc	agaactgcga	actggtcggc	540
ctcgcggaac	tcgggaccgg	caaggagtac	gtccgcccag	gcacgcgcga	ctacatgaac	600
gacctgctgt	cgctgggcgt	cgccggcttc	cgtatcgacg	ccgccaagca	catgcccgcc	660
gacgacctgc	gggccatcag	ggccaagctg	aacaaccgga	acgcctactg	gaagcaggaa	720
gtcatccatg	gcgcccgcga	ggccgtccag	cccaggaggt	acctcggcgc	cggcgacgtc	780
caggagtccc	gctacgcgca	cgacatgaag	cgcatgttcc	agcaggaacg	catcgcccac	840
ctgcggacct	tcggcgagag	ctggggcttc	atgccagctc	cgcagtcggc	ggtcttcgtc	900
gacaaccacg	acaccgagcg	caacggcacc	accctcagct	acaaggacgg	gtcggcctac	960
acgctcgcca	acgtcttcat	gctggcctgg	ccctacggct	ccccggacgt	ccactccggg	1020
taccggttca	gcaacttcga	cgcgggccct	cccggcaacg	gccaggtgag	ccagtgtctg	1080
cagaacggct	gggagtgcga	gcacgcctgg	cccagatcgc	agtccatggt	cgccttcctc	1140
aacgccaccc	gcggccagca	ggtcaccaac	tgggtgggaca	acggcaacaa	cgtcacgcgc	1200
ttcggccggg	gcagccaggg	ctacgtggcg	atcaaccggg	agagcggacc	ggtcaccgcg	1260
accttcacga	cctccctggc	cgccggtgac	tactgcgacg	tccagtccgg	gtccacggtc	1320
accgtgaacg	gctcggggca	gttcaccgcc	accgtggccg	gcggaaccgc	gctcgccttg	1380
cacaccggcg	cccggaaactg	caccggcaac	ggcggcggca	ccccccagga	gccggtggac	1440
ggcggcgtct	ccttcgccgt	caacgcctcc	acgcagtggg	gcgagcacct	ctatgtcacc	1500
ggcaaccgcc	ccgaactggg	caactgggac	ccgcagaacg	ccgtccggct	cggctcggac	1560
tactgacggg	tctggaagcg	cgaactggcg	atgcccgcgc	acacctcctt	ccagtacaag	1620
tacctgcggc	gcaccgacgg	cggaaaccgtg	acctgggagc	agggcggcaa	ccgcaccgcg	1680
accgtccccg	cctccgggac	cctgcggctg	aacgacacct	ggcgcagcta	g	1731

<210> 337
 <211> 576
 <212> PRT
 <213> Unknown

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<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(35)

<400> 337

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Met Thr Glu Arg His Leu Ser Arg Arg Ala Lys Gly Ala Leu Ala Gly
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Leu Ala Ala Ala Val Ala Ala Val Thr Leu Ala Ala Pro Pro Ala
      20
Ala Gln Ala Ser Pro Pro Gly Asn Lys Asp Val Thr Ala Val Leu Phe
      35
Ser Trp Asn Phe Asp Ser Ile Ala Arg Glu Cys Thr Asp Arg Leu Gly
 50      55
Pro Ala Gly Tyr Gly Phe Val Gln Val Ser Pro Pro Gln Glu His Ile
 65      70
Gln Gly Ser Ala Trp Thr Gln Tyr Gln Pro Val Ser Tyr Arg Leu
      85
Glu Ser Arg Leu Gly Asn Arg Ala Gln Phe Arg Asn Met Val Asp Thr
      100
Cys Asn Ser Ala Gly Val Gly Val Val Val Asp Thr Val Ile Asn His
      115
Met Thr Ala Gly Ser Gly Thr Gly Thr Gly Gly Ser Gln Tyr Ser Lys
 130      135
Tyr His Tyr Pro Asp Ala Pro Tyr Ser Asp Gly Asp Phe Asn Asp Cys
 145      150
Arg Arg Asp Ile Ser Asn Tyr Arg Asp Arg Tyr Asp Val Gln Asn Cys
      165
Glu Leu Val Gly Leu Ala Asp Leu Arg Thr Gly Lys Glu Tyr Val Arg
      180
Gln Arg Ile Ala Asp Tyr Met Asn Asp Leu Leu Ser Leu Gly Val Ala
      195
Gly Phe Arg Ile Asp Ala Ala Lys His Met Pro Ala Asp Asp Leu Arg
      210
Ala Ile Arg Ala Lys Leu Asn Asn Pro Asp Ala Tyr Trp Lys Gln Glu
 225      230
Val Ile His Gly Ala Gly Glu Ala Val Gln Pro Glu Glu Tyr Leu Gly
      245
Ala Gly Asp Val Gln Glu Phe Arg Tyr Ala His Asp Met Lys Arg Met
      260
Phe Gln Gln Glu Arg Ile Ala His Leu Arg Thr Phe Gly Glu Ser Trp
      275
Gly Phe Met Pro Ser Ser Gln Ser Ala Val Phe Val Asp Asn His Asp
      290
Thr Glu Arg Asn Gly Thr Thr Leu Ser Tyr Lys Asp Gly Ser Ala Tyr
 305      310
Thr Leu Ala Asn Val Phe Met Leu Ala Trp Pro Tyr Gly Ser Pro Asp
      325
Val His Ser Gly Tyr Arg Phe Ser Asn Phe Asp Ala Gly Pro Pro Gly
      340
Asn Gly Gln Val Ser Gln Cys Trp Gln Asn Gly Trp Glu Cys Gln His
      355
Ala Trp Pro Glu Ile Glu Ser Met Val Ala Phe Arg Asn Ala Thr Arg
      370
Gly Gln Gln Val Thr Asn Trp Trp Asp Asn Gly Asn Asn Val Ile Ala
 385      390
Phe Gly Arg Gly Ser Gln Gly Tyr Val Ala Ile Asn Arg Glu Ser Gly
      405
Pro Val Thr Arg Thr Phe Gln Thr Ser Leu Ala Ala Gly Asp Tyr Cys
      420
Asp Val Gln Ser Gly Ser Thr Val Thr Val Asn Gly Ser Gly Gln Phe
      435
Thr Ala Thr Val Ala Gly Gly Thr Ala Leu Ala Leu His Thr Gly Ala
      450
Arg Asn Cys Thr Gly Asn Gly Gly Gly Thr Pro Gln Glu Pro Val Asp
 465      470      480

```

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Gly Gly Val Ser Phe Ala Val Asn Ala Ser Thr Gln Trp Gly Glu His
485 490 495
Leu Tyr Val Thr Gly Asn Arg Pro Glu Leu Gly Asn Trp Asp Pro Gln
500 505 510
Asn Ala Val Arg Leu Gly Ser Asp Asp Tyr Pro Val Trp Lys Ala Glu
515 520 525
Leu Ala Met Pro Ala Asp Thr Ser Phe Gln Tyr Lys Tyr Leu Arg Arg
530 535 540
Thr Asp Gly Gly Thr Val Thr Trp Glu Gln Gly Asn Arg Thr Ala
545 550 555 560
Thr Val Pro Ala Ser Gly Thr Leu Arg Leu Asn Asp Thr Trp Arg Ser
565 570 575

<210> 338
<211> 1704
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 338
atgccgcacc aatacaacta caggccgaaa atacaccgat tatttttcagg tgctccaatg 60
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gccttctcac tatcctgtag cctgcaagcc aattcattgc cgcagatccc ggcagagcag 180
gtcaacaata ccatgtatca gatgttctac tgggatgcct atcccggact ttgggccgat 240
ctgccagcca tggcggaacc attggctgaa cgggggatca cctccatgtg gctgcccgcg 300
gcggcaaaag gcatgaatgg caccaacagc gttggctacg atgtctatga cttctgggat 360
ctgggcgaat tctacaaaa agggaccatc gccaccggtt acggcacccg ccagcagctg 420
gaccaggcgc tgagtgtctt ggatcaactg ggtgttcagg cctattttga tgtggtcttt 480
aaccaccgca tgggcgccga tgcccaggaa tacattcctg gctatggact ggcctggacc 540
gagtaccagc tgcagggtcg gcaggtgcat tataccagc aaaactgggg ctatttgttg 600
cagcactttg actggaactg gaccgcgttt aatggctccg acaatcagtt gtatcctggt 660
aaatggtggg gcaatacctt ccacttcctt tatttgatgg gcgaggatgt cgattacaac 720
cgctttgaag tgcagcagga aatgaaagcc tggggggagt ggatcatcaa cagcgttggc 780
tttagcggct ttcggatgga tgccatcgcc catgtcgata ccgattttac ccgtgactgg 840
atcaatcacg tgcagtgggc caccagttag gatgtgttct ttgtcgctga agcctgggtc 900
agcgatatca acggctatct ggatgccgtc aatacgccgc atttgcgcgc ttttgatttc 960
aatgtgcgcg aagactttgt tgctttaagc agtggcagca aagacatgcg ctggtggggg 1020
ggtctggtca atagccagca ccgtgatcgg gcggtcacct ttgtcgataa ccacgatacc 1080
agccgggccc gcaaccctta tggcatgccg caggtgatca actacaagaa ccaggcctac 1140
gcttacattc tgttgctgta gcatgggggt cgcaccgtgt ttgcccgcga ttacgacgaa 1200
tttggtcattg cgccaacgct ggataaactg attgaagcgc gccgttactt tgcctatggc 1260
cctggccatg agtactccgg caatactgaa gccgtctatg cctatgtgcg cgaggggcta 1320
agtactgtgc cgggcaccgg tttggtgatg ctgatgtcgg gtcgtaactg ggggtggctag 1380
cagtcgttca ccatcaacag ccaccagccg aataccacct tttacgatta caccggcaat 1440
gtcagtggca ccgtaaccac caatgcgcag ggctatggca gcttcccggg caatatgacg 1500
gaaagtaccg gttggtcagt ctgggtacca caatccattg gtggcactca gccgggaacc 1560
attaccctgc gtagtaccaa ggatgttggc tatggctttt cactgttctt caccggcagc 1620
agtgcggagc tgaccaattg gggcggcggg gttgaaggca cctggacatc cggtaattgtg 1680
tggaagtga ccattcccga tcca 1704

<210> 339
<211> 568
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(50)

<400> 339
Met Pro His Gln Tyr Asn Tyr Arg Pro Lys Ile His Arg Leu Phe Ser
1 5 10 15
Gly Ala Pro Met Gln Thr Asn Lys Lys Thr Phe Leu Ala His Cys Leu
20 25 30

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Arg	Gln	Ser	Leu	Val	Ala	Leu	Gly	Ala	Phe	Ser	Leu	Ser	Cys	Ser	Leu
Gln	Ala	Asn	Ser	Leu	Pro	Gln	Ile	Pro	Ala	Glu	Gln	Val	Asn	Asn	Thr
Met	Tyr	Gln	Met	Phe	Tyr	Trp	Asp	Ala	Tyr	Pro	Gly	Leu	Trp	Ala	Asp
Leu	Pro	Ala	Met	Ala	Glu	Pro	Leu	Ala	Glu	Arg	Gly	Ile	Thr	Ser	Met
Trp	Leu	Pro	Pro	Ala	Ala	Lys	Gly	Met	Asn	Gly	Thr	Asn	Ser	Val	Gly
Tyr	Asp	Val	Tyr	Asp	Phe	Trp	Asp	Leu	Gly	Glu	Phe	Tyr	Gln	Lys	Gly
Thr	Ile	Ala	Thr	Arg	Tyr	Gly	Thr	Arg	Gln	Gln	Leu	Asp	Gln	Ala	Leu
Ser	Ala	Leu	Asp	Gln	Leu	Gly	Val	Gln	Ala	Tyr	Phe	Asp	Val	Val	Phe
Asn	His	Arg	Met	Gly	Ala	Asp	Ala	Gln	Glu	Tyr	Ile	Pro	Gly	Tyr	Gly
Leu	Ala	Trp	Thr	Glu	Tyr	Gln	Leu	Gln	Gly	Arg	Gln	Val	His	Tyr	Thr
Gln	Gln	Asn	Trp	Gly	Tyr	Leu	Trp	His	Asp	Phe	Asp	Trp	Asn	Trp	Thr
Ala	Phe	Asn	Gly	Ser	Asp	Asn	Gln	Leu	Tyr	Pro	Gly	Lys	Trp	Trp	Gly
Asn	Thr	Phe	His	Phe	Pro	Tyr	Leu	Met	Gly	Glu	Asp	Val	Asp	Tyr	Asn
Arg	Phe	Glu	Val	Gln	Gln	Glu	Met	Lys	Ala	Trp	Gly	Glu	Trp	Ile	Ile
Asn	Ser	Val	Gly	Phe	Ser	Gly	Phe	Arg	Met	Asp	Ala	Ile	Ala	His	Val
Asp	Thr	Asp	Phe	Thr	Arg	Asp	Trp	Ile	Asn	His	Val	Gln	Trp	Ala	Thr
Ser	Glu	Asp	Val	Phe	Phe	Val	Ala	Glu	Ala	Trp	Val	Ser	Asp	Ile	Asn
Gly	Tyr	Leu	Asp	Ala	Val	Asn	Thr	Pro	His	Leu	Arg	Ala	Phe	Asp	Phe
Asn	Leu	Arg	Glu	Asp	Phe	Val	Ala	Leu	Ser	Ser	Gly	Ser	Lys	Asp	Met
Arg	Trp	Trp	Gly	Gly	Leu	Val	Asn	Ser	Gln	His	Arg	Asp	Arg	Ala	Val
Thr	Phe	Val	Asp	Asn	His	Asp	Thr	Ser	Arg	Ala	Gly	Asn	Pro	Tyr	Gly
Met	Pro	Gln	Val	Ile	Asn	Tyr	Lys	Asn	Gln	Ala	Tyr	Ala	Tyr	Ile	Leu
Leu	Arg	Glu	His	Gly	Val	Pro	Thr	Val	Phe	Ala	Arg	Asp	Tyr	Asp	Glu
Phe	Gly	Met	Ala	Pro	Thr	Leu	Asp	Lys	Leu	Ile	Glu	Ala	Arg	Arg	Tyr
Phe	Ala	Tyr	Gly	Pro	Gly	His	Glu	Tyr	Ser	Gly	Asn	Thr	Glu	Ala	Val
Tyr	Ala	Tyr	Val	Arg	Glu	Gly	Leu	Ser	Thr	Val	Pro	Gly	Thr	Gly	Leu
Val	Met	Leu	Met	Ser	Gly	Arg	Asn	Trp	Gly	Gly	Gln	Gln	Ser	Phe	Thr
Ile	Asn	Ser	His	Gln	Pro	Asn	Thr	Thr	Phe	Tyr	Asp	Tyr	Thr	Gly	Asn
Val	Ser	Gly	Thr	Val	Thr	Thr	Asn	Ala	Gln	Gly	Tyr	Gly	Ser	Phe	Pro
Val	Asn	Met	Thr	Glu	Ser	Thr	Gly	Trp	Ser	Val	Trp	Val	Pro	Gln	Ser
Ile	Gly	Gly	Thr	Gln	Pro	Gly	Thr	Ile	Thr	Leu	Arg	Met	Thr	Lys	Asp
Val	Gly	Tyr	Gly	Phe	Ser	Leu	Phe	Phe	Thr	Gly	Ser	Ser	Ala	Glu	Leu
Thr	Asn	Trp	Gly	Gly	Gly	Val	Glu	Gly	Thr	Trp	Thr	Ser	Gly	Asn	Val
Trp	Glu	Val	Thr	Ile	Pro	Asp	Pro								

<210> 340
 <211> 1848
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 ctggactatc tcaaggagat ggggtgtacag gctttatggc tgtcaccaat ccattcctgca 180
 acatcctatc acggctatga cgtggaggac tacagttcgg taaatccagc ctacggcacg 240
 gaggcggatt tcgagtccct gctgaaagcg gcgcactcga aaggaatgaa aatctacctt 300
 gatttcgtgc tgaaccacac atccaaggaa catccgtggg tccttgaggg aaagtccgat 360
 ccggacagca ggtaccgaca ctattaccat ttctccacga cggccaagga cggatattcg 420
 cagaccgttt cgggcaccga cccagggaaa atcaacgtga agttcacctt taaatgcaat 480
 tcttccggca cacctcagac gcttaaggcc gaaaggggtg agagcgcggg caacgagggc 540
 actccgaatt ccggaaaaa cctctggtag ggttccgtga ccgaaagcac aatgccggaa 600
 ttctattctg ccggcgagga cacatatacc cttgcaatag aaaacttcga aaccgactgg 660
 ggcgtactgg tacgcaccag caagactgaa tggggcgcaa agaaattcgg cgcccaatcc 720
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 tgcgcagctg ctgacaaatg gataaaaatg ggcgttgacg gcttccgtct cgacgccgtc 960
 aagcacatat atgacaacga gaactccgac gagaaccgga ccttcctgaa gaaattctac 1020
 gaccattgca ataccaccta taaggctgca ggtcattcag gaaatatcta tatggtcggg 1080
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 cagggcgaa agcttggtta ctggggcacg aaaagcaatg gggacgaata tgtccgcacc 1440
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 ccggcaggag acaagctttc ggcaatgata ggctcaaacg gaagcgcgac ggtaaagaac 1800
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<210> 341
 <211> 615
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 Asp Phe Lys Gly Ile Glu Ser Lys Leu Asp Tyr Leu Lys Glu Met Gly
 35 40 45
 Val Gln Ala Leu Trp Leu Ser Pro Ile His Pro Ala Thr Ser Tyr His
 50 55 60
 Gly Tyr Asp Val Glu Asp Tyr Ser Ser Val Asn Pro Ala Tyr Gly Thr
 65 70 75 80
 Glu Ala Asp Phe Glu Ser Leu Leu Lys Ala Ala His Ser Lys Gly Met
 85 90 95
 Lys Ile Tyr Leu Asp Phe Val Leu Asn His Thr Ser Lys Glu His Pro
 100 105 110
 Trp Phe Leu Glu Gly Lys Ser Asp Pro Asp Ser Arg Tyr Arg His Tyr
 115 120 125
 Tyr His Phe Ser Thr Thr Ala Lys Asp Gly Tyr Ser Gln Thr Val Ser

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Gly 130	Thr 135	Asp 140	Pro 145	Gly 150	Lys 155	Ile 160	Asn 165	Val 170	Lys 175	Phe 180	Thr 185	Leu 190	Lys 195	Cys 200	Asn 205
Ser 145	Ser 150	Gly 155	Thr 160	Pro 165	Gln 170	Thr 175	Leu 180	Lys 185	Ala 190	Glu 195	Arg 200	Val 205	Glu 210	Ser 215	Ala 220
Val 155	Asn 160	Glu 165	Gly 170	Thr 175	Pro 180	Asn 185	Ser 190	Gly 195	Lys 200	Tyr 205	Leu 210	Trp 215	Tyr 220	Gly 225	Ser 230
Val 165	Thr 170	Glu 175	Ser 180	Thr 185	Met 190	Pro 195	Glu 200	Phe 205	Tyr 210	Ser 215	Ala 220	Gly 225	Glu 230	Asp 235	Thr 240
Tyr 175	Thr 180	Leu 185	Ala 190	Ile 195	Glu 200	Asn 205	Phe 210	Glu 215	Thr 220	Asp 225	Trp 230	Gly 235	Val 240	Leu 245	Val 250
Arg 185	Thr 190	Ser 195	Lys 200	Thr 205	Glu 210	Trp 215	Gly 220	Ala 225	Lys 230	Lys 235	Phe 240	Gly 245	Ala 250	Gln 255	Ser 260
Glu 195	Ser 200	Ala 205	Gln 210	Met 215	Phe 220	Val 225	Trp 230	Gly 235	Val 240	Pro 245	Leu 250	Asn 255	Leu 260	Lys 265	Ser 270
Asn 205	Ser 210	Asp 215	Tyr 220	Asp 225	Ile 230	Leu 235	Met 240	Pro 245	Trp 250	Met 255	Gly 260	Lys 265	Ile 270	Trp 275	Tyr 280
Gln 215	Ser 220	Val 225	Phe 230	Gly 235	Ser 240	Tyr 245	Met 250	Pro 255	Asp 260	Leu 265	Asn 270	Tyr 275	Gly 280	Ala 285	Ala 290
Ala 225	Ser 230	Cys 235	Glu 240	Glu 245	Ser 250	Glu 255	Ala 260	Phe 265	Lys 270	Glu 275	Val 280	Cys 285	Ala 290	Ala 295	Ala 300
Asp 235	Lys 240	Trp 245	Ile 250	Lys 255	Met 260	Gly 265	Val 270	Asp 275	Gly 280	Phe 285	Arg 290	Leu 295	Asp 300	Ala 305	Val 310
Lys 245	His 250	Ile 255	Tyr 260	Asp 265	Asn 270	Glu 275	Asn 280	Ser 285	Asp 290	Glu 295	Asn 300	Pro 305	Thr 310	Phe 315	Leu 320
Lys 255	Lys 260	Phe 265	Tyr 270	Asp 275	His 280	Cys 285	Asn 290	Thr 295	Thr 300	Tyr 305	Lys 310	Ala 315	Ala 320	Gly 325	His 330
Ser 265	Gly 270	Asn 275	Ile 280	Tyr 285	Met 290	Val 295	Gly 300	Glu 305	Gln 310	Trp 315	Ser 320	Glu 325	Pro 330	Asn 335	Tyr 340
Val 275	Thr 280	Pro 285	Tyr 290	Tyr 295	Lys 300	Gly 305	Leu 310	Asn 315	Ala 320	Phe 325	Phe 330	Glu 335	Phe 340	Ala 345	Phe 350
Cys 285	Trp 290	Arg 295	Leu 300	Thr 305	Asp 310	Ala 315	Ile 320	Asn 325	Gly 330	Ala 335	Lys 340	Gly 345	Ala 350	Gly 355	Phe 360
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Ala 315	Ser 320	Thr 325	Leu 330	Gly 335	Arg 340	Asn 345	Ala 350	Ser 355	Lys 360	Leu 365	Lys 370	Leu 375	Ala 380	Ala 385	Ala 390
Val 325	Leu 330	Leu 335	Thr 340	Ala 345	Gly 350	Gly 355	Glu 360	Pro 365	Tyr 370	Ile 375	Tyr 380	Gln 385	Gly 390	Glu 395	Glu 400
Leu 335	Gly 340	Tyr 345	Trp 350	Gly 355	Thr 360	Lys 365	Ser 370	Asn 375	Gly 380	Asp 385	Glu 390	Tyr 395	Val 400	Arg 405	Thr 410
Pro 345	Ile 350	Leu 355	Trp 360	Thr 365	Ala 370	Asp 375	Ala 380	Gly 385	Ser 390	Ala 395	Ala 400	Ala 405	Gly 410	Ala 415	Leu 420
Asn 355	Gly 360	Lys 365	Ile 370	Asp 375	Arg 380	Asn 385	Met 390	Leu 395	Thr 400	Ser 405	Asp 410	Ile 415	Ser 420	Val 425	Glu 430
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Gly 375	Val 380	Leu 385	Arg 390	Asp 395	Cys 400	Tyr 405	Lys 410	Ala 415	Leu 420	Ala 425	Lys 430	Gly 435	Ser 440	Phe 445	Gln 450
Glu 385	Lys 390	Asn 395	Gly 400	Ile 405	Ser 410	Asn 415	Gln 420	Ala 425	Ile 430	Cys 435	Ala 440	Trp 445	Tyr 450	Arg 455	Glu 460
Tyr 395	Asp 400	Gly 405	Gln 410	Lys 415	Ile 420	Leu 425	Val 430	Val 435	His 440	Asn 445	Phe 450	Ser 455	Ala 460	Ala 465	Pro 470
Val 405	Thr 410	Phe 415	Ala 420	Pro 425	Ala 430	Gly 435	Asp 440	Lys 445	Leu 450	Ser 455	Ala 460	Met 465	Ile 470	Gly 475	Ser 480
Asn 415	Gly 420	Ser 425	Ala 430	Thr 435	Val 440	Lys 445	Asn 450	Asp 455	Asn 460	Leu 465	Thr 470	Gly 475	Gly 480	Gly 485	Tyr 490
Ala 425	Ser 430	Ala 435	Leu 440	Phe 445	Ala 450	Thr 455	Val 460	Lys 465	Leu 470	Thr 475	Leu 480	Thr 485	Leu 490	Gly 495	Tyr 500
Ala 435	Ser 440	Ala 445	Leu 450	Phe 455	Ala 460	Thr 465	Val 470	Lys 475	Leu 480	Thr 485	Leu 490	Thr 495	Leu 500	Gly 505	Tyr 510
Ala 445	Ser 450	Ala 455	Leu 460	Phe 465	Ala 470	Thr 475	Val 480	Lys 485	Leu 490	Thr 495	Leu 500	Thr 505	Leu 510	Gly 515	Tyr 520
Ala 455	Ser 460	Ala 465	Leu 470	Phe 475	Ala 480	Thr 485	Val 490	Lys 495	Leu 500	Thr 505	Leu 510	Thr 515	Leu 520	Gly 525	Tyr 530
Ala 465	Ser 470	Ala 475	Leu 480	Phe 485	Ala 490	Thr 495	Val 500	Lys 505	Leu 510	Thr 515	Leu 520	Thr 525	Leu 530	Gly 535	Tyr 540
Ala 475	Ser 480	Ala 485	Leu 490	Phe 495	Ala 500	Thr 505	Val 510	Lys 515	Leu 520	Thr 525	Leu 530	Thr 535	Leu 540	Gly 545	Tyr 550
Ala 485	Ser 490	Ala 495	Leu 500	Phe 505	Ala 510	Thr 515	Val 520	Lys 525	Leu 530	Thr 535	Leu 540	Thr 545	Leu 550	Gly 555	Tyr 560
Ala 495	Ser 500	Ala 505	Leu 510	Phe 515	Ala 520	Thr 525	Val 530	Lys 535	Leu 540	Thr 545	Leu 550	Thr 555	Leu 560	Gly 565	Tyr 570
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Ala 515	Ser 520	Ala 525	Leu 530	Phe 535	Ala 540	Thr 545	Val 550	Lys 555	Leu 560	Thr 565	Leu 570	Thr 575	Leu 580	Gly 585	Tyr 590
Ala 525	Ser 530	Ala 535	Leu 540	Phe 545	Ala 550	Thr 555	Val 560	Lys 565	Leu 570	Thr 575	Leu 580	Thr 585	Leu 590	Gly 595	Tyr 600
Ala 535	Ser 540	Ala 545	Leu 550	Phe 555	Ala 560	Thr 565	Val 570	Lys 575	Leu 580	Thr 585	Leu 590	Thr 595	Leu 600	Gly 605	Tyr 610
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<210> 342
 <211> 2061
 <212> DNA
 <213> Unknown

<220>

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<223> Obtained from an environmental sample

<400> 342

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<210> 343

<211> 686

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(23)

<400> 343

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35     40     45
Lys Gly Tyr Ala Ala Val Gln Val Ser Pro Pro Gln Lys Ser Val Glu
50     55     60
Gly Ser Phe Trp Trp Thr Arg Tyr Gln Pro Val Ser Tyr Ser Ile Glu
65     70     75     80
Gly Arg Ser Gly Thr Arg Thr Glu Phe Ala Ser Met Val Ser Arg Cys
85     90     95
Lys Ala Val Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn His Met
100    105    110
Ala Ala Gly Asn Arg Asn Phe Pro Glu Val Pro Tyr Gly Val Asn Asp
115    120    125
Phe His Thr Cys Thr Gly Asn Ile Asp Tyr Thr Asn Ala Trp Gln Val

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Gly	Val	Ala	Gly	Phe	Arg	165	Ile	Asp	Ala	Ala	Lys	170	His	Ile	Ala	Ser	Ala	
Asp	Ile	Ala	Asn	Ile	Val	180	Ser	Arg	Leu	Asn	Gly	185	Asn	Pro	Tyr	Ile	Phe	
Gln	Glu	Val	Ile	Gly	Ala	195	Ala	Asn	Glu	Pro	Ile	200	Thr	Pro	Ala	Gln	Tyr	
210	Thr	Tyr	Ile	Gly	Asp	215	Val	Thr	Glu	Phe	Asn	220	Phe	Ser	Asn	Thr	Leu	
225	Asn	Tyr	Phe	Lys	Gly	230	Arg	Ala	Pro	Leu	Lys	235	Asp	Leu	Arg	Asn	Ile	
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Asn	His	Asp	Asn	Gln	Arg	260	Gln	Asn	Thr	Ser	Asn	265	Ile	Val	Thr	His	Lys	
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385	Ala	Gly	Ser	Tyr	Cys	390	Asn	Val	Ile	Ser	Gly	395	Asp	Phe	Ala	Asn	Gly	
Cys	Thr	Gly	Ala	Ser	Ile	405	Thr	Val	Ala	Ala	Asn	410	Gly	Thr	Ala	Thr	Phe	
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465	Ile	Trp	Met	Gly	Ser	460	Ala	Thr	Phe	Ser	Gly	465	Ser	Ser	Thr	Asp	Arg	
Lys	Phe	Asp	Ala	Tyr	Gly	470	Asn	Trp	Val	Thr	Asn	475	Trp	Gly	Asn	Asn	Asn	
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580	Ala	Ile	Lys	Leu	Glu	570	Pro	Thr	Ser	Tyr	Pro	575	Ile	Trp	Thr	Gly	Thr	
595	Gln	Leu	Pro	Ser	Asn	585	Thr	Ala	Val	Glu	Trp	590	Lys	Cys	Leu	Lys	Arg	
610	Glu	Asn	Asn	Ala	Ala	595	Gly	Val	Val	Trp	Glu	600	Ser	Gly	Ser	Asn	Thr	
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675

680

685

<210> 344
 <211> 1980
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 tgcgaaaacc atctcggccc aaaaggttat gcggcggtag aagtatcacc gccacaaaaa 180
 tctatcagcg gcaaccaatg gtggacacgc taccaaccgg tcagctattc cattgaaggc 240
 cgtttaggaa cccgcgcgga attcgccagc atggtttcac gctgtaacgc agtgggagtg 300
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 gccggtacct attgcgatgt gatcaagggt gatctttcta acggcgtagt taccggtccg 1260
 accattaccg tcggtgccga cggcaatgcg agttttgctg tcggatccaa aggtgtcttcg 1320
 gctattcatg ctgacgtgt agtcggtccg attaccggca atgaatgcgt ttacgacacc 1380
 atgaatttac gcggcacatt taacagctgg gaatcacacg ccatgacatg tgccaatggt 1440
 atctggatcg gcagcgccaa cttttccggc aacagcaatg aacgattcaa gtttgacgtc 1500
 tatggtgatt ggcaaacc aa ttttggtgac gccaataacg atggtgttgc caatctcggc 1560
 ggcgcggata tcgcgatagc ctctgccggg gattacacca tcacctttaa cgatgcgagc 1620
 tttgcttata cttttacccc ggcaaccagc agcagctcaa gttcgaattc cagcagcaca 1680
 cccggtactg tggcggtgaa tttcacttgt gaaaatggca gtacttatgt gggacaaagt 1740
 gtttatgtgg tgggtaacaa tgcagccatc ggcggttggt ctccaacgaa cgcggtaaag 1800
 cttaatcccc tcagttatcc cacctggacc ggcacgattc aattaccggc gaacacagct 1860
 attgaatgga aatgtttgaa gcgagaagaa aataatcctg cagcgggtat tgagtgggaa 1920
 gcaggtggaa ataatagtct aaatacaggg agcgatgcga atacgacggg cgggttttga 1980

<210> 345
 <211> 659
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(22)

<400> 345
 Met Lys Lys Leu Leu Ile Thr Leu Leu Ile Phe Pro Gly Phe Leu Phe
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 Ser Phe Ser Ala Val Ala Gln Thr Thr Phe Val His Leu Phe Glu Trp
 20 25 30
 Gln Trp Asn Ala Ile Ala Ser Glu Cys Glu Asn His Leu Gly Pro Lys
 35 40 45
 Gly Tyr Ala Ala Val Gln Val Ser Pro Pro Gln Lys Ser Ile Ser Gly
 50 55 60
 Asn Gln Trp Trp Thr Arg Tyr Gln Pro Val Ser Tyr Ser Ile Glu Gly
 65 70 75 80
 Arg Leu Gly Thr Arg Ala Glu Phe Ala Ser Met Val Ser Arg Cys Asn

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90

				85				90					95		
Ala	Val	Gly	Val	Asp	Ile	Tyr	Val	Asp	Ala	Val	Ile	Asn	His	Met	Ser
			100					105					110		
Ala	Gly	Asn	Arg	Ser	Phe	Pro	Glu	Val	Pro	Phe	Gly	Thr	Asn	Asp	Phe
		115					120					125			
His	Thr	Cys	Thr	Ser	Asp	Ile	Asp	Tyr	Ser	Asp	Val	Trp	Ser	Ile	Gln
						135					140				
Asn	Cys	Asp	Leu	Val	Gly	Leu	Asn	Asp	Leu	Lys	Thr	Glu	Ser	Asp	Tyr
					150					155				160	
Val	Arg	Gly	Lys	Ile	Ala	Asp	Tyr	Met	Asn	Asp	Leu	Ile	Ser	Leu	Gly
				165					170					175	
Val	Thr	Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His	Met	Pro	Ser	Ser	Asp
			180					185					190		
Ile	Ala	Asn	Ile	Val	Ser	Arg	Leu	Asn	Gly	Asp	Pro	Tyr	Ile	Phe	Gln
		195					200				205				
Glu	Val	Ile	Gly	Ala	Gly	Gly	Glu	Pro	Ile	Gln	Pro	Ser	Gln	Tyr	Thr
		210				215					220				
Tyr	Val	Gly	Asp	Val	Thr	Glu	Phe	Asn	Phe	Ser	Asn	Thr	Leu	Gly	His
					230					235					240
Tyr	Phe	Lys	Gly	Arg	Ala	Thr	Leu	His	Glu	Leu	Arg	Asn	Ile	Gly	Val
				245					250					255	
Trp	Ser	Gly	Trp	Leu	Asn	Ser	Ala	Asp	Ala	Ile	Thr	Phe	Val	Ala	Asn
			260					265					270		
His	Asp	Asn	Gln	Arg	Gln	Asn	Thr	Asn	Asn	Ile	Ile	Thr	His	Lys	Asp
		275					280					285			
Gly	Met	Asn	Leu	Asn	Asn	Ile	Ala	His	Val	Phe	Ala	Leu	Ala	Trp	Pro
		290				295					300				
Tyr	Gly	Tyr	Pro	Lys	Val	Met	Ser	Ser	Tyr	Asp	Trp	Asn	Asp	His	Asp
					310					315				320	
Gln	Gly	Pro	Pro	Thr	Asn	Ala	Ala	Asn	Thr	Cys	Ser	Asn	Gly	Trp	Leu
				325					330					335	
Cys	Glu	His	Arg	Asn	Arg	Glu	Ile	Ala	Asn	Met	Val	Gly	Phe	Arg	Asn
			340					345					350		
Ala	Thr	Gln	Ser	Ala	Phe	Tyr	Val	Ser	Asn	Trp	Trp	Asp	Asn	Gly	Ala
		355					360					365			
Asn	Gln	Ile	Ala	Phe	Gly	Arg	Gly	Asp	Arg	Gly	Phe	Val	Val	Ile	Asn
		370				375					380				
Gly	Glu	Ser	Gly	Gly	Thr	Leu	Asn	Thr	Thr	Leu	Gln	Thr	Gly	Leu	Ala
					390					395					400
Ala	Gly	Thr	Tyr	Cys	Asp	Val	Ile	Lys	Gly	Asp	Phe	Ser	Asn	Gly	Val
				405					410					415	
Cys	Thr	Gly	Pro	Thr	Ile	Thr	Val	Gly	Ala	Asp	Gly	Asn	Ala	Ser	Phe
			420					425					430		
Ala	Val	Gly	Ser	Lys	Gly	Ala	Ser	Ala	Ile	His	Ala	Asp	Ala	Val	Val
		435													

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625 Ala Gly Gly Asn Asn 630 Ser Leu Asn Thr Gly 635 Ser Asp Ala Asn Thr 640 Thr
645 650 655
Gly Gly Phe

<210> 346
<211> 1398
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 346
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gccgacgtca tactgcacgc cttcaactgg caatacagcg aagtcaccgc caaggccgat 120
ctcatcaagg ccgccggcta caagcagggtg ctcatctcgc cgcccctcaa atcatcgggc 180
aacgagtggg gggcacgcta tcaaccgcag gatctgcgcc tcatcgatac tccccttggc 240
aacaagcagg atctcgagca gctgatcgcc gccatgcagg cacgaggcat agccgtctat 300
gccgatattg tgctcaacca catggccaac gagagctgga agcgagcga tctcaactac 360
cccggcagcg agctgctcgg ccagtatgcg gccaaccccg cctacttcga acgccagaag 420
ctgttcggcg atctcgcca gaacttgctg gccggatcgg attttcaccc cgaaggctgc 480
atcaccgact ggagcgatcc gggccacgtt cagtactggc gcctgtgtgg cggcgccggt 540
gacaaggggc tgccggatct tgatcccaac aactgggtag tgagccagca gcaggcttat 600
ctcaaggcgc tcaaggggat ggggatcaag ggctttcggg tcgatgcggt caagcacatg 660
agcgattacc agatcaatgc cgtgttcacc ccagagataa aacagggcag gcacgtcttt 720
ggcgaggtaa tcaccacggg gggcgccggc agcacggatt acgaacgctt ccttaaacc 780
tatctcgaca gcagcggta gggggcctat gactttccgc tgtttgctc cctgcgcggg 840
gcgctggggt atggcggcag catgagcctg ctggccgacc ccggtgccta cggtcaggcc 900
ctgccgggca gccgcgccgt caccttcgcc atcaccacg acattcccac caacgatggc 960
tttcgtacc agatcctgaa ccagaccgac gagaagctgg cctatgccta cctgctcggg 1020
cgcgatgggt gctcgccgt tgtctattcc gatcacggcg agaccaggc caaggatggg 1080
ctgcgctggc aggattatta ccagcgcgcg gatctcaaag gcatgatccg cttccataac 1140
gcgctgcagg gccagcccat gcaactggtc ggcagcggcg actgcttcgt gctgttcaaa 1200
cgtggcaagc aggggctggt cgggtgtcaac aagtgcgagt acgagcaaga gttctggctc 1260
gataccgcca aattcgagat gaactgggtat cgcaactacc gggatgtgct cgatcaaaac 1320
gccgtcatca atgtgcagag ccagtgggtg cgggtcgcca ttcccggccg cagcgcccgg 1380
ctctggctgc aagaataa 1398

<210> 347
<211> 465
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(21)

<400> 347
Met Lys Lys Thr Ala Gly Ile Trp Val Ile Ala Gly Met Leu Ile Ala
1 5 10 15
Pro Leu Ala His Ala Asp Val Ile Leu His Ala Phe Asn Trp Gln Tyr
20 25 30
Ser Glu Val Thr Ala Lys Ala Asp Leu Ile Lys Ala Ala Gly Tyr Lys
35 40 45
Gln Val Leu Ile Ser Pro Leu Lys Ser Ser Gly Asn Glu Trp Trp
50 55 60
Ala Arg Tyr Gln Pro Gln Asp Leu Arg Leu Ile Asp Thr Pro Leu Gly
65 70 75 80
Asn Lys Gln Asp Leu Glu Gln Leu Ile Ala Ala Met Gln Ala Arg Gly
85 90 95
Ile Ala Val Tyr Ala Asp Ile Val Leu Asn His Met Ala Asn Glu Ser
100 105 110
Trp Lys Arg Ser Asp Leu Asn Tyr Pro Gly Ser Glu Leu Leu Gly Gln
115 120 125

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Tyr Ala Ala Asn Pro Ala Tyr Phe Glu Arg Gln Lys Leu Phe Gly Asp
 130 135 140
 Leu Gly Gln Asn Leu Leu Ala Gly Ser Asp Phe His Pro Glu Gly Cys
 145 150 155 160
 Ile Thr Asp Trp Ser Asp Pro Gly His Val Gln Tyr Trp Arg Leu Cys
 165 170 175
 Gly Gly Ala Gly Asp Lys Gly Leu Pro Asp Leu Asp Pro Asn Asn Trp
 180 185 190
 Val Val Ser Gln Gln Gln Ala Tyr Leu Lys Ala Leu Lys Gly Met Gly
 195 200 205
 Ile Lys Gly Phe Arg Val Asp Ala Val Lys His Met Ser Asp Tyr Gln
 210 215 220
 Ile Asn Ala Val Phe Thr Pro Glu Ile Lys Gln Gly Met His Val Phe
 225 230 235 240
 Gly Glu Val Ile Thr Thr Gly Gly Ala Gly Ser Thr Asp Tyr Glu Arg
 245 250 255
 Phe Leu Lys Pro Tyr Leu Asp Ser Ser Gly Gln Gly Ala Tyr Asp Phe
 260 265 270
 Pro Leu Phe Ala Ser Leu Arg Gly Ala Leu Gly Tyr Gly Ser Met
 275 280 285
 Ser Leu Leu Ala Asp Pro Gly Ala Tyr Gly Gln Ala Leu Pro Gly Ser
 290 295 300
 Arg Ala Val Thr Phe Ala Ile Thr His Asp Ile Pro Thr Asn Asp Gly
 305 310 315 320
 Phe Arg Tyr Gln Ile Leu Asn Gln Thr Asp Glu Lys Leu Ala Tyr Ala
 325 330 335
 Tyr Leu Leu Gly Arg Asp Gly Gly Ser Pro Leu Val Tyr Ser Asp His
 340 345 350
 Gly Glu Thr Gln Ala Lys Asp Gly Leu Arg Trp Gln Asp Tyr Tyr Gln
 355 360 365
 Arg Ala Asp Leu Lys Gly Met Ile Arg Phe His Asn Ala Leu Gln Gly
 370 375 380
 Gln Pro Met Gln Leu Val Gly Ser Gly Asp Cys Phe Val Leu Phe Lys
 385 390 395 400
 Arg Gly Lys Gln Gly Leu Val Gly Val Asn Lys Cys Glu Tyr Glu Gln
 405 410 415
 Glu Phe Trp Leu Asp Thr Ala Lys Phe Glu Met Asn Trp Tyr Arg Asn
 420 425 430
 Tyr Arg Asp Val Leu Asp Gln Asn Ala Val Ile Asn Val Gln Ser Gln
 435 440 445
 Trp Val Arg Val Ala Ile Pro Ala Arg Ser Ala Arg Leu Trp Leu Gln
 450 455 460
 Glu
 465

<210> 348
 <211> 1488
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 348
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 agcctatggc agcaagtggc agagaaagca gaggaattag ctgcggtggg ggctcacgtcg 120
 ctctggttgc cgcctgccta caaggaacc gggggcggtt atgatgtggg ctacggcggtt 180
 tatgacctgt ttgatctggg tgagtttgac caaaaaggct cgggtgcgcac aaagtatggc 240
 actaaggatg aatacctcgc tgcgattaag caggcacaac gtttaggcac caggatctat 300
 gccgatgttg tgtttaacca caagttgggt gcggatgaag aagaggaagt ggaagccacg 360
 cccttcaatc ctgacaatcg caatcatacg gttggcgact atcagaaaat caaagcctgg 420
 acgggggttca cttttcccg tgcaggtgac aaatactcca gcatgaagtg gcattggtgg 480
 cactttgatg cgatcgacta taacgcttac aaaccggaag agaacgctat ctacttgctc 540
 aagggttaagg agttcgatcg taacgttgac ctcgaaaaag gtaacttcga ttacttgatg 600
 ggatgtgatt tagatatgga tcatccggaa gtgattggag agttgaaata ctggggtgaa 660
 tggatcatca agacaaccaa tgtcgatggg ttccggtttg atgccgtaaa gcatgtgtct 720
 gctgacttct ttcgtgaatg gttgaacacc atgcgggtag agatgtcttt 780
 gccgtaggtg agtactggtc ctatgatgtc gaggcactac acagcttcac cgaacaacg 840

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aatgggaaag	tcacgttggt	tgatgcccc	ctccactaca	actttcactg	cgccagtacc	900
tcaggcaaca	gttatgacat	gcgccaatc	tttgacggca	ccctggtgca	gcaacaacct	960
gctctggcag	tcaccctggt	agaaaaccat	gattctcaac	ccctgcaatc	gctggaatcc	1020
atcgttgaag	ggtggttcaa	gcccctggca	tatgccttga	ttttgctgag	gcgggaaggg	1080
tatccctgta	ttttttatgc	tgactactac	ggtgctcact	acaaagatca	gggcagggat	1140
ggccaggagt	atgagatttg	gatggacagc	caccaattct	tgatcgacaa	aatgctgggt	1200
gctcggcaaa	cctttgccta	tggcgatcaa	tacgattatt	ttgaccacgc	caatacgatc	1260
ggctggacac	gcctaggcac	agaggagcat	cctggaggca	tggcggttgt	gctcagtaat	1320
ggcgatgctg	gcagcaaata	catggaagtt	gggcagccta	actgcaccta	tgttgacatc	1380
acggaacata	ttaatgaacc	gatcgtcacc	aatgccgatg	gatgggctga	ctttcgtctg	1440
gagcctgggt	cagtatcagt	ttgggttccc	caagcagcgg	cacaatag		1488

<210> 349
 <211> 495
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 349

Met	Ser	Glu	Leu	Asn	Gly	Val	Met	Met	Gln	Tyr	Phe	His	Trp	Tyr	Ser
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Ala	Ala	Asp	Gly	Ser	Leu	Trp	Gln	Gln	Val	Ala	Glu	Lys	Ala	Glu	Glu
			20					25					30		
Leu	Ala	Ala	Val	Gly	Val	Thr	Ser	Leu	Trp	Leu	Pro	Pro	Ala	Tyr	Lys
		35					40					45			
Gly	Thr	Gly	Gly	Gly	Tyr	Asp	Val	Gly	Tyr	Gly	Val	Tyr	Asp	Leu	Phe
	50					55					60				
Asp	Leu	Gly	Glu	Phe	Asp	Gln	Lys	Gly	Ser	Val	Arg	Thr	Lys	Tyr	Gly
65					70					75					80
Thr	Lys	Asp	Glu	Tyr	Leu	Ala	Ala	Ile	Lys	Gln	Ala	Gln	Arg	Leu	Gly
				85					90					95	
Ile	Arg	Ile	Tyr	Ala	Asp	Val	Val	Phe	Asn	His	Lys	Leu	Gly	Ala	Asp
			100					105					110		
Glu	Glu	Glu	Glu	Val	Glu	Ala	Thr	Pro	Phe	Asn	Pro	Asp	Asn	Arg	Asn
		115					120					125			
His	Thr	Val	Gly	Asp	Tyr	Gln	Lys	Ile	Lys	Ala	Trp	Thr	Gly	Phe	Thr
	130					135					140				
Phe	Pro	Gly	Arg	Gly	Asp	Lys	Tyr	Ser	Ser	Met	Lys	Trp	His	Trp	Trp
145					150					155					160
His	Phe	Asp	Ala	Ile	Asp	Tyr	Asn	Ala	Tyr	Lys	Pro	Glu	Glu	Asn	Ala
				165					170					175	
Ile	Tyr	Leu	Leu	Lys	Gly	Lys	Glu	Phe	Asp	Arg	Asn	Val	Asp	Leu	Glu
			180					185					190		
Lys	Gly	Asn	Phe	Asp	Tyr	Leu	Met	Gly	Cys	Asp	Leu	Asp	Met	Asp	His
		195					200					205			
Pro	Glu	Val	Ile	Gly	Glu	Leu	Lys	Tyr	Trp	Gly	Glu	Trp	Tyr	Ile	Lys
	210					215					220				
Thr	Thr	Asn	Val	Asp	Gly	Phe	Arg	Phe	Asp	Ala	Val	Lys	His	Val	Ser
225					230					235					240
Ala	Asp	Phe	Phe	Arg	Glu	Trp	Leu	Asp	His	Val	Glu	His	His	Ala	Gly
				245					250					255	
Arg	Asp	Val	Phe	Ala	Val	Gly	Glu	Tyr	Trp	Ser	Tyr	Asp	Val	Glu	Ala
			260					265					270		
Leu	His	Ser	Phe	Ile	Glu	Thr	Thr	Asn	Gly	Lys	Val	Thr	Leu	Phe	Asp
		275					280					285			
Ala	Pro	Leu	His	Tyr	Asn	Phe	His	Cys	Ala	Ser	Thr	Ser	Gly	Asn	Ser
	290					295					300				
Tyr	Asp	Met	Arg	Gln	Ile	Phe	Asp	Gly	Thr	Leu	Val	Gln	Gln	Gln	Pro
305					310					315					320
Ala	Leu	Ala	Val	Thr	Leu	Val	Glu	Asn	His	Asp	Ser	Gln	Pro	Leu	Gln
				325					330					335	
Ser	Leu	Glu	Ser	Ile	Val	Glu	Gly	Trp	Phe	Lys	Pro	Leu	Ala	Tyr	Ala
			340					345					350		
Leu	Ile	Leu	Leu	Arg	Arg	Glu	Gly	Tyr	Pro	Cys	Ile	Phe	Tyr	Ala	Asp
		355					360					365			
Tyr	Tyr	Gly	Ala	His	Tyr	Lys	Asp	Gln	Gly	Arg	Asp	Gly	Gln	Glu	Tyr

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370	375	380															
Glu Ile Trp Met Asp Ser His Gln Phe Leu Ile Asp Lys Met Leu Val																	
385	390	395	400														
Ala Arg Gln Thr Phe Ala Tyr Gly Asp Gln Tyr Asp Tyr Phe Asp His																	
405	410	415															
Ala Asn Thr Ile Gly Trp Thr Arg Leu Gly Thr Glu Glu His Pro Gly																	
420	425	430															
Gly Met Ala Val Val Leu Ser Asn Gly Asp Ala Gly Ser Lys Tyr Met																	
435	440	445															
Glu Val Gly Gln Pro Asn Cys Thr Tyr Val Asp Ile Thr Glu His Ile																	
450	455	460															
Asn Glu Pro Ile Val Thr Asn Ala Asp Gly Trp Ala Asp Phe Arg Cys																	
465	470	475	480														
Glu Pro Gly Ser Val Ser Val Trp Val Pro Gln Ala Ala Ala Gln																	
485	490	495															

<210> 350
 <211> 2001
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 350

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gagtgcgaaa	catttttagg	gccaaatggt	tttgccgcag	ttcaagtgtc	tccgccaaac	180
gagcacattc	agggttcgca	gtggtggacc	cgctatcaac	cagttagtta	tcaactcgag	240
agtcgtggtg	gtagccgtgc	cgcattttatc	aacatgggtgc	aacgttgtaa	tgctgcgggg	300
gtgtcaattt	atgtcgacgc	cgtgatcaac	catatggcca	atggcagtgg	cgttggtacc	360
gccggcagta	attacggcaa	ccgccaatat	ccaattttact	caggccaaga	tttccatgct	420
tcttggtgca	ttaatggctc	tgattacggc	aataatcggt	ggcgtgtaca	gaattgcgaa	480
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caagaagtga	tagaccaagg	tggcgaagca	attagctcaa	gtgagtacac	aggtcagggc	720
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aaagacggcc	ggatgtatga	cttagccaac	gtgttttatgc	tggcttacc	ctacggctat	960
acccaagtta	tgctcgagtta	tgactataaa	ggcgataccg	acgctggcgg	tcctggcatt	1020
cgggtccatc	aaaaatggta	ggtgaactgt	tttggcactg	actggaagtg	tgagcatcgt	1080
tcggagttata	tcgcaggggc	agtgcaattt	cgcaacaaca	ctgttaatga	atggcgcgtc	1140
accaattggt	gggacaacgg	caacaaccag	attgcgtttg	gtcgtgctgc	tgctgggttt	1200
gtcgcgatca	acaaagaaag	ttatgcatta	aataccagct	tggcaaccag	tatggcgcca	1260
ggtcagtatt	gtaacgtgct	caaaggtaaa	cgcagcagtg	acaaaaatc	ttgtacaggt	1320
gaggtgatta	ctgtgggttg	cgacggtcgg	atccaagcga	atgtcgccgc	ttgggatgcg	1380
tttgcgattc	atcaagatag	caaattgacg	actggcggcg	gtacgccggg	cgcggaattg	1440
cagcgactcg	tgggttttat	ccaagcacag	actctgagtg	gccaagatat	gtttgtccgc	1500
ggtggtcttg	atcatcaagt	ggcactgaac	caaaaagggt	taagtgtcac	cgcgacgaac	1560
ttcttggtg	caatgcctat	tcgccatcgg	aatctgaaaa	acgcgactac	agcgccatgg	1620
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gaaggtacgc	cgctcgattg	gaccactaat	ctttggccaa	atagttgggg	cacattacgc	1740
acagtgccgg	tggttggtta	tggtcaagaa	gcccttaaca	cgtttgcca	acattactgg	1800
atgatggatg	tagatatgga	ctgcagcaaa	accctcaacg	gttggtttga	actgaaggct	1860
tttgtagaaa	acggtcaagg	ttgggaaggc	gatatcagcc	aagcaaatcc	accttacgcg	1920
acaaaaacc	atttagcgca	atgcggcaaa	ctaaacaagt	ttgaatttgg	cagctcggcg	1980
gcattcatca	ccaacttgta	a				2001

<210> 351
 <211> 666
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<221> SIGNAL
<222> (1) . . . (21)

<400> 351

Met 1	Lys	Leu	Ser	His 5	Leu	Ala	Ala	Ala	Leu 10	Leu	Leu	Ala	Ser	Ser 15	Ser
Thr	Ala	Ser	Leu 20	Ala	Ala	Ser	Pro	Thr 25	Thr	Phe	Val	His	Leu 30	Phe	Glu
Trp	Arg	Trp 35	Ser	Asp	Val	Ala	Gln 40	Glu	Cys	Glu	Thr	Phe 45	Leu	Gly	Pro
Asn	Gly 50	Phe	Ala	Ala	Val	Gln 55	Val	Ser	Pro	Pro	Asn 60	Glu	His	Ile	Gln
Gly 65	Ser	Gln	Trp	Trp	Thr 70	Arg	Tyr	Gln	Pro	Val 75	Ser	Tyr	Gln	Leu	Glu 80
Ser	Arg	Gly	Gly	Ser 85	Arg	Ala	Ala	Phe	Ile 90	Asn	Met	Val	Gln	Arg 95	Cys
Asn	Ala	Ala	Gly 100	Val	Ser	Ile	Tyr	Val 105	Asp	Ala	Val	Ile	Asn 110	His	Met
Ala	Asn	Gly 115	Ser	Gly	Val	Gly	Thr 120	Ala	Gly	Ser	Asn	Tyr 125	Gly	Asn	Arg
Gln	Tyr 130	Pro	Ile	Tyr	Ser	Gly 135	Gln	Asp	Phe	His	Ala 140	Ser	Cys	Ala	Ile
Asn 145	Gly	Ser	Asp	Tyr	Gly 150	Asn	Asn	Arg	Trp	Arg 155	Val	Gln	Asn	Cys	Glu 160
Leu	Ser	Gly	Leu	Pro 165	Asp	Leu	Asn	Thr	Gly 170	Ser	Pro	Tyr	Val	Gln 175	Asn
Thr	Ile	Ala	Ala 180	Tyr	Met	Asn	Glu	Leu 185	Thr	Ser	Leu	Gly	Val 190	Lys	Gly
Phe	Arg	Leu 195	Asp	Ala	Ala	Lys	His 200	Met	Ser	Val	Glu	Asp 205	Ile	Ala	Ala
Ile	Arg 210	Gly	Lys	Leu	Thr	Gly 215	Ala	Pro	Leu	Ile	Tyr 220	Gln	Glu	Val	Ile
Asp 225	Gln	Gly	Gly	Glu	Ala 230	Ile	Ser	Ser	Ser	Glu 235	Tyr	Thr	Gly	Gln	Gly 240
Leu	Val	Ile	Glu	Phe 245	Lys	Tyr	Ser	Thr	Gln 250	Leu	Gly	Asn	Val	Phe 255	Lys
Thr	Gly	Arg	Leu 260	Ala	Ser	Leu	Arg	Asn 265	Phe	Gly	Glu	Gly	Trp 270	Gly	Phe
Leu	Pro	Ser 275	Ser	Arg	Ala	Val	Val 280	Phe	Val	Asp	Asn	His 285	Asp	Asn	Gln
Arg	Gly 290	His	Gly	Gly	Ala	Gly 295	Asn	Val	Val	Thr	Tyr 300	Lys	Asp	Gly	Arg
Met 305	Tyr	Asp	Leu	Ala	Asn 310	Val	Phe	Met	Leu	Ala 315	Tyr	Pro	Tyr	Gly	Tyr 320
Thr	Gln	Val	Met	Ser 325	Ser	Tyr	Asp	Tyr	Lys 330	Gly	Asp	Thr	Asp	Ala 335	Gly
Gly	Pro	Gly	Ile 340	Pro	Val	His	Gln	Asn 345	Gly	Gln	Leu	Asn	Cys 350	Phe	Gly
Thr	Asp	Trp 355	Lys	Cys	Glu	His	Arg 360	Trp	Ser	Tyr	Ile	Ala 365	Gly	Ala	Val
Gln	Phe	Arg	Asn	Asn	Thr	Val 375	Asn	Glu	Trp	Arg	Val 380	Thr	Asn	Trp	Trp
Asp 385	Asn	Gly	Asn	Asn	Gln 390	Ile	Ala	Phe	Gly	Arg 395	Ala	Ala	Ala	Gly	Phe 400
Val	Ala	Ile	Asn	Lys 405	Glu	Ser	Tyr	Ala	Leu 410	Asn	Thr	Ser	Leu	Ala 415	Thr
Ser	Met	Ala	Pro 420	Gly	Gln	Tyr	Cys	Asn 425	Val	Leu	Lys	Gly	Lys 430	Arg	Ser
Ser	Asp	Gln 435	Lys	Ser	Cys	Thr	Gly 440	Glu	Val	Ile	Thr	Val 445	Gly	Gly	Asp
Gly	Arg 450	Ile	Gln	Ala	Asn	Val 455	Ala	Ala	Trp	Asp	Ala 460	Phe	Ala	Ile	His
Gln 465	Asp	Ser	Lys	Leu	Thr 470	Thr	Gly	Gly	Gly	Thr 475	Pro	Gly	Ala	Asp	Trp 480
Gln	Arg	Thr	Val	Val 485	Phe	Ile	Gln	Ala	Gln 490	Thr	Leu	Ser	Gly	Gln 495	Asp
Met	Phe	Val	Arg 500	Gly	Gly	Leu	Asp	His 505	Gln	Val	Ala	Leu	Asn 510	Gln	Lys

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Gly	Leu	Ser 515	Cys	Thr	Ala	Thr	Asn 520	Phe	Leu	Cys	Ala	Met 525	Pro	Ile	Arg
His	Arg 530	Asn	Leu	Lys	Asn	Ala 535	Thr	Thr	Ala	Pro	Trp 540	Lys	Ala	Asn	Asp
Asn 545	Tyr	Leu	Asp	Trp	Tyr 550	Gly	Arg	Glu	Ala	Ser 555	Gln	Ser	Ala	Thr	Ala 560
Glu	Gly	Thr	Pro	Leu 565	Asp	Trp	Thr	Thr	Asn 570	Leu	Trp	Pro	Asn	Ser 575	Trp
Gly	Thr	Leu	Arg 580	Thr	Val	Pro	Val	Asp 585	Gly	Tyr	Gly	Gln	Glu 590	Ala	Leu
Asn	Thr	Phe 595	Gly	Gln	His	Tyr	Trp 600	Met	Met	Asp	Val	Asp 605	Met	Asp	Cys
Ser	Lys 610	Thr	Leu	Asn	Gly	Trp 615	Phe	Glu	Leu	Lys	Ala 620	Phe	Val	Lys	Asn
Gly 625	Gln	Gly	Trp	Glu	Gly 630	Asp	Ile	Ser	Gln	Ala 635	Asn	Pro	Pro	Tyr	Ala 640
Thr	Lys	Asn	His	Leu 645	Ala	Gln	Cys	Gly	Lys 650	Leu	Asn	Lys	Phe	Glu 655	Phe
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<211>	1263
<212>	DNA
<213>	Unknown

<220>
<223> obtained from an environmental sample

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tccgacatca	aggccagcgg	cttcaccatg	gtgtggttcc	caccctccgg	ggactcggcg		240
gccaatgagg	gctaccgtcc	cgccgagctc	tacgtgcaga	gcagtaagta	cggcacggac		300
gcgcagctca	agagcgccat	gggtctgctg	cacacctacg	gggtgaaggc	catcgcggac		360
atcgtggtca	accaccgcgt	gggcaccacc	aactgggcgg	acttcaccaa	ccccacctgg		420
ggctcgtggt	cggtggtgaa	gggcgacgag	tggacgagcg	ccaccggcaa	ctacgacacc		480
ggtgatggct	tcagcgcgcg	gcgcgacctg	gaccacacga	acggcacctg	gcagagcgac		540
ctgaaggcct	ggatgaactg	gctcaaggtc	agcatcggct	acgacggctg	gcgctatgac		600
tacgtgaagg	gctacggcgg	ctcgtacgtg	ggcggttaca	acagcgccac	ggtgccctac		660
ttctccgtgg	gcgagctgtg	gacggacctc	aacctcaacg	acgtcaaccc	gcaccgccag		720
ctcatcatga	actggatcaa	cgccaccggg	ggcaactcgg	gcgcgttcga	cttcaccacc		780
aaggggcatcc	tccagcaggc	ggtgcagtac	aacgagttct	ggcggctcaa	ggccagcgat		840
ggaaagcccc	agggcgccat	cggctggtgg	ccggccaggt	cggtgacgtt	catcgacaac		900
cacgacacgg	gcccgagcca	cccgagcggc	ggccagaacc	actggccctt	ccccagcgac		960
aagggtgatgc	agggctacgc	gtacatcctc	acgcaccggg	ggattccctg	tgtctactgg		1020
gtgcacttct	atgactgggg	ccacgcagcg	gccatcaaga	gcctcatcgc	ggcgcgcaag		1080
gagaagggcg	tcacctccac	gtccgcggtg	aacatcgtgg	ccgcggacac	gagcaagtac		1140
gccgccatca	tcaccggcaa	cacgggcagc	ctggcgatga	agattggtgt	cggcgccctgg		1200
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tag							1263

<210>	353
<211>	420
<212>	PRT
<213>	Unknown

<220>
<223> obtained from an environmental sample

<221> SIGNAL
<222> (1) . . . (27)

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Leu Gly Leu Val Ala Ser Thr Ala Ala Ser Ala Gly Pro Leu Asp Gly

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Asn	Ser	Ser	20	Val	Met	Leu	Gln	25	Gly	Phe	His	Trp	Arg	30	Ser	Thr	Glu
Thr	Tyr	Pro	35	Trp	Trp	Gly	Val	40	Ile	Gln	Gly	Lys	Ala	45	Ser	Asp	Ile
Ala	Ser	Gly	50	Phe	Thr	Met	Val	55	Trp	Phe	Pro	Pro	Ser	60	Gly	Asp	Ser
Ala	Asn	Glu	65	Gly	Tyr	Leu	Pro	70	Arg	Gln	Leu	Tyr	Val	75	Gln	Ser	80
Tyr	Gly	Thr	85	Asp	Ala	Gln	Leu	90	Ser	Ala	Ile	Gly	Ser	95	Leu	His	Thr
Tyr	Gly	Val	100	Lys	Ala	Ile	Ala	105	Ile	Val	Val	Asn	His	110	Arg	Val	Gly
Thr	Thr	Asn	115	Trp	Ala	Asp	Phe	120	Thr	Asn	Pro	Thr	Trp	125	Gly	Ser	Trp
Val	Val	Lys	130	Gly	Asp	Glu	Trp	135	Thr	Ser	Ala	Thr	Gly	140	Asn	Tyr	Asp
Gly	Asp	Gly	145	Phe	Ser	Ala	Ala	150	Arg	Asp	Leu	Asp	His	155	Thr	Asn	Gly
Val	Gln	Ser	160	Asp	Leu	Lys	Gly	165	Trp	Met	Asn	Trp	Leu	170	Lys	Ser	Ile
Gly	Tyr	Asp	175	Gly	Trp	Arg	Tyr	180	Asp	Tyr	Val	Lys	Gly	185	Tyr	Gly	Ser
Tyr	Val	Gly	190	Gly	Tyr	Asn	Ser	195	Ala	Thr	Val	Pro	Tyr	200	Phe	Ser	Val
Glu	Leu	Trp	205	Thr	Asp	Leu	Asn	210	Asp	Val	Asn	Pro	His	215	Arg	Gln	220
Leu	Ile	Met	225	Asn	Trp	Ile	Asn	230	Ala	Thr	Gly	Gly	Asn	235	Ser	Gly	Ala
Asp	Phe	Thr	240	Thr	Lys	Gly	Ile	245	Leu	Gln	Gln	Ala	Val	250	Gln	Tyr	Asn
Phe	Trp	Arg	255	Leu	Lys	Ala	Ser	260	Asp	Gly	Lys	Pro	Gln	265	Gly	Ala	Ile
Trp	Trp	Pro	270	Ala	Arg	Ser	Val	275	Thr	Phe	Ile	Asp	Asn	280	His	Asp	Thr
Pro	Ser	His	285	Pro	Ser	Gly	Gly	290	Gln	Asn	His	Trp	Pro	295	Phe	Pro	Ser
Lys	Val	Met	300	Gln	Gly	Tyr	Ala	305	Ile	Leu	Thr	His	Pro	310	Gly	Ile	Pro
Cys	Val	Tyr	315	Trp	Val	His	Phe	320	Asp	Trp	Gly	His	Ala	325	Ala	Ala	Ile
Lys	Ser	Leu	330	Ile	Ala	Ala	Arg	335	Glu	Lys	Gly	Val	Thr	340	Ser	Thr	Ser
Ala	Val	Asn	345	Ile	Val	Ala	Ala	350	Asp	Thr	Ser	Lys	Tyr	355	Ala	Ile	Ile
Thr	Gly	Asn	360	Thr	Gly	Ser	Leu	365	Met	Lys	Ile	Gly	Val	370	Gly	Ala	Trp
Ser	Pro	Gly	375	Thr	Gly	Trp	Thr	380	Val	Asn	Ser	Gly	Thr	385	Asn	Trp	Ala
Val	Trp	Lys	390	Gln	400	410	415										

<210> 354

<211> 2577

<212> DNA

<213> Unknown

<220>

<223> obtained from an environmental sample

<400> 354

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aacgagtgcg	agacagtgc	cggtcccaag	ggattcgggtg	gtgtccagat	cactccgccc	180
gcagagcaca	agcaggggag	ccaggtcttg	tggaccgtct	accagcccgt	cagcttcaag	240
aacttcaaca	gcttcggcgg	cagcgaggca	gagctcagga	gcatgatcac	ccgctgcaat	300
gcggccgggg	tcaaggtcta	tgccgacgcc	gtcttcaacc	agctggcctc	gggcagcggg	360
acagccaccg	gcggcggcag	ctacaactcg	gggcaatatc	agtatcccca	gttcggctac	420

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aacgactttcc accacagcgg tgacatcacc aactacggcg acagcaacaa cgtctggaac 480
ggcgccctct acggcatgcc ggatctcaat accggctctc cctacgtgca ggatcagata 540
gccacctaca tgaagaccct gctgggttgg ggcgtggcgg gctttcgcat cगतgcggcc 600
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tacctggaag tgataggcgc aggtggagag tcggcagaca tccagccggg tcgctacacc 720
tatatcgaca ctgtcaccga cttcaaatac ggcacagatc tggccgccaa cttcaacggc 780
cagatcaaga acctcaagac cctgggcgag agctggggcc tgtccctc cagcaaggcc 840
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accatcatga acatggccct gttccacaac agaaccgagg gtcaagctgt cagtacttgg 1140
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aacaacgaga gcggcagcct ggtggcctcg gtgcagaccg gcctgccagc gggcgagtac 1260
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gatgggggtg ccgaacagg gggcgccgac atagtgacc cggtcagcgg ccagtaccgg 2220
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aagaacatcg ccagctgaa catccgtggc accaccaatg gctggggcac cactccatg 2340
acgctgggtg gggatcacca gtggcaggcc ggtgtcacct tcacgggggc cggtgatgcc 2400
agcggcggcc agcgcttcaa gttcgacgtc aagggtgact ggaccagaa ctacggcgac 2460
accaacaagg acggcgtggc cgagctggcc ggagccgaca tcaccaccgc agtggtcgg 2520
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<210> 355
 <211> 858
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(24)

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Gln Trp Lys Phe Asn Asp Ile Ala Asn Glu Cys Glu Thr Val Leu Gly
35     40     45
Pro Lys Gly Phe Gly Gly Val Gln Ile Thr Pro Pro Ala Glu His Lys
50     55     60
Gln Gly Ser Gln Val Trp Trp Thr Val Tyr Gln Pro Val Ser Phe Lys
65     70     75     80
Asn Phe Asn Ser Phe Gly Gly Ser Glu Ala Glu Leu Arg Ser Met Ile
85     90     95
Thr Arg Cys Asn Ala Ala Gly Val Lys Val Tyr Ala Asp Ala Val Phe
100    105    110
Asn Gln Leu Ala Ser Gly Ser Gly Thr Ala Thr Gly Gly Gly Ser Tyr
115    120    125
Asn Ser Gly Gln Tyr Gln Tyr Pro Gln Phe Gly Tyr Asn Asp Phe His
130    135    140
His Ser Gly Asp Ile Thr Asn Tyr Gly Asp Ser Asn Asn Val Trp Asn

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145	Gly	Ala	Leu	Tyr	Gly	150	Met	Pro	Asp	Leu	Asn	155	Thr	Gly	Ser	Pro	Tyr	160	Val
	Gln	Asp	Gln	Ile	Ala	165	Thr	Tyr	Met	Lys	Thr	170	Leu	Leu	Gly	Trp	Gly	175	Val
	Ala	Gly	Phe	Arg	Ile	180	Asp	Ala	Ala	Lys	His	185	Met	Ala	Pro	Gly	Glu	190	Val
	Lys	Ala	Ile	Leu	Asp	195	Lys	Ala	Ala	Gly	Ser	200	Pro	Lys	Ala	Tyr	Leu	205	Val
	Ile	Gly	Ala	Gly	Gly	210	Glu	Ser	Ala	Asp	Ile	215	Gln	Pro	Gly	Arg	Tyr	220	Thr
225	Tyr	Ile	Asp	Thr	Val	230	Thr	Asp	Phe	Lys	Tyr	235	Gly	Thr	Asp	Leu	Ala	240	Ala
	Asn	Phe	Asn	Gly	Gln	245	Ile	Lys	Asn	Leu	Lys	250	Thr	Leu	Gly	Glu	Ser	255	Trp
	Gly	Leu	Leu	Pro	Ser	260	Ser	Lys	Ala	Phe	Val	265	Phe	Val	Val	Asn	His	270	Asp
	Arg	Glu	Arg	Gly	His	275	Gly	Gly	Gly	Gly	Met	280	Leu	Thr	Phe	Met	Ser	285	Gly
	Ala	Arg	Tyr	Asp	Leu	290	Ala	Asn	Thr	Phe	Met	295	Met	Ala	Trp	Pro	Tyr	300	Gly
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	Glu	Thr	Asp	Lys	Gly	325	Ala	Pro	Gly	Ser	Thr	330	Pro	Cys	Thr	Asp	Gly	335	Gln
	Trp	Asn	Cys	Glu	Gln	340	Arg	Arg	Pro	Thr	Ile	345	Met	Asn	Met	Ala	Leu	350	Phe
	His	Asn	Arg	Thr	Glu	355	Gly	Gln	Ala	Val	Ser	360	His	Trp	Trp	Asp	Asn	365	Gly
	Asn	Asn	Gln	Ile	Ala	370	Phe	Gly	Arg	Gly	Asp	375	Lys	Gly	Phe	Val	Ala	380	Ile
385	Asn	Asn	Glu	Ser	Gly	390	Ser	Leu	Val	Ala	Ser	395	Val	Gln	Thr	Gly	Leu	400	Pro
	Ala	Gly	Glu	Tyr	Cys	405	Asn	Leu	Leu	Gly	Gly	410	Asn	Asp	Tyr	Cys	Ser	415	Gly
	Gly	Tyr	Val	Thr	Val	420	Asp	Gly	Ser	Gly	Lys	425	Ala	Ser	Leu	Asn	Val	430	Pro
	Gly	Met	Lys	Ala	Ala	435	Ala	Ile	Ala	Gly	Cys	440	Gly	Lys	Ala	Asn	Pro	445	
	Cys	Gly	Gly	Thr	Ala	450	Leu	Pro	Gly	Asn	Lys	455	Phe	Ser	Ser	Met	Asn	460	Leu
465	Arg	Gly	Thr	His	Asn	470	Ala	Trp	Gly	Asn	Thr	475	Pro	Met	Thr	Val	Asp	480	Ala
	Asn	Arg	Val	Trp	Ser	485	Ala	Thr	Leu	Thr	Leu	490	Thr	Gly	Ser	Gly	Asp	495	Ala
	Thr	Gly	Ala	Gln	Arg	500	Phe	Lys	Phe	Asp	Val	505	Phe	Gly	Asn	Trp	Thr	510	Glu
	Ser	Tyr	Gly	Asp	Asn	515	Glu	Gly	Asp	Gly	Ile	520	Ala	Asp	Lys	Gly	Ser	525	Ser
	Lys	Asp	Ile	Leu	Val	530	Ser	Gly	Thr	Gly	Ser	535	Tyr	Arg	Ile	Thr	Leu	540	Asn
545	Glu	Gly	Asp	Leu	Arg	550	Tyr	Thr	Val	Thr	Pro	555	Leu	Thr	Ser	Asn	Gln	560	Ala
	Pro	Val	Ala	Ala	Leu	565	Ser	Pro	Lys	Val	Ser	570	Ser	Val	Lys	Thr	Gly	575	Glu
	Ser	Val	Val	Phe	Asp	580	Gly	Ser	Ala	Ser	Thr	585	Asp	Asp	Glu	Ala	Val	590	Ala
	Ser	Tyr	Ser	Trp	Ser	595	Thr	Gly	Gly	Asn	Ala	600	Pro	Thr	Glu	Thr	Val	605	Gln
	Phe	Asp	Thr	Pro	Gly	610	Thr	His	Thr	Val	Thr	615	Leu	Thr	Val	Thr	Asp	620	Ala
625	Glu	Gly	Leu	Thr	Ser	630	Ser	Ala	Ser	Ala	Thr	635	Val	Thr	Val	Thr	Asp	640	Ser
	Asn	Gly	Ala	Tyr	Asn	645	Ser	Val	Leu	Pro	Thr	650	Leu	His	Phe	Arg	Gly	655	Thr
	Pro	Asn	Gly	Trp	Gly	660	Thr	Leu	Ala	Met	Thr	665	Leu	Val	Ala	Asp	Asn	670	Gln
	Trp	Glu	Ala	Leu	Ala	675	Thr	Phe	Asn	Gly	Gln	680	Ala	Asn	Gln	Arg	Phe	685	Lys

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690	Phe	Asp	Val	Lys	Gly	Asp	Trp	Thr	Gln	Asn	Tyr	Gly	Asp	Thr	Asn	Lys
705	Asp	Gly	Val	Ala	Glu	Gln	Gly	Gly	Ala	Asp	Ile	Val	Thr	Pro	Val	Ser
	Gly	Gln	Tyr	Arg	Val	Arg	Phe	Asn	Asp	Gln	Thr	Leu	Gln	Tyr	Ser	Leu
	Thr	Pro	Val	Ser	Val	Gly	Tyr	Ala	Lys	Asn	Ile	Ala	Ser	Leu	Asn	Ile
	Arg	Gly	Thr	Thr	Asn	Gly	Trp	Gly	Thr	Thr	Pro	Met	Thr	Leu	Val	Gly
	Asp	His	Gln	Trp	Gln	Ala	Gly	Val	Thr	Phe	Thr	Gly	Ala	Gly	Asp	Ala
	Ser	Gly	Gly	Gln	Arg	Phe	Lys	Phe	Asp	Val	Lys	Gly	Asp	Trp	Thr	Gln
	Asn	Tyr	Gly	Asp	Thr	Asn	Lys	Asp	Gly	Val	Ala	Glu	Leu	Ala	Gly	Ala
	Asp	Ile	Thr	Thr	Ala	Val	Val	Gly	Ala	Tyr	Val	Val	Arg	Phe	Asn	Asp
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 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 356

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ctgattgagg	ctatccaggc	gcttaatgca	gcggatgtta	aagccatggc	tgatattgtt	180
atcaatcacc	gctgcggaca	gtatcagatt	aacggtaact	ggcacggata	tgaacagccg	240
gactggcagg	aatgggctgt	tgttgcta	gatggcggca	cgggtgctca	ggataccggt	300
gagggatacg	gtcctgcacc	ggacatagac	cacaccaacc	ctacagtcca	ggcggacatt	360
atggcctgga	tgaactggct	gcggactaca	ataggatttg	atggctggcg	ctatgattat	420
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<211> 1625

<212> PRT

<213> Unknown

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<223> Obtained from an environmental sample

<400> 357

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35     40     45
Asn Ala Ala Asp Val Lys Ala Met Ala Asp Ile Val Ile Asn His Arg
50     55     60
Cys Gly Gln Tyr Gln Ile Asn Gly Asn Trp His Gly Tyr Glu Gln Pro
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Asp Trp Gln Glu Trp Ala Val Val Ala Asn Asp Gly Gly Thr Gly Ala
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Gln Asp Thr Gly Glu Gly Tyr Gly Pro Ala Pro Asp Ile Asp His Thr

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Ser	Val	Gly	Glu	165	Trp	Pro	Asp	170	Ile	Thr	Gly	Ser	Tyr	175	Tyr	Ala	Thr
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Leu	Gln	Leu	195	Ala	Phe	Glu	Arg	215	Thr	Glu	Phe	Trp	Arg	220	Met	Gly	Val
Pro	Gly	Leu	210	Ile	Gly	Trp	Trp	230	Pro	Glu	Arg	Ser	Val	235	Thr	Phe	Ile
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Phe	Pro	Ala	240	Val	His	Val	Ala	265	Ala	Gly	Tyr	Ala	Tyr	270	Ile	Leu	Thr
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Glu	Asp	Ile	415	Pro	Ile	Thr	Glu	440	Ala	Gly	Arg	Tyr	Thr	445	Ile	Thr	Leu
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 35 40 45
 Glu Ile Lys Asp Leu Gly Tyr Lys Ala Val Leu Val Ala Pro Pro Leu
 50 55 60
 Lys Ser Asn Ala Ala Asn Cys Ala Trp Trp Gln Arg Tyr Gln Pro Gln
 65 70 75 80
 Asp Ile Arg Val Ile Asp His Cys Lys Gly Asn Lys Gln Ala Phe Val
 85 90 95
 Asn Met Ile Asn Ala Leu Asn Asp Ala Asn Pro Ala Arg Lys Val Asp
 100 105 110
 Val Tyr Ala Asp Ile Val Leu Asn His Met Ala Asn Glu Arg Ser Gly
 115 120 125
 Ala Thr Asp Phe Pro Gly Ser Ala Ala Val Asn Ser Tyr Gly Ser Asn
 130 135 140
 Ser Ser Tyr Trp Asn Asn Gln Gln Leu Phe Gly Asn Leu Thr Gln Gly
 145 150 155 160
 Ile Phe Ser Ala Gly Asp Phe Asn Pro Ala Asn Cys Ile Ser Asn Tyr
 165 170 175
 Asn Asp Val Trp Gln Val Gln Asn Tyr Arg Leu Cys Gly Gly Ala Gly
 180 185 190
 Asp Thr Gly Leu Pro Asp Leu Asn Pro Asn Ser Trp Val Val Gln Gln
 195 200 205
 Gln Arg Ser Tyr Leu Thr Ala Leu Lys Asn Leu Gly Val Lys Gly Phe
 210 215 220
 Arg Val Asp Ala Ala Lys His Met Thr Ile Trp His Ile Asn Glu Ile
 225 230 235 240
 Phe Thr Ser Ser Ile Lys Ser Gly Ser Tyr Leu Phe Gly Glu Ile Ile
 245 250 255
 Thr Ser Gly Gly Ala Gly Asn Asn Glu Tyr Asp Ser Phe Leu Ala Pro
 260 265 270

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Tyr Leu Asn Tyr Thr Asp His Lys Ala Tyr Asp Phe Pro Leu Phe Asn
275 280 285
Ser Ile Arg Asn Ala Phe Ser Phe Gly Gly Ser Leu Ser Gln Leu Val
290 295 300
Asn Pro Ala Ala Tyr Gly Gln Ala Leu Gln Asn Ser Arg Ala Val Thr
305 310 315
Phe Thr Ile Thr His Asp Ile Pro Thr Asn Asp Gly Phe Arg Tyr Leu
325 330 335
Ile Met Asp Pro Thr Asp Glu Tyr Leu Ala Tyr Val Met Gly
340 345 350
Arg Asp Gly Gly Lys Pro Leu Ile Phe Ser Asp Ser Thr Gly Thr Asp
355 360 365
Asn Asn Arg Trp Val Asn Ala Tyr Lys Ala Asp His Ile Ser Lys Met
370 375 380
Leu Asn Phe His Asn Arg Met Gln Gly Gln Gly Met Glu Met Leu Ala
385 390 395
Trp Asn Asp Cys Ala Ile Leu Phe Arg Arg Gly Gln Glu Gly Ile Val
405 410 415
Gly Ile Asn Lys Cys Ser Gly Ser Gln Ser Phe Ser Val Asn Thr His
420 425 430
Gly Arg Phe Tyr Trp Tyr Arg Asn Tyr Arg Asp Val Leu Thr Gly Gly
435 440 445
Asn Leu Val Tyr Ile Asn Gly Gly Asn Tyr His Phe Ser Ile Pro Ala
450 455 460
Arg Gly Ala Arg Met Trp Tyr Ala Asp
465 470

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<210> 360
 <211> 1938
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 360
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gaaagttata acatcttttg cgcacatgtt gtgacggaag atgagattca aggggtacga 120
tttacagtctt gggctcctca tgcgaaagca atgagtgttg ttggagattt taatgagtgg 180
gattatgagc aacataagat gctacaagtg acagaagaag ggattttggc cttattttata 240
ccgcatattg aagaaggaga aatatataaa tatgcgattg aaacgttggc tggtgacgctc 300
attttaaagg cagatccgta tgctatatat gcagaagtaa gaccgaatac ggcattctgta 360
gtttttgata taaaaggata tgaatggaat gataaaaact ggaatcgtaa gaaaaagaaa 420
aaaccgattt ataaagaagc gatgacagtt tatgaattac attttggttc ttggaaaaag 480
aaagaagatg gaacactata ttcttacagg gaaatggcag aagaactcat tccgtatgtg 540
gtggaacatc aatttacaca tattgaaatt atgccgcttg ttgagcatcc atatgatcgt 600
tcttggggat atcaaggaa gggatattat gcagcgacga gtagattttg tacaccgcat 660
gatttaattg attttgctga cgaatgtcat aaatatggaa tcggtgtcat tttagatttg 720
gtaccggggc atttttgtaa agatgctcac ggtttatatt tgtttgatgg cacaccgacc 780
tatgaatata aagataaaga tgtacaagaa aatctagtat ggggaactgt caattttgat 840
ttaggaaaga gagaggtag taatttctta atttcaaag cgttattttg gatgagatat 900
ttccatattg atggtttcag agtagatgca gttgcgaaca tgttgtagtg gaataaagaa 960
ggacaagagc aaagtaatga gtacgctgtg tcatTTTTTaa gggagttaaa tgaagcagtg 1020
tttgagagg atgaagattt tcttatgacg gcagaggatt caacagcttg gccacttgta 1080
acagaccaa cgtatgaagg tgggcttgga aatggaatat gggctggatg 1140
aatgacgtgc tgaaatatat ggagtgtgag ccagagtaca gaaaatatat tcatgagaaa 1200
atgacgttct cttactata tgcttactct gaaaacttca tattaccctt ttctcatgat 1260
gaagtctgtc atgggaaaaa gtcgttatta aataaaatgc caggtgatta ctgggataag 1320
tttgctcagc ttcgtttatt atatggatat ttctttactc acccaggaaa gaaattactt 1380
ttcatgggag gagaattcgg gcagtttgat gagtggaaag acctgaaga tttagatttg 1440
aatttcatag attttgaaat gcatcgttat atgcatgatt actttaaaga gctcatagca 1500
ttgtataagc gctcaaacc actttggcag cttgatcatt cgcctgaagg ttttcagtgg 1560
attgatgcta ataataacga gcaaagtatt ttctcgttta tccgccaagg ggataaacia 1620
gaagatgctg tagttatcgt atgtaatttt acgaaagcta catatgaaa ctataaagta 1680
ggtgtaccag atttcgagta ttataacgag attttaaaca gtgatgcaca gcaatatggc 1740
ggttcgggac aagtgaataa gaaacgtctt aagacgattc tagaaccgta ccataatcaa 1800
gcagcacatg tagagattac aattccacca tttggcgatt ccatattacg accagtgaa 1860
acgagaaagg ggagcaaaaa acaagatggc tcaaaaacia aagtgcgtag caatgttact 1920

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1938

agcaggggga aaaggtag

<210> 361
<211> 645
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 361
Leu Ser Val Ile Asn Cys Glu Glu Val Lys Arg Asp Glu Phe His Thr
1 5 10 15
Glu Lys Tyr Tyr Glu Ser Tyr Asn Ile Phe Gly Ala His Val Thr
20 25 30
Glu Asp Glu Ile Gln Gly Val Arg Phe Thr Val Trp Ala Pro His Ala
35 40 45
Lys Ala Met Ser Val Val Gly Asp Phe Asn Glu Trp Asp Tyr Glu Gln
50 55 60
His Lys Met Leu Gln Val Thr Glu Glu Gly Ile Trp Ser Leu Phe Ile
65 70 75 80
Pro His Ile Glu Glu Gly Glu Ile Tyr Lys Tyr Ala Ile Glu Thr Leu
85 90 95
Ala Gly Asp Val Ile Leu Lys Ala Asp Pro Tyr Ala Ile Tyr Ala Glu
100 105 110
Val Arg Pro Asn Thr Ala Ser Val Phe Asp Ile Lys Gly Tyr Glu
115 120 125
Trp Asn Asp Lys Asn Trp Asn Arg Lys Lys Lys Lys Pro Ile Tyr
130 135 140
Lys Glu Ala Met Thr Val Tyr Glu Leu His Phe Gly Ser Trp Lys Lys
145 150 155 160
Lys Glu Asp Gly Thr Leu Tyr Ser Tyr Arg Glu Met Ala Glu Glu Leu
165 170 175
Ile Pro Tyr Val Val Glu His Gln Phe Thr His Ile Glu Ile Met Pro
180 185 190
Leu Val Glu His Pro Tyr Asp Arg Ser Trp Gly Tyr Gln Gly Thr Gly
195 200 205
Tyr Tyr Ala Ala Thr Ser Arg Phe Gly Thr Pro His Asp Leu Met Tyr
210 215 220
phe Val Asp Glu Cys His Lys Tyr Gly Ile Gly Val Ile Leu Asp Trp
225 230 235 240
Val Pro Gly His Phe Cys Lys Asp Ala His Gly Leu Tyr Leu Phe Asp
245 250 255
Gly Thr Pro Thr Tyr Glu Tyr Lys Asp Lys Asp Val Gln Glu Asn Leu
260 265 270
Val Trp Gly Thr Val Asn Phe Asp Leu Gly Lys Arg Glu Val Arg Asn
275 280 285
Phe Leu Ile Ser Asn Ala Leu Phe Trp Met Arg Tyr Phe His Ile Asp
290 295 300
Gly Phe Arg Val Asp Ala Val Ala Asn Met Leu Tyr Trp Asn Lys Glu
305 310 315 320
Gly Gln Glu Gln Ser Asn Glu Tyr Ala Val Ser Phe Leu Arg Glu Leu
325 330 335
Asn Glu Ala Val Phe Ala Glu Asp Glu Asp Phe Leu Met Thr Ala Glu
340 345 350
Asp Ser Thr Ala Trp Pro Leu Val Thr Ala Pro Thr Tyr Glu Gly Gly
355 360 365
Leu Gly Phe Asn Tyr Lys Trp Asn Met Gly Trp Met Asn Asp Val Leu
370 375 380
Lys Tyr Met Glu Cys Ala Pro Glu Tyr Arg Lys Tyr Ile His Glu Lys
385 390 395 400
Met Thr Phe Ser Leu Leu Tyr Ala Tyr Ser Glu Asn Phe Ile Leu Pro
405 410 415
Leu Ser His Asp Glu Val Val His Gly Lys Lys Ser Leu Leu Asn Lys
420 425 430
Met Pro Gly Asp Tyr Trp Asp Lys Phe Ala Gln Leu Arg Leu Leu Tyr
435 440 445
Gly Tyr Phe Phe Thr His Pro Gly Lys Lys Leu Leu Phe Met Gly Gly

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450 455 460
 Glu Phe Gly Gln Phe Asp Glu Trp Lys Asp Leu Asp Leu Asp Trp
 465 470 475 480
 Asn Leu His Asp Phe Glu Met His Arg Tyr Met His Asp Tyr Phe Lys
 485 490 495
 Glu Leu Ile Ala Leu Tyr Lys Arg Ser Lys Pro Leu Trp Gln Leu Asp
 500 505 510
 His Ser Pro Glu Gly Phe Gln Trp Ile Asp Ala Asn Asn Asn Glu Gln
 515 520 525
 Ser Ile Phe Ser Phe Ile Arg Gln Gly Asp Lys Gln Glu Asp Ala Leu
 530 535 540
 Val Ile Val Cys Asn Phe Thr Lys Ala Thr Tyr Glu Asn Tyr Lys Val
 545 550 555 560
 Gly Val Pro Asp Phe Glu Tyr Tyr Asn Glu Ile Leu Asn Ser Asp Ala
 565 570 575
 Gln Gln Tyr Gly Gly Ser Gly Gln Val Asn Lys Lys Arg Leu Lys Thr
 580 585 590
 Ile Leu Glu Pro Tyr His Asn Gln Ala Ala His Val Glu Ile Thr Ile
 595 600 605
 Pro Pro Phe Gly Val Ser Ile Leu Arg Pro Val Lys Thr Arg Lys Gly
 610 615 620
 Ser Lys Lys Gln Asp Gly Ser Lys Thr Lys Val Arg Ser Asn Val Thr
 625 630 635 640
 Ser Arg Gly Lys Arg
 645

<210> 362
 <211> 2094
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<400> 362
 atgaaaaaaa ttacttttctt attggcatta ttgctttttgg taagcgcaac aatattttgccc 362
 aacgacccca aagacgacat aatgctgcaa tcattttgggt gggacgaata cgaccagccc 120
 aaaataaaagg ctgctggtaa cttctacaaa tttgttgagt cgaaagctag cgagtggaaa 180
 gctgctgggt tcgatatgat atggatgcca ccccttagtc gctccacagg tgggtgtgggt 240
 tatttgccca ccgagttaca caacttcaac tcaacatggg gcagcgaggc cgacttacgt 300
 gctgtacgg ctgcactcaa agcacatggc atgtttccta ttgccgatgt ggttgccaat 360
 catcgcaatg gcaccacggc atggaccgac ttacgaacc ccgatggag ctgtgcaacc 420
 atcacatcta ccgacgaggc taatgtaaat tgggttgcat cgctggccc aaagccttgt 480
 ggcgaccgg atacaggcga cgattttaat ggcgggcgcg acctcgacca taccaatatc 540
 gaaacacgca acggcataaa aacttttctc aaaaaactaa aagaacaagg cttcgaaggc 600
 tggcgttggg acatgaccaa aggttttagt gccagttatg taggcgagta taacactgca 660
 agccaaccct atttctcggg aggcgaatat tgggacggaa actcgaacac cctcaaaaac 720
 tgggtagatg gcacaggcaa aaagtcggca acattcgatt tttcgcaata ttattcgatg 780
 gagaatgcat tcaaaaataa tgcattggagt gcattaggtt cggaagcag catggccggt 840
 ttggcgggtg tttttggcta ctccgactat gccgttactt ttgtcgacaa ccacgatact 900
 tttgtgcatg gcagcgcgcc actcggcgat aatattatga aagcctatgc ctacatactt 960
 acgaccccg gcataccgag cgtattttatt gcgcattatt acggtggaac atatcgtaaa 1020
 gatggtgtta ctgcactta cacaagccac aaagataaaa tcgaccccat aatggcagtg 1080
 cgcaaagcca atcgtattga cgcttgagc acattacaag ttgcaacaaa ttcgggcatg 1140
 tatgccgctt atatcaaacg ccgccacgac gatgccgctg cttcggtagc cgtaaaaatt 1200
 ggcccaggaa gttggtcgcc cgaaggtagt ggctggattt tagctgcatc gggaacagac 1260
 tacgccgtgt ggtcgaaagc gcaaatagtt gtagcaccta ccatgctcat tggtagtgga 1320
 acttacgaat taggacaaac actcagcatt acagctccgt cgggctatag catacgctac 1380
 accaccgatg gtagcgagcc aacagcggcg tcgacacttt acactgcgcc cattacattg 1440
 ccacaaggca caaccgttat caaagctgct tcgtttgtaa atggtgttat gtcggccgta 1500
 gtaaccaaca cttacactgc acaagcaaaa gcaaccagca taaaagtgcg tttcaaagct 1560
 cccgcttcgt ggacagcaag caaagcctat gtgtgggaaa atcgtaatgg cacaacacc 1620
 aaccttgtag gcgcatggcc aggaacaaca gctacaaaag acgctgatgg attttatgtg 1680
 tacaccataa caaatcacac acaatttagg gtaaatatta ttttcaacaa tgccagtggc 1740
 acaaccgcca gcgagcaaac cgaagacctt tcggtaacgg ccgacatatg ttggcaggca 1800
 gtaggcagca acaaatatgg tgtagaaaca gtgaactgcc caggcaccaa tgtcgaatat 1860
 aattacatac gtttattcaa catctatcca aatccgggca aagaacagtg gcgtataaac 1920
 accaacgaaa cattgcgcgc agtaaccatg tacgatagca tgggacgcat gcaccgcatg 1980

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caattaaatg ccgacaatag tttggacata agcaatatgg ccgccggact gtatcagatg 2040
caaattgaaa cagaaagcgg acgaaaagaa actcataaat ttgtaaaatt gtag 2094

<210> 363
<211> 697
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(20)

<400> 363
Met Lys Lys Ile Thr Phe Leu Leu Ala Leu Leu Leu Leu Val Ser Ala
1 5 10 15
Thr Ile Phe Ala Asn Asp Pro Lys Asp Asp Ile Met Leu Gln Ser Phe
20 25 30
Gly Trp Asp Glu Tyr Asp Gln Pro Lys Ile Lys Ala Ala Gly Asn Phe
35 40 45
Tyr Lys Phe Val Glu Ser Lys Ala Ser Glu Trp Lys Ala Ala Gly Phe
50 55 60
Asp Met Ile Trp Met Pro Pro Pro Ser Arg Ser Thr Gly Gly Val Gly
65 70 75 80
Tyr Leu Pro Thr Glu Leu His Asn Phe Asn Ser Thr Trp Gly Ser Glu
85 90 95
Ala Asp Leu Arg Ala Ala Thr Ala Ala Leu Lys Ala His Gly Met Phe
100 105 110
Pro Ile Ala Asp Val Val Ala Asn His Arg Asn Gly Thr Thr Ala Trp
115 120 125
Thr Asp Phe Thr Asn Pro Ala Trp Ser Cys Ala Thr Ile Thr Ser Thr
130 135 140
Asp Glu Ala Asn Val Asn Trp Val Ala Ser Arg Gly Pro Lys Pro Cys
145 150 155 160
Gly Ala Pro Asp Thr Gly Asp Asp Phe Asn Gly Gly Arg Asp Leu Asp
165 170 175
His Thr Asn Ile Glu Thr Arg Asn Gly Ile Lys Thr Phe Leu Thr Lys
180 185 190
Leu Lys Glu Gln Gly Phe Glu Gly Trp Arg Trp Asp Met Thr Lys Gly
195 200 205
Phe Ser Ala Ser Tyr Val Gly Glu Tyr Asn Thr Ala Ser Gln Pro Tyr
210 215 220
Phe Ser Val Gly Glu Tyr Trp Asp Gly Asn Ser Asn Thr Leu Lys Asn
225 230 235 240
Trp Val Asp Gly Thr Gly Lys Lys Ser Ala Thr Phe Asp Phe Ser Gln
245 250 255
Tyr Tyr Ser Met Glu Asn Ala Phe Lys Asn Asn Ala Trp Ser Ala Leu
260 265 270
Gly Ser Gly Ser Ser Met Ala Gly Leu Ala Gly Val Phe Gly Tyr Ser
275 280 285
Asp Tyr Ala Val Thr Phe Val Asp Asn His Asp Thr Phe Val His Gly
290 295 300
Ser Ala Pro Leu Gly Asp Asn Ile Met Lys Ala Tyr Ala Tyr Ile Leu
305 310 315 320
Thr His Pro Gly Ile Pro Ser Val Phe Ile Ala His Tyr Tyr Gly Gly
325 330 335
Thr Tyr Arg Lys Asp Gly Val Thr Arg Thr Tyr Thr Ser His Lys Asp
340 345 350
Lys Ile Asp Pro Ile Met Ala Val Arg Lys Ala Asn Arg Ile Asp Ala
355 360 365
Trp Ser Thr Leu Gln Val Ala Thr Asn Ser Gly Met Tyr Ala Ala Tyr
370 375 380
Ile Lys Arg Arg His Asp Asp Ala Ala Ala Ser Val Ala Val Lys Ile
385 390 395 400
Gly Pro Gly Ser Trp Ser Pro Glu Gly Ser Gly Trp Ile Leu Ala Ala
405 410 415
Ser Gly Thr Asp Tyr Ala Val Trp Ser Lys Ala Gln Ile Val Val Ala

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Pro	Thr	Met	420	Leu	Ile	Gly	Ser	Gly	425	Thr	Tyr	Glu	Leu	Gly	430	Gln	Thr	Leu
Ser	Ile	Thr	435	Ala	Pro	Ser	Gly	Tyr	440	Ser	Ile	Arg	Tyr	Thr	445	Thr	Asp	Gly
Ser	Glu	Pro	450	Thr	Ala	Ala	Ser	Thr	455	Leu	Tyr	Thr	Ala	Pro	460	Ile	Thr	Leu
465	Pro	Gln	Gly	Thr	Thr	Val	Ile	Lys	470	Ala	Ala	Ser	Phe	Val	475	Asn	Gly	480
Met	Ser	Ala	485	Val	Val	Thr	Asn	Thr	490	Tyr	Thr	Ala	Gln	Ala	495	Lys	Ala	Thr
Ser	Ile	Lys	500	Val	Arg	Phe	Lys	Ala	505	Pro	Ala	Ser	Trp	Thr	510	Ala	Ser	Lys
Ala	Tyr	Val	515	Trp	Glu	Asn	Arg	Asn	520	Gly	Thr	Asn	Thr	Asn	525	Leu	Ala	Gly
Ala	Trp	Pro	530	Gly	Thr	Thr	Ala	Thr	535	Lys	Asp	Ala	Asp	Gly	540	Phe	Tyr	Val
545	Tyr	Thr	Ile	Thr	Asn	His	Thr	Gln	550	Phe	Arg	Val	Asn	Ile	555	Ile	Phe	Asn
Asn	Ala	Ser	565	Gly	Thr	Thr	Ala	Ser	570	Glu	Gln	Thr	Glu	Asp	575	Leu	Ser	Val
Thr	Ala	Asp	580	Ile	Cys	Trp	Gln	Ala	585	Val	Gly	Ser	Asn	Lys	590	Tyr	Gly	Val
Glu	Thr	Val	595	Asn	Cys	Pro	Gly	Thr	600	Asn	Val	Glu	Asn	Asn	605	Tyr	Ile	Arg
Leu	Phe	Asn	610	Ile	Tyr	Pro	Asn	Pro	615	Gly	Lys	Glu	Thr	Val	620	Arg	Ile	Asn
625	Thr	Asn	Glu	Thr	Leu	Arg	Ala	Val	630	Thr	Met	Tyr	Asp	Ser	635	Met	Gly	Arg
Met	His	Arg	645	Met	Gln	Leu	Asn	Ala	650	Asp	Asn	Ser	Leu	Asp	655	Ile	Ser	Asn
Met	Ala	Ala	660	Gly	Leu	Tyr	Gln	Met	665	Gln	Ile	Glu	Thr	Glu	670	Ser	Gly	Arg
Lys	Glu	Thr	675	His	Lys	Phe	Val	Lys	680	Lys	Leu				685			
690									695									

<210> 364
 <211> 1536
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 364					
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agttttaatg	ctgctatagt	tcaagcagct	accaatggca	caatgtttca	atactttgaa
tggtacttac	caaatgacgg	aacgctatgg	gataaattag	gtaatgatgc	tccaaactta
aagagtatag	gtgtaacagc	tgtttgata	ccaccagcat	ataagggag	ttcacaggct
gatgttggat	atggtaccta	tgatttatat	gacctagggtg	aatttaatat	aaaagggtaca
gtgagaacaa	agtatggaac	aaagagccaa	ttaatcagtg	cagtagataa	actacataat
aatggaatac	aagtatatgg	agatgtttgta	ttaaatacata	agatgggtgc	tgatggaact
gaaacagttt	gggctgttga	agtaaatacca	agcaatagaa	atcaagaaac	atcaggagat
tatcaaattt	cagcatggac	aaaatttgat	tttccaggaa	gaggaaacac	atactcaagc
ttcaaatgga	gatggtatca	ttttgatgga	ggtgattggg	accaaagcag	aagcttaagt
agaatatata	aattaagagg	agactataaa	ggatgggact	gggaagtga	tactgaaaat
ggaaattatg	actattttaa	gggtgcggat	gtagacatgc	aacaccctga	agttgtaaat
gaattaaata	actggggtaa	gtggtttacc	caaacagtaa	aattagatgg	atatagattg
gatgcagtaa	agcatattaa	gtttgatttc	ttaatacat	ggcttgatta	tcaaagacaa
caatcaggaa	aggaactatt	tacagttagt	gaatactggt	cagatgattt	gggaaaactt
caaaattata	taagtaaaac	aggaagaaga	atgtctctat	ttgatgttcc	actacaaaga
aatctctata	atgcttcaaa	tggtgggtgga	tattatgata	tgagaaatat	aaagagtgga
acattaataa	gtgttgaccc	aactaaagca	gttacatttg	ttgacaacca	tgatacagaa
ccaggacaat	cactacaaag	atgtgttcaa	gattggttta	agccacttgc	atatacattt
attttaacaa	gagaagaagg	atatccatgt	atattctatg	gtgattatta	tggaattcca
gcaaagggag	tatcagcaaa	gaaggtagca	attgataaga	taatgcttgc	aagaaagaat
tatgcttatg	tgactattta	gaccattggg	atataatagg	ttggacaaga	ttggacactga
gaaggagata	atgaacatgc	taattcaggt	atggcagcac	ttataactga	tggaactggt

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ggttcaaaat ggatgtatgt aggacaaaaa ttcataaggaa agacattcta cgatattaca 1440
ggaaacagaa cagatacagt tggtataaat tcatacaggtt ggggtgaatt taaagtaa 1500
ggtggttcac actcaatctg ggttccaaag aattaa 1536

<210> 365
<211> 511
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(29)

<400> 365
Met Lys Asn Phe Lys Lys Lys Leu Ser Ala Leu Leu Leu Ser Phe Ile
1 5 10 15
Leu Ile Phe Ser Phe Asn Ala Ala Ile Val Gln Ala Ala Thr Asn
20 25 30
Gly Thr Met Phe Gln Tyr Phe Glu Trp Tyr Leu Pro Asn Asp Gly Thr
35 40 45
Leu Trp Asp Lys Leu Gly Asn Asp Ala Pro Asn Leu Lys Ser Ile Gly
50 55 60
Val Thr Ala Val Trp Ile Pro Pro Ala Tyr Lys Gly Ser Ser Gln Ala
65 70 75 80
Asp Val Gly Tyr Gly Thr Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn
85 90 95
Gln Lys Gly Thr Val Arg Thr Lys Tyr Gly Thr Lys Ser Gln Leu Ile
100 105 110
Ser Ala Val Asp Lys Leu His Asn Asn Gly Ile Gln Val Tyr Gly Asp
115 120 125
Val Val Leu Asn His Lys Met Gly Ala Asp Gly Thr Glu Thr Val Trp
130 135 140
Ala Val Glu Val Asn Pro Ser Asn Arg Asn Gln Glu Thr Ser Gly Asp
145 150 155 160
Tyr Gln Ile Ser Ala Trp Thr Lys Phe Asp Phe Pro Gly Arg Gly Asn
165 170 175
Thr Tyr Ser Ser Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Val Asp
180 185 190
Trp Asp Gln Ser Arg Ser Leu Ser Arg Ile Tyr Lys Leu Arg Gly Asp
195 200 205
Tyr Lys Gly Trp Asp Trp Glu Val Asp Thr Glu Asn Gly Asn Tyr Asp
210 215 220
Tyr Leu Met Gly Ala Asp Val Asp Met Gln His Pro Glu Val Val Asn
225 230 235 240
Glu Leu Asn Asn Trp Gly Lys Trp Phe Thr Gln Thr Val Lys Leu Asp
245 250 255
Gly Tyr Arg Leu Asp Ala Val Lys His Ile Lys Phe Asp Phe Asn
260 265 270
Thr Trp Leu Asp Tyr Gln Arg Gln Gln Ser Gly Lys Glu Leu Phe Thr
275 280 285
Val Gly Glu Tyr Trp Ser Asp Asp Leu Gly Lys Leu Gln Asn Tyr Ile
290 295 300
Ser Lys Thr Gly Arg Arg Met Ser Leu Phe Asp Val Pro Leu Gln Arg
305 310 315 320
Asn Phe Tyr Asn Ala Ser Asn Gly Gly Gly Tyr Tyr Asp Met Arg Asn
325 330 335
Ile Lys Ser Gly Thr Leu Ile Ser Val Asp Pro Thr Lys Ala Val Thr
340 345 350
Phe Val Asp Asn His Asp Thr Glu Pro Gly Gln Ser Leu Gln Arg Phe
355 360 365
Val Gln Asp Trp Phe Lys Pro Leu Ala Tyr Thr Phe Ile Leu Thr Arg
370 375 380
Glu Glu Gly Tyr Pro Cys Ile Phe Tyr Gly Asp Tyr Tyr Gly Ile Pro
385 390 395 400
Ala Lys Gly Val Ser Ala Lys Lys Val Ala Ile Asp Lys Ile Met Leu
405 410 415

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Ala Arg Lys Asn Tyr Ala Tyr Gly Lys Gln Asn Asp Tyr Leu Asp His
420 425 430
Trp Asp Ile Gly Trp Thr Arg Gly Asp Asn Glu His Ala Asn
435 440 445
Ser Gly Met Ala Ala Leu Ile Thr Asp Gly Pro Gly Gly Ser Lys Trp
450 455 460
Met Tyr Val Gly Gln Lys Phe Ile Gly Lys Thr Phe Tyr Asp Ile Thr
465 470 475 480
Gly Asn Arg Thr Asp Thr Val Val Ile Asn Ser Ser Gly Trp Gly Glu
485 490 495
Phe Lys Val Asn Gly Gly Ser Tyr Ser Ile Trp Val Pro Lys Asn
500 505 510

<210> 366
<211> 1992
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 366
atgaaaatct ctagttttta ctgcgcgaca cttggcgag gggtattact ttccctccgca 60
caggtaagcg cagcgccaac gacctttgtg cattttattcg attggaattg gcaagatatc 120
gctaccgagt gtgaaaccta tttaggtcca caaggttacg ctgccgtgca ggtttcgcct 180
ccaaatgaac atattgaagt gtctgagtgg tgggcgcgat atcagcctgt aagctacgtt 240
ctgcaaagcc gcagtggcag ccgcagtgag tttatcaata tggttgagcg ctgtgatgca 300
gtcgggtgtag atattttacgt cgatgctgtc attaaccaca tggccgctgg aagcgggtact 360
ggaactgcag gcaatacatt tggcaataag tcttatccta tctactctcc acaggacttt 420
catgctacat gcgctattaa tgactacggc gatcgctggc aagtacagaa ctgcgagtta 480
gtcgggttgg ccgacttaaa taccgccgat aattatgtac agacgacgct ggccggatat 540
ttaaagtatc ttgtgaacat tgggtttgcc ggtttccgtc tggacgcgtc aaaacacatg 600
gctgcaaccg atattgaagg tatcttacag agagtaaattg gcgatccgct tatttttcag 660
gaagtgattg atcaaggcgg cgaagcggta agtgcttcag agtatttttg gaacggcctg 720
gtaaccgaat ttaagtattc tgtagaaata ggaaatacat ttcgcaacgg ctcgctggca 780
tggctgagta acttcggcga agcgtgggga ttcatgccta gctatcaggc agtcgtgttt 840
gttgataatc acgacaatca gcgtgggtcat ggtggagggt gaaatgtcat cacccttgag 900
gatggcgagc tgtatgattt agccaatgta tttatgctgg cgtggccgta cggctaccct 960
aaagtaattg ccagctttga ctttcatggc gatacagatg cgggacctcc tggcgttccg 1020
gtccataacg acggcaacct cgaatgcttt ggcagtaact ggaagtgtga acaccgctgg 1080
agctacattg caggcggtgt aaactttaga aatagcaccg cggacaattt cactgtcacc 1140
gactgggtgga gcaacggtgc caaccaaact gccttcggcc ggcttgacag tggctttgtt 1200
gtcattaata aagaatctta tgaccttaac accactctcg aaacgtcaat ggcaccggga 1260
gagtattgtg atgttctcac aggcgagctc aatgcacatg gcgatggctg taccggcaat 1320
aagctttcgg tagatagcaa tgggcgcctt tctgtctccg tcccatcctg ggaagcgatt 1380
gccattcacc ataagtccaa agtggcagcc acatcaaatt cggcagactg gcaacgaacg 1440
atgatattca tcgaagccga aactgccagt ggtcaagata tgtttgttcg gggcgacatt 1500
gaccatcaag ccgcagcgaa ccaactcggc atcacctgta cagatacaaa ttacgcttgt 1560
gcagtgcga ttgtgcataa taacttgccg aatacgacta cggcttcctg gaaggcaaat 1620
gacaactatc ttgactggta tgggtgtgaa gactcccaga gtactgacgc agaaggcagt 1680
gcgctggact ggacaaccaa ctcttgcccc gcagaatggg gaaccttaaa aaccgtgatt 1740
gacgatgggt tcggtgaaac gccctataat acctggggct cgcattttctg gatgctcgac 1800
gtacaaatgg attgcagtaa gacagtgaat ggggtggttcg aacttaaagc ctatgtcaaa 1860
aacgggcagg gctgggaagg cgatatcagt cagagcggta caccctaccc atcaaccaac 1920
cacatggcgc aatgcggtaa aaagaatatg ttccggtttg gacaaagcac ggtgcaaatc 1980
agcgatctgt aa 1992

<210> 367
<211> 663
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(24)

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<400> 367
Met Lys Ile Ser Ser Phe Asn Cys Ala Thr Leu Gly Ala Gly Leu Leu
1 5 10 15
Leu Ser Ser Ala Gln Val Ser Ala Ala Pro Thr Thr Phe Val His Leu
20 25 30
Phe Asp Trp Asn Trp Gln Asp Ile Ala Thr Glu Cys Glu Thr Tyr Leu
35 40 45
Gly Pro Gln Gly Tyr Ala Ala Val Gln Val Ser Pro Pro Asn Glu His
50 55 60
Ile Glu Val Ser Glu Trp Trp Ala Arg Tyr Gln Pro Val Ser Tyr Val
65 70 75 80
Leu Gln Ser Arg Ser Gly Ser Arg Ser Glu Phe Ile Asn Met Val Glu
85 90 95
Arg Cys Asp Ala Val Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn
100 105 110
His Met Ala Ala Gly Ser Gly Thr Gly Thr Ala Gly Asn Thr Phe Gly
115 120 125
Asn Lys Ser Tyr Pro Ile Tyr Ser Pro Gln Asp Phe His Ala Thr Cys
130 135 140
Ala Ile Asn Asp Tyr Gly Asp Arg Trp Gln Val Gln Asn Cys Glu Leu
145 150 155 160
Val Gly Leu Ala Asp Leu Asn Thr Ala Asp Asn Tyr Val Gln Thr Thr
165 170 175
Leu Ala Gly Tyr Leu Asn Asp Leu Val Asn Ile Gly Val Ala Gly Phe
180 185 190
Arg Leu Asp Ala Ser Lys His Met Ala Ala Thr Asp Ile Glu Gly Ile
195 200 205
Leu Gln Arg Val Asn Gly Asp Pro Leu Ile Phe Gln Glu Val Ile Asp
210 215 220
Gln Gly Gly Glu Ala Val Ser Ala Ser Glu Tyr Phe Gly Asn Gly Leu
225 230 235 240
Val Thr Glu Phe Lys Tyr Ser Val Glu Ile Gly Asn Thr Phe Arg Asn
245 250 255
Gly Ser Leu Ala Trp Leu Ser Asn Phe Gly Glu Ala Trp Gly Phe Met
260 265 270
Pro Ser Tyr Gln Ala Val Val Phe Val Asp Asn His Asp Asn Gln Arg
275 280 285
Gly His Gly Gly Gly Gly Asn Val Ile Thr Phe Glu Asp Gly Arg Leu
290 295 300
Tyr Asp Leu Ala Asn Val Phe Met Leu Ala Trp Pro Tyr Gly Tyr Pro
305 310 315 320
Lys Val Met Ser Ser Phe Asp Phe His Gly Asp Thr Asp Ala Gly Pro
325 330 335
Pro Gly Val Pro Val His Asn Asp Gly Asn Leu Glu Cys Phe Gly Ser
340 345 350
Asn Trp Lys Cys Glu His Arg Trp Ser Tyr Ile Ala Gly Gly Val Asn
355 360 365
Phe Arg Asn Ser Thr Ala Asp Asn Phe Thr Val Thr Asp Trp Trp Ser
370 375 380
Asn Gly Ala Asn Gln Ile Ala Phe Gly Arg Ser Asp Ser Gly Phe Val
385 390 395 400
Val Ile Asn Lys Glu Ser Tyr Asp Leu Asn Thr Thr Leu Glu Thr Ser
405 410 415
Met Ala Pro Gly Glu Tyr Cys Asp Val Leu Thr Gly Glu Leu Asn Ala
420 425 430
Ser Gly Asp Gly Cys Thr Gly Asn Lys Leu Ser Val Asp Ser Asn Gly
435 440 445
Arg Leu Ser Val Ser Val Pro Ser Trp Glu Ala Ile Ala Ile His His
450 455 460
Asn Ala Lys Val Ala Ala Thr Ser Asn Ser Ala Asp Trp Gln Arg Thr
465 470 475 480
Met Ile Phe Ile Glu Ala Glu Thr Ala Ser Gly Gln Asp Met Phe Val
485 490 495
Arg Gly Asp Ile Asp His Gln Ala Ala Asn Gln Leu Gly Ile Thr
500 505 510
Cys Thr Asp Thr Asn Tyr Ala Cys Ala Val Pro Ile Val His Asn Asn
515 520 525
Leu Arg Asn Thr Thr Thr Ala Ser Trp Lys Ala Asn Asp Asn Tyr Leu

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530      535      540
Asp Trp Tyr Gly Val Glu Asp Ser Gln Ser Thr Asp Ala Glu Gly Ser
545      550      555      560
Ala Leu Asp Trp Thr Thr Asn Ser Trp Pro Ala Glu Trp Gly Thr Leu
565      570      575
Lys Thr Val Ile Asp Asp Gly Phe Gly Glu Thr Pro Tyr Asn Thr Trp
580      585      590
Gly Ser His Phe Trp Met Leu Asp Val Gln Met Asp Cys Ser Lys Thr
595      600      605
Val Asn Gly Trp Phe Glu Leu Lys Ala Tyr Val Lys Asn Gly Gln Gly
610      615      620
Trp Glu Gly Asp Ile Ser Gln Ser Gly Thr Pro Tyr Pro Ser Thr Asn
625      630      635      640
His Met Ala Gln Cys Gly Lys Lys Asn Met Phe Arg Phe Gly Gln Ser
645      650      655
Thr Val Gln Ile Ser Asp Leu
660

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<210> 368
 <211> 1257
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 368
atgtttatattt acaatctttt ccctaggcctt tatgggccat ttagcaagtg gatagacgac      60
ttagatagaa taaaggagtt gggagtagac tggatatata taaaccccgt atcttatccc      120
gggttttccg gcagcctata ctcaataaaa gatcattaca aattcaaccc tctgttccta      180
gactctaatt caaaaaaatc accagaggaa caattcagag atttcattaa agcgtgtaaa      240
tcaagaggaa taaaagtaat cgttgatttg gtagtcaatc acacagctat agattccgta      300
cttgtagaca aacatcctaa ctggtacaaa agagacgaaa acgggcaaat aaagagacca      360
ggcgcatggg ataatagcgt atgggttgaa tggggagatt tagcagaact agacaacgag      420
aacagtcctg ataaaagagc attatggaac tattggaaag acctagtaga gtggcatata      480
gacttgggag ttgatgggtt tagatgcgac tatgcttata atgtccctgt agaattatgg      540
gacttcctaa taaagaacgc aaaatccaag aaagaagatg ttatattcct agcagagata      600
ctcggaggtc ctttgatag aaacctagaa gttgcaaaag ctggccttga ttatgtattt      660
agcagtgcta aatgggtggaa tttctcagac gcttggttca tagaacaac aagagagttc      720
tcaaaatatg ttaaagcaat agcatttcct gaatctcatg acacaccaag gttgaattcg      780
gagtacgaag aagatttagc aaaaataaag caaagattca tatttacagc attcataaat      840
gaaggattaa tgatacctgt aggatttgag tttggcttca ttaacaagct agatgtggta      900
aaaacaaccc cagactgggt ggaaatacca tcttatgaca tttctaagtt cataaaacac      960
atagttgaaa tcaaaaaaaaa gcatcccata ttgtcatatg acgctacatc agtagatatg      1020
ctagaaaaca cccctgacat ctcaatgttt gtcaagagga acaagctaac agaccagatt      1080
gcccttttcg tagttaacaa aaacacaacc tcaaaagtaa agagaataat agatttcctt      1140
agtgtattcg gttattatga tgctatagat gtaatagata acgttgtaat gcctaagtac      1200
gctgaagtaa cagtaccgag agcagggtata aaagttttcg taaacagaat acactaa      1257

```

<210> 369
 <211> 418
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 369
Met Phe Ile Tyr Asn Leu Phe Pro Arg Leu Tyr Gly Pro Phe Ser Lys
1      5      10      15
Trp Ile Asp Asp Leu Asp Arg Ile Lys Glu Leu Gly Val Asp Trp Ile
20      25      30
Tyr Ile Asn Pro Val Ser Tyr Pro Gly Phe Ser Gly Ser Leu Tyr Ser
35      40      45
Ile Lys Asp His Tyr Lys Phe Asn Pro Leu Phe Leu Asp Ser Asn Ser
50      55      60
Lys Lys Ser Pro Glu Glu Gln Phe Arg Asp Phe Ile Lys Ala Cys Lys
65      70      75      80

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Ser Arg Gly Ile Lys Val Ile Val Asp Leu Val Val Asn His Thr Ala
 85 90 95
 Ile Asp Ser Val Leu Val Asp Lys His Pro Asn Trp Tyr Lys Arg Asp
 100 105 110
 Glu Asn Gly Gln Ile Lys Arg Pro Gly Ala Trp Asp Asn Ser Val Trp
 115 120 125
 Val Glu Trp Gly Asp Leu Ala Glu Leu Asp Asn Glu Asn Ser Pro Asp
 130 135 140
 Lys Arg Ala Leu Trp Asn Tyr Trp Lys Asp Leu Val Glu Trp His Ile
 145 150 155 160
 Asp Leu Gly Val Asp Gly Phe Arg Cys Asp Tyr Ala Tyr Asn Val Pro
 165 170 175
 Val Glu Leu Trp Asp Phe Leu Ile Lys Asn Ala Lys Ser Lys Lys Glu
 180 185 190
 Asp Val Ile Phe Leu Ala Glu Ile Leu Gly Gly Pro Phe Asp Arg Asn
 195 200 205
 Leu Glu Val Ala Lys Ala Gly Phe Asp Tyr Val Phe Ser Ser Ala Lys
 210 215 220
 Trp Trp Asn Phe Ser Asp Ala Trp Phe Ile Glu Gln Thr Arg Glu Phe
 225 230 235 240
 Ser Lys Tyr Val Lys Ala Ile Ala Phe Pro Glu Ser His Asp Thr Pro
 245 250 255
 Arg Leu Asn Ser Glu Tyr Glu Glu Asp Leu Ala Lys Ile Lys Gln Arg
 260 265 270
 Phe Ile Phe Thr Ala Phe Ile Asn Glu Gly Leu Met Ile Pro Val Gly
 275 280 285
 Phe Glu Phe Gly Phe Ile Asn Lys Leu Asp Val Val Lys Thr Thr Pro
 290 295 300
 Asp Trp Trp Glu Ile Pro Ser Tyr Asp Ile Ser Lys Phe Ile Lys His
 305 310 315 320
 Ile Val Glu Ile Lys Lys His Pro Ile Leu Ser Tyr Asp Ala Thr
 325 330 335
 Ser Val Asp Met Leu Glu Asn Thr Pro Asp Ile Ser Met Phe Val Lys
 340 345 350
 Arg Asn Lys Leu Thr Asp Gln Ile Ala Leu Phe Val Val Asn Lys Asn
 355 360 365
 Thr Thr Ser Lys Val Lys Arg Ile Ile Asp Phe Pro Ser Val Phe Gly
 370 375 380
 Tyr Tyr Asp Ala Ile Asp Val Ile Asp Asn Val Val Met Pro Lys Tyr
 385 390 395 400
 Ala Glu Val Thr Val Pro Arg Ala Gly Ile Lys Val Phe Val Asn Arg
 405 410 415
 Ile His

<210> 370
 <211> 1614
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 370
 atgaaacgct acttaacact caccctactg cttaccctgt tgatacttgc gccgctggct 60
 ttggcacaaag agtccgatga gggcgcagca tgggctaacg agactgtttt ttatgagatt 120
 tttgtgcgta gcttttacga tagcgacggg gacggtaaag gggatattcg cggcgttatc 180
 gaaaaactcg attacctcaa cgacggcgac ccgaatacca caaccgactt aggcattacc 240
 ggcattttggt tgatgcccgt catggagtcg cccagttatc acggctacga cgcgaccgat 300
 tactacatgc tcgataaaga ctacggcaca aacgacgact tcaagctgtt catggaagaa 360
 gcgcacaaac gcggaatccg tgtcatcggt gacttgatgc tcaatcacac ctcggtgcag 420
 catgagtgggt tcattcgaatc ggggtgacccg caaagcgaaa aagccgattg gtacatctgg 480
 gaagatgaga acccgaatta tcgcggcccc gataatcagc aggtgtggca cggcctacgc 540
 gggcgctatt actacggcgt gttctgggtca gggatgccgg acttgaactt caccaatccc 600
 gatgtcaccg cgcaaattga cgatgtcacg cgcttttggc tagaagagat gaacgtggac 660
 ggcttccgcg ttgacgcgat taaacacatc atcgaggatg gcaccattca agagaacacc 720
 ccgcgcacgc gccaatggct ggccgattac cgcgcgttcg tcaagtccat caagcctgac 780
 gcgctgttgg tcggtgaggt ctggtcgccg acgttcgcta tcgcgccgta tgtgggcaca 840

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```

gcggttgaca tcgcgtttga gttcgacttg gccgaggcgc tgatacgcgg cgcgagcttt 900
ggcacgcccc acctcgtcac gcgacagcta gagacggtct ttgagagtta cccgccaac 960
gctttcgcca cattcttgac taaccacgat cagaaccgct tcatgacca aattcgcggg 1020
gacgcggcag cggccaaaaa cggcgcacgc ttgctactca cgttgcccg ggtaccgttc 1080
atttactacg gtgaagaaat cggcatgacc ggcataaaac ctgatgagga catccgaaa 1140
ccgatgcagt gggacgatac ccccaagacc gctggtttca ccagtgggta gcgcgtgtgg 1200
cgtgctgtga ataacgatgc attcgagggc gtcaatgtgg ccgcacaaac cgaaaacccc 1260
gactcacttc taagccacta ccgcgcgctg gttcaagcac ggggtgagcag ccccgcgtaa 1320
caaagcggcg acatgatttt ggttgaatcg tctaaccgta agctgtttac tttcttgcg 1380
caaagcgagg cccaaacggg gctgggtgat attaacctg acgaccgtcc cgtgaccgag 1440
tacacgctca cgctggaaag tggcacactc gcggaaggcg tgacagcatc ggcgctcctc 1500
gcatcagggt aggtaaacgc gccacaggtc aacgggcaag gcggctttgc ggattacacc 1560
ccgcttgacg aactcgcgcc caattcggtc acgctgattc agcttgggga ataa 1614

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<210> 371
 <211> 537
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(22)

<400> 371

Met	Lys	Arg	Tyr	Leu	Thr	Leu	Thr	Leu	Leu	Leu	Thr	Leu	Leu	Ile	Leu
1				5				10						15	
Ala	Pro	Leu	Ala	Leu	Ala	Gln	Glu	Ser	Asp	Glu	Gly	Ala	Ala	Trp	Ala
			20					25					30		
Asn	Glu	Thr	Val	Phe	Tyr	Glu	Ile	Phe	Val	Arg	Ser	Phe	Tyr	Asp	Ser
		35					40					45			
Asp	Gly	Asp	Gly	Lys	Gly	Asp	Ile	Arg	Gly	Val	Ile	Glu	Lys	Leu	Asp
	50					55					60				
Tyr	Leu	Asn	Asp	Gly	Asp	Pro	Asn	Thr	Thr	Thr	Asp	Leu	Gly	Ile	Thr
65					70					75				80	
Gly	Ile	Trp	Leu	Met	Pro	Val	Met	Glu	Ser	Pro	Ser	Tyr	His	Gly	Tyr
			85						90					95	
Asp	Ala	Thr	Asp	Tyr	Tyr	Met	Leu	Asp	Lys	Asp	Tyr	Gly	Thr	Asn	Asp
			100					105					110		
Asp	Phe	Lys	Leu	Phe	Met	Glu	Glu	Ala	His	Lys	Arg	Gly	Ile	Arg	Val
		115					120					125			
Ile	Val	Asp	Leu	Met	Leu	Asn	His	Thr	Ser	Val	Gln	His	Glu	Trp	Phe
	130					135					140				
Ile	Glu	Ser	Gly	Asp	Pro	Gln	Ser	Glu	Lys	Ala	Asp	Trp	Tyr	Ile	Trp
145					150					155				160	
Glu	Asp	Glu	Asn	Pro	Asn	Tyr	Arg	Gly	Pro	Asp	Asn	Gln	Gln	Val	Trp
			165						170					175	
His	Gly	Leu	Arg	Gly	Arg	Tyr	Tyr	Tyr	Gly	Val	Phe	Trp	Ser	Gly	Met
		180						185					190		
Pro	Asp	Leu	Asn	Phe	Thr	Asn	Pro	Asp	Val	Thr	Ala	Gln	Met	Tyr	Asp
		195					200					205			
Val	Thr	Arg	Phe	Trp	Leu	Glu	Glu	Met	Asn	Val	Asp	Gly	Phe	Arg	Leu
	210					215					220				
Asp	Ala	Ile	Lys	His	Ile	Glu	Asp	Gly	Thr	Ile	Gln	Glu	Asn	Thr	
225					230				235					240	
Pro	Ala	Thr	Arg	Gln	Trp	Leu	Ala	Asp	Tyr	Arg	Ala	Phe	Val	Lys	Ser
			245						250					255	
Ile	Lys	Pro	Asp	Ala	Leu	Leu	Val	Gly	Glu	Val	Trp	Ser	Pro	Thr	Phe
			260					265					270		
Ala	Ile	Ala	Pro	Tyr	Val	Gly	Thr	Ala	Val	Asp	Ile	Ala	Phe	Glu	Phe
		275					280					285			
Asp	Leu	Ala	Glu	Ala	Leu	Ile	Arg	Gly	Ala	Ser	Phe	Gly	Thr	Pro	Asn
	290					295					300				
Leu	Val	Thr	Arg	Gln	Leu	Glu	Thr	Val	Phe	Glu	Ser	Tyr	Pro	Pro	Asn
305					310					315					320
Ala	Phe	Ala	Thr	Phe	Leu	Thr	Asn	His	Asp	Gln	Asn	Arg	Phe	Met	Thr
				325					330					335	

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Gln Ile Arg Gly Asp Ala Ala Ala Ala Lys Asn Gly Ala Ser Leu Leu
 340 345 350
 Leu Thr Leu Pro Gly Val Pro Phe Ile Tyr Tyr Gly Glu Glu Ile Gly
 355 360 365
 Met Thr Gly Met Lys Pro Asp Glu Asp Ile Arg Lys Pro Met Gln Trp
 370 375 380
 Asp Asp Thr Pro Lys Thr Ala Gly Phe Thr Ser Gly Glu Arg Val Trp
 385 390 395 400
 Arg Ala Val Asn Asn Asp Ala Phe Glu Gly Val Asn Val Ala Ala Gln
 405 410 415
 Thr Glu Asn Pro Asp Ser Leu Leu Ser His Tyr Arg Ala Leu Val Gln
 420 425 430
 Ala Arg Val Ser Ser Pro Ala Leu Gln Ser Gly Asp Met Ile Leu Val
 435 440 445
 Glu Ser Ser Asn Arg Lys Leu Phe Thr Phe Leu Arg Gln Ser Glu Ala
 450 455 460
 Gln Thr Val Leu Val Met Ile Asn Leu Asp Asp Arg Pro Val Thr Glu
 465 470 475 480
 Tyr Thr Leu Thr Leu Glu Ser Gly Thr Leu Ala Glu Gly Val Thr Ala
 485 490 495
 Ser Ala Leu Leu Ala Ser Gly Glu Val Asn Ala Pro Gln Val Asn Gly
 500 505 510
 Gln Gly Gly Phe Ala Asp Tyr Thr Pro Leu Ala Glu Leu Ala Pro Asn
 515 520 525
 Ser Val Thr Leu Ile Gln Leu Gly Glu
 530 535

<210> 372
 <211> 1437
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 372
 atgttcaaaa tagtgccgct cctttctctc gtcacggcct tcgtcatcgg cgcatacccg 60
 ctgcgcgcgc agtccgctat agaggacctg gtaaaggcct ctccggtccg ggaccaggcc 120
 gtaccggccc cctccgagcc caaatccgcc cacacgccca gcgccctggc gcgccaggac 180
 tggaaaacgg ccgtaaccct ctattcggcc aagaacttca atcctccgcc cgcgcggcgg 240
 ctgaacctcg aggcctcgcc ggtgggtcatg ctgcagggct tccactggta cgccgacaat 300
 tactggctca acctgccccg cggctgggtg ggcaggctgg ccgagaaagc cccggaggtc 360
 ggcgccggcg gcttcgacct ggtctggttc cccccggtat cccgcggcag ctattacccc 420
 accgagtggg acgacctga ttcccagtgg ggcaagaaag acgtcctgat agaggccgtg 480
 cgggcatga agggccacgg ggtcagcccc gtggccgaca tcatactcaa ccaccgcaac 540
 ggcaccctcg actgggcgga tttcaccagc cccgactggc cgaccacggc gatagtcagg 600
 gacgacgagt ggcagggcat gaaaggcccc aattacgacg agggacaggg agagggcggc 660
 tgccgcgacc tggatcacia caacgccctg gtgcgcgagg acgccaaggt cttcctgcgc 720
 tggctgcgct acaccgtcgg tttcgaaggc tggcgctacg acatggtcaa gggcttcctg 780
 cccacacagga tcaaggacta caaccgcgct tcggcgccgg ccttctcggg aggggaatac 840
 tacgacacca accgccagct gctgggtcaac tgggtggacg gcacggccga cggcgacaag 900
 tcggaagcct ccacggtctt cgatttcacc acccgctaca acctggtggc cgccgtagag 960
 agcgagcgtt acgacctgct cagcgaccac ggccgcccc tccggtccat cggctggtgg 1020
 ccggccaagt cggtcacctt cgtggagaac cagcacacct cgccccgcga ccccaagttc 1080
 ctgcagaacg ccccgcgcga gtaccgcgtc cagcgccaga tgggctacgc ctacatcctt 1140
 accatccccg gcacgcccag cgtcttcttg cccatttct tcgactgggg ttccgaatac 1200
 agggccaagc tgcaggagct cgtcaatatc cgcaagtcgg ccggcataac cgccgtcagc 1260
 ccggtgcaga tactcgcggc ctacaacgag ctctacgccg cgatgataac cggcagcgat 1320
 aagtacgtcg tattgaagct cggcaagaac tggggctggc agcccgcca cgactggaag 1380
 ctggaagcca gcggcgagcg ctgggcccgtc tggaccagc ccgtcaccag gccctga 1437

<210> 373
 <211> 478
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(25)

<400> 373

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Met Phe Lys Ile Val Pro Leu Leu Ser Leu Val Thr Ala Phe Val Ile
1      5      10      15
Gly Ala Tyr Pro Leu Arg Ala Gln Ser Ala Ile Glu Asp Leu Val Lys
20      25      30
Ala Ser Pro Val Arg Asp Gln Ala Val Pro Ala Pro Ser Glu Pro Lys
35      40      45
Ser Ala His Thr Pro Ser Ala Leu Ala Arg Gln Asp Trp Lys Thr Ala
50      55      60
Val Thr Leu Tyr Ser Ala Lys Asn Phe Asn Pro Pro Ala Arg Arg
65      70      75      80
Leu Asn Leu Glu Ala Ser Pro Val Val Met Leu Gln Gly Phe His Trp
85      90      95
Tyr Ala Asp Asn Tyr Trp Leu Asn Leu Pro Arg Gly Trp Trp Gly Arg
100     105
Leu Ala Glu Lys Ala Pro Glu Val Gly Ala Ala Gly Phe Asp Leu Val
115
Trp phe Pro Pro Val Ser Arg Gly Ser Tyr Tyr Pro Thr Glu Trp Tyr
120     125     130     135     140
Asp Leu Asp Ser Gln Trp Gly Lys Lys Asp Val Leu Ile Glu Ala Val
145     150     155
Arg Ala Met Lys Ala His Gly Val Ser Pro Val Ala Asp Ile Ile Leu
160     165     170     175
Asn His Arg Asn Gly Thr Leu Asp Trp Ala Asp Phe Thr Ser Pro Asp
180     185     190
Trp Pro Thr Thr Ala Ile Val Arg Asp Asp Glu Trp Gln Gly Met Lys
195     200     205
Gly Pro Asn Tyr Asp Glu Gly Gln Gly Glu Gly Cys Arg Asp Leu
210     215     220
Asp His Asn Asn Ala Leu Val Arg Glu Asp Ala Lys Val Phe Leu Arg
225     230     235     240
Trp Leu Arg Tyr Thr Val Gly Phe Glu Gly Trp Arg Tyr Asp Met Val
245     250     255
Lys Gly Phe Leu Pro His Arg Ile Lys Asp Tyr Asn Arg Ala Ser Ala
260     265     270
Pro Ala phe Ser Val Gly Glu Tyr Tyr Asp Thr Asn Arg Gln Leu Leu
275     280     285
Val Asn Trp Val Asp Gly Thr Ala Asp Gly Asp Lys Ser Glu Ala Ser
290     295     300
Thr Val Phe Asp Phe Thr Thr Arg Tyr Asn Leu Val Ala Ala Val Glu
305     310     315     320
Ser Glu Arg Tyr Asp Leu Leu Ser Asp His Gly Arg Pro Ser Gly Ser
325     330     335
Ile Gly Trp Trp Pro Ala Lys Ser Val Thr Phe Val Glu Asn His Asp
340     345     350
Thr Ser Pro Arg Asp Pro Lys Phe Leu Gln Asn Ala Pro Arg Glu Tyr
355     360     365
Arg Val Gln Arg Gln Met Gly Tyr Ala Tyr Ile Leu Thr His Pro Gly
370     375     380
Thr Pro Ser Val Phe Trp Pro His Phe Phe Asp Trp Gly Ser Glu Tyr
385     390     395     400
Arg Ala Lys Leu Gln Glu Leu Val Asn Ile Arg Lys Ser Ala Gly Ile
405     410     415
Thr Ala Val Ser Pro Val Gln Ile Leu Ala Ala Tyr Asn Glu Leu Tyr
420     425     430
Ala Ala Met Ile Thr Gly Ser Asp Lys Tyr Val Val Leu Lys Leu Gly
435     440     445
Lys Asn Trp Gly Trp Gln Pro Gly His Asp Trp Lys Leu Glu Ala Ser
450     455     460
Gly Glu Arg Trp Ala Val Trp Thr Gln Pro Val Thr Arg Pro
465     470     475

```

<210> 374

<211> 1398

<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 374
atgaaacaca cagcgggaat atgggcgatc gcaggcatgc tgatcgcccc cttggcgcac 60
gccgacgtca tactgcacgc cttcaactgg aagtacagtg aagtgaccgc caaagccgat 120
cttatcaagt cagccgggta caaacaggta ctcatctcac cacctctcaa gtcatacaggc 180
aacgagtggg gggctcggtt ccagccgcag gatctgcgtc ttgtcgactc cctctttggc 240
aacaagcagg atctcgagca gctgatcgcc gccatgaagg cgcgaggtat cgccgtctat 300
gccgatgtgg tgctcaacca catggccaac gaaagctgga aacgcagcga tctcaactat 360
cccggcaacg agctgctgca aagctacgcc agcaaccggg cctatttcga gcgccagaaa 420
ctgtttggcg atctggggca aaacgtcctc tccggtcagg attttcatcc cgaaggggtgc 480
atcagcgact ggagcgatcc gggatcatgtc cagtattggc gactctgtgg tgggtgccggg 540
gacaaggggc tgccggatct ggatcccaac aactgggtgg tcagccagca acaagcctat 600
ctcaaggcgc tcaaggggat ggggatcaag ggttttcggg tcgatgcggg caaacacatg 660
agcgattacc agatcaatgc tgtgttcacc cccgagatca aacaggggat gcacgtgttt 720
ggcgaggtag tcaccacagg aggggcaggc agcaccgatt acgagcgctt cctcaaacc 780
tacctcgaca gcagcgcca gggagcctat gactttccgc tgtttgctc cctgcgtggg 840
gcgttaagtt acggtggcag catgaacctg ctggcggatc cgggcgccta tggtcaggcg 900
ctaccgggca atcgcgccgt cacctttgcc gtgaccacg acatccccac caacgacggc 960
ttccgctatc agatcctgag ccagactgac gagaagctgg cctatgccta cctgctcggc 1020
cgcgatgggt gctcaccact ggtctattcg gatcacggcg agaccaggc caaggatgga 1080
ctgcgtggc aggattacta ccagcgcgcc gatctcaagg ggatgatccg tttccacaat 1140
gcggtgcagg gccagcccat gcagctggc ggcagtggcg actgcttcgt gctgtttcaa 1200
cggggcaaac aggggctggg cggcatcaac aagtgcgact atgagcagga gtactggctc 1260
gataccgcca gggtcgagat gaactgggat cgcaactatc gggacgtgct cgaccaaggg 1320
gccgtagtca acgtgcagag ccagtgggta cgggtcgcca ttcctgcccg cagcgccaga 1380
atgtggctgc aggagtga 1398

<210> 375
<211> 465
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(21)

<400> 375
Met Lys His Thr Ala Gly Ile Trp Ala Ile Ala Gly Met Leu Ile Ala
1 5 10 15
Pro Leu Ala His Ala Asp Val Ile Leu His Ala Phe Asn Trp Lys Tyr
20 25 30
Ser Glu Val Thr Ala Lys Ala Asp Leu Ile Lys Ser Ala Gly Tyr Lys
35 40 45
Gln Val Leu Ile Ser Pro Pro Leu Lys Ser Ser Gly Asn Glu Trp Trp
50 55 60
Ala Arg Tyr Gln Pro Gln Asp Leu Arg Leu Val Asp Ser Pro Leu Gly
65 70 75 80
Asn Lys Gln Asp Leu Glu Gln Leu Ile Ala Ala Met Lys Ala Arg Gly
85 90 95
Ile Ala Val Tyr Ala Asp Val Val Leu Asn His Met Ala Asn Glu Ser
100 105 110
Trp Lys Arg Ser Asp Leu Asn Tyr Pro Gly Asn Glu Leu Leu Gln Ser
115 120 125
Tyr Ala Ser Asn Pro Ala Tyr Phe Glu Arg Gln Lys Leu Phe Gly Asp
130 135 140
Leu Gly Gln Asn Val Leu Ser Gly Gln Asp Phe His Pro Glu Gly Cys
145 150 155 160
Ile Ser Asp Trp Ser Asp Pro Gly His Val Gln Tyr Trp Arg Leu Cys
165 170 175
Gly Gly Ala Gly Asp Lys Gly Leu Pro Asp Leu Asp Pro Asn Asn Trp
180 185 190

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Val Val Ser Gln Gln Gln Ala Tyr Leu Lys Ala Leu Lys Gly Met Gly
 195 200 205
 Ile Lys Gly Phe Arg Val Asp Ala Val Lys His Met Ser Asp Tyr Gln
 210 215 220
 Ile Asn Ala Val Phe Thr Pro Glu Ile Lys Gln Gly Met His Val Phe
 225 230 235 240
 Gly Glu Val Ile Thr Thr Gly Gly Ala Gly Ser Thr Asp Tyr Glu Arg
 245 250 255
 Phe Leu Lys Pro Tyr Leu Asp Ser Ser Gly Gln Gly Ala Tyr Asp Phe
 260 265 270
 Pro Leu Phe Ala Ser Leu Arg Gly Ala Leu Ser Tyr Gly Gly Ser Met
 275 280 285
 Asn Leu Leu Ala Asp Pro Gly Ala Tyr Gly Gln Ala Leu Pro Gly Asn
 290 295 300
 Arg Ala Val Thr Phe Ala Val Thr His Asp Ile Pro Thr Asn Asp Gly
 305 310 315 320
 Phe Arg Tyr Gln Ile Leu Ser Gln Thr Asp Glu Lys Leu Ala Tyr Ala
 325 330 335
 Tyr Leu Leu Gly Arg Asp Gly Gly Ser Pro Leu Val Tyr Ser Asp His
 340 345 350
 Gly Glu Thr Gln Ala Lys Asp Gly Leu Arg Trp Gln Asp Tyr Tyr Gln
 355 360 365
 Arg Ala Asp Leu Lys Gly Met Ile Arg Phe His Asn Ala Val Gln Gly
 370 375 380
 Gln Pro Met Gln Leu Val Gly Ser Gly Asp Cys Phe Val Leu Phe Gln
 385 390 395 400
 Arg Gly Lys Gln Gly Leu Val Gly Ile Asn Lys Cys Asp Tyr Glu Gln
 405 410 415
 Glu Tyr Trp Leu Asp Thr Ala Arg Phe Glu Met Asn Trp Tyr Arg Asn
 420 425 430
 Tyr Arg Asp Val Leu Asp Gln Gly Ala Val Val Asn Val Gln Ser Gln
 435 440 445
 Trp Val Arg Val Ala Ile Pro Ala Arg Ser Ala Arg Met Trp Leu Gln
 450 455 460
 Glu
 465

<210> 376
 <211> 1551
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<400> 376
 atgtcacaat tcctaagggt cgcagtcagg cgcataatgta ctcagtgggtg gtgccgtacc 60
 actgggatta ttcttctggt ggttttcgcg ggcgtgtcgg cggcgttggg ccaggccggt 120
 ttgcagatg accgtgtcat gctgcaggga ttttattggg aatcctatcg gcatggacac 180
 ctggagaagt tccccaatta cggcgataaa cactgggtacg tgattgtaca aaacctcgca 240
 ccggaaatcc gagccggccg cttcaatctg atctggctgc cgcgcacctc gtacgcgggc 300
 ggcttttagcg ccggctataa tccaaaggag tacttttcgcc ttgataacag ctacggcacg 360
 ttcgatgagc accgcgccgc gcttgaagcg ctccctcgga acggaatcga gccagtggct 420
 gatctagtca tcaatcacgg cgacggtaac caggcgtggg cggatttcaa aaatcccgat 480
 tggggaactt gggcgatctg ccgaacggac gaggcattca caaatgtgaa ctccggcatc 540
 accaatacgc ggaacgatca acgcggtaac tgtgaggaag aagccgatta tcggcccagag 600
 ggcacgtata attattcgga ttttcgcgac attgcgcagt ctgatccgcg ggtgcggcgc 660
 gacattgaac gatattctgct ccagttaaaa tctttgggtt accgggggtg gcgttacgat 720
 atgggtgcacg gctataacgc gagatgggtt gcgctctata atcgactcac gcaaccact 780
 ttctcggtag gagagtacga ttgggacaag cagcgcgacg agcgtggctg gatattggga 840
 accgcaacga acccgactcc aacgggtgca gatcacctca gaacttcgag cagcgtcttt 900
 gatttcacga gccagttcag tctgaaatca atcaacagcg gaaactacac ggcactgtat 960
 ggcttcggaa acggcatcgg tctgggtggc gatacaacag atgggctgcc gtggaagaat 1020
 cgcgccgtga cgtttctcga aaaccacgat accgggtacc gcaccaacga ggacggcacg 1080
 ccgcagcagg gccaccaatt cgattcgttt gcgaacaact ggcaggtcga gcaggggtat 1140
 gcgcagattt tgacacatcc cgggtgtgcc actgtctact ggaagcatta ctttgactgg 1200
 ggatcggacc tgcagaataa gatcataatg ctcggaaggt agcgggggtt 1260
 aacgcgggca gtaaggtaga cccacaggac aatgccaaat actcgggcgt atacgcggct 1320

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agaatcagcg gacgtaatgg cgagttgtac gtgcgcatag gagggagcga ccagcaatgg 1380
cagcctagtg cttcgggata tgccgattac cgcgagtacg cgcaaggagc cggctggaaa 1440
gtctgggtta aattacctgg aaatcccccg gtgcagcagg tgcctcttcc cagcgcgttg 1500
cctgttccgc agtaccggcc tccggaccaaa atcgaagcccc caccgcaatg a 1551

<210> 377
<211> 516
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(37)

<400> 377
Met Ser Gln Phe Leu Arg Phe Ala Val Arg Arg Ile Cys Thr Gln Trp
1 5 10 15
Trp Cys Arg Thr Gly Ile Ile Leu Val Val Phe Ala Gly Val
20 25 30
Ser Ala Ala Leu Gly Gln Ala Gly Phe Asp Asp Asp Arg Val Met Leu
35 40 45
Gln Gly Phe Tyr Trp Glu Ser Tyr Arg His Gly His Leu Glu Lys Phe
50 55 60
Pro Asn Tyr Gly Asp Lys His Trp Tyr Val Ile Val Gln Asn Leu Ala
65 70 75 80
Pro Glu Ile Arg Ala Gly Arg Phe Asn Leu Ile Trp Leu Pro Pro Pro
85 90 95
Ser Tyr Ala Gly Gly Phe Ser Ala Gly Tyr Asn Pro Lys Glu Tyr Phe
100 105 110
Arg Leu Asp Asn Ser Tyr Gly Thr Phe Asp Glu His Arg Ala Ala Leu
115 120 125
Glu Ala Leu Leu Arg Asn Gly Ile Glu Pro Val Ala Asp Leu Val Ile
130 135 140
Asn His Arg Asp Gly Asn Gln Ala Trp Ala Asp Phe Lys Asn Pro Asp
145 150 155 160
Trp Gly Thr Trp Ala Ile Cys Arg Thr Asp Glu Ala Phe Thr Asn Val
165 170 175
Asn Ser Gly Ile Thr Asn Thr Pro Asn Asp Gln Arg Gly Asn Cys Glu
180 185 190
Glu Glu Ala Asp Tyr Arg Pro Glu Gly Thr Tyr Asn Tyr Ser Asp Phe
195 200 205
Arg Asp Ile Ala His Ala Asp Pro Arg Val Arg Arg Asp Ile Glu Arg
210 215 220
Tyr Leu Leu Gln Leu Lys Ser Leu Gly Tyr Arg Gly Trp Arg Tyr Asp
225 230 235 240
Met Val His Gly Tyr Asn Ala Arg Trp Val Ala Leu Tyr Asn Arg Leu
245 250 255
Thr Gln Pro Thr Phe Ser Val Gly Glu Tyr Asp Trp Asp Lys His Ala
260 265 270
Gln Gln Arg Gly Trp Ile Trp Gly Thr Ala Thr Asn Pro Thr Pro Thr
275 280 285
Gly Ala Asp His Leu Arg Thr Ser Ser Ser Val Phe Asp Phe Thr Ser
290 295 300
Gln Phe Ser Leu Lys Ser Ile Asn Ser Gly Asn Tyr Thr Ala Leu Tyr
305 310 315 320
Gly Phe Gly Asn Gly Ile Gly Leu Val Gly Asp Thr Thr Asp Gly Leu
325 330 335
Pro Trp Lys Asn Arg Ala Val Thr Phe Leu Glu Asn His Asp Thr Gly
340 345 350
Tyr Arg Thr Asn Glu Asp Gly Thr Pro Gln Gln Gly His Gln Phe Asp
355 360 365
Ser phe Ala Asn Asn Trp Gln Val Glu Gln Gly Tyr Ala Gln Ile Leu
370 375 380
Thr His Pro Gly Val Pro Thr Val Tyr Trp Lys His Tyr Phe Asp Trp
385 390 395 400
Gly Ser Asp Leu Gln Asn Lys Ile Arg Ala Leu Ile Asn Ala Arg Lys

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Val Ala Gly Val 405 Asn Ala Gly Ser Lys 410 Val Asp Pro Gln Asp 415 Asn Ala
420 425 430
Lys Tyr Ser Gly Val Tyr Ala Ala Arg Ile Ser Gly Arg Asn Gly Glu
435 440 445
Leu Tyr Val Arg Ile Gly Gly Ser Asp Gln Gln Trp Gln Pro Ser Ala
450 455 460
Ser Gly Tyr Ala Asp Tyr Arg Glu Tyr Ala Gln Gly Ala Gly Trp Lys
465 470 475 480
Val Trp Val Lys Leu Pro Gly Asn Pro Pro Val Gln Gln Val Pro Leu
485 490 495
Pro Ser Ala Leu Pro Val Pro Gln Tyr Arg Pro Pro Asp Gln Ile Glu
500 505 510
Pro Pro Pro Glu
515

```

<210> 378
 <211> 1269
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

```

<400> 378
atgttgaagc agttcacgaa ggcctgatac accctgacga gcctgctggc gctcgctctc 60
gtcgcaccgt tggccagcgc gggcccgcgt gatggcaaca gcagcgacgt catgttgtag 120
ggcttccatt ggtactcgta ccagtcgttt ccgtgggtggg gcgtcatcaa gaacaacgcg 180
gcgagcatca aggccgacgg cttcaccatg gtgtggctgc cgccgcccag cgacgcggcc 240
tccaacgagg gctacctgcc gcgccggctc gagctgctgg acagcaagta tggcaccggc 300
acggacctgg tcaacgccct gtccgcgctg aatgccaatg gtgtgaagcc cattgcggac 360
atcgtcatca accaccgcgt gggcaccacg ggctgggcgg acttcacgct gcctccgtgg 420
ggctcgaacg cgggtgtgccg cggcgacgag tggagcgggg ccacgggcaa cgcggtatgc 480
ggcgtatggc tcaacgccgg gcgcgacatc gatcacacgc agaccttcgt gcaggacggc 540
atcgtcacct ggatgaacaa ctcgctgaag agcgtcgggt tcgcgggttg gcggtatgac 600
tacgtgaagg gctacagcgg ctcctacgtc ggctcgtaca acaccgcac gacgccgtac 660
ttctccgtgg gcgagctgtg gacggacctg gacctgaaca accccaacc ccaccgccag 720
ctgatcatga attggtatga cgcgacgggt ggcgggtccg cggcgttcga cttcacgacc 780
aagggcctgc tgcagcaggc ggtgcagtac aacgagttct ggcggtgaa ggatgcggcg 840
ggcgcgccag cgggtgccat tggttggtgg gcagcgaagt ccgtgacctt catcgacaat 900
cacgacacgg gcccgagcta tccgagcggc ggccagaacc actggccgtt ccctgggtgac 960
aagatcctcc aggggtacgc ctacatcctg actcactctg gcatcccctg cgtgtactgg 1020
gtgcactaca aggactgggg ccaggcgaac acggacgcca tcaagaagct gatcagcatc 1080
cgcaagtcca agggcatcac cagcacctcc tcggtgagca tccaggccgc ggacagctcg 1140
aagtacgccg ccatcatcac cggcaacaac ggcaagggtg ccgtgaagat cggcttcggc 1200
gcctggtctc cgccgggcac ctggacgctg gccacctccg gcaacaacta cgccgtctgg 1260
acgcagtaa 1269

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<210> 379
 <211> 422
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(27)

```

<400> 379
Met Leu Lys Gln Phe Thr Lys Arg Leu Ile Thr Leu Thr Ser Leu Leu
1 5 10 15
Ala Leu Val Leu Val Ala Pro Leu Ala Ser Ala Gly Pro Leu Asp Gly
20 25 30
Asn Ser Ser Asp Val Met Leu Gln Gly Phe His Trp Tyr Ser Tyr Gln
35 40 45
Ser Phe Pro Trp Trp Gly Val Ile Lys Asn Asn Ala Ala Ser Ile Lys
50 55 60

```

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Ala 65	Asp	Gly	Phe	Thr	Met 70	Val	Trp	Leu	Pro	Pro 75	Pro	Ser	Asp	Ala	Ala 80
Ser	Asn	Glu	Gly	Tyr 85	Leu	Pro	Arg	Arg	Leu 90	Glu	Leu	Leu	Asp	Ser 95	Lys
Tyr	Gly	Thr	Arg 100	Thr	Asp	Leu	Val	Asn 105	Ala	Leu	Ser	Ala	Leu 110	Asn	Ala
Asn	Gly	Val 115	Lys	Pro	Ile	Ala	Asp 120	Ile	Val	Ile	Asn	His 125	Arg	Val	Gly
Thr	Thr 130	Gly	Trp	Ala	Asp	Phe 135	Thr	Leu	Pro	Pro	Trp 140	Gly	Ser	Asn	Ala
Val 145	Cys	Arg	Gly	Asp	Glu 150	Trp	Ser	Gly	Ala	Thr 155	Gly	Asn	Ala	Asp	Thr 160
Gly	Asp	Gly	Phe	Asn 165	Ala	Gly	Arg	Asp	Ile 170	Asp	His	Thr	Gln	Thr 175	Phe
Val	Gln	Asp	Gly 180	Ile	Val	Thr	Trp	Met 185	Asn	Asn	Ser	Leu	Lys 190	Ser	Val
Gly	Phe 195	Ala	Gly	Trp	Arg	Tyr	Asp 200	Tyr	Val	Lys	Gly	Tyr 205	Ser	Gly	Ser
Tyr	Val 210	Gly	Ser	Tyr	Asn	Thr 215	Arg	Thr	Thr	Pro	Tyr 220	Phe	Ser	Val	Gly
Glu 225	Leu	Trp	Thr	Asp	Leu 230	Asp	Leu	Asn	Asn	Pro 235	Asn	Pro	His	Arg	Gln 240
Leu	Ile	Met	Asn	Trp 245	Ile	Asp	Ala	Thr	Gly 250	Gly	Arg	Ser	Ala	Ala 255	Phe
Asp	Phe	Thr	Thr 260	Lys	Gly	Leu	Leu	Gln 265	Ala	Val	Gln	Tyr 270	Asn	Glu	
Phe	Trp	Arg 275	Leu	Lys	Asp	Ala	Ala 280	Gly	Ala	Pro	Ala	Gly 285	Ala	Ile	Gly
Trp	Trp 290	Ala	Ala	Lys	Ser	Val 295	Thr	Phe	Ile	Asp	Asn 300	His	Asp	Thr	Gly
Pro 305	Ser	Tyr	Pro	Ser	Gly 310	Gly	Gln	Asn	His	Trp 315	Pro	Phe	Pro	Gly	Asp 320
Lys	Ile	Leu	Gln	Gly 325	Tyr	Ala	Tyr	Ile	Leu 330	Thr	His	Ser	Gly	Ile 335	Pro
Cys	Val	Tyr	Trp 340	Val	His	Tyr	Lys	Asp 345	Trp	Gly	Gln	Ala	Asn	Thr	Asp
Ala	Ile	Lys 355	Lys	Leu	Ile	Ser	Ile	Arg	Lys	Ser	Lys	Gly 365	Ile	Thr	Ser
Thr	Ser	Ser	Val	Ser	Ile	Gln 375	Ala	Ala	Asp	Ser	Ser	Lys	Tyr	Ala	Ala
Ile 385	Ile	Thr	Gly	Asn	Asn 390	Gly	Lys	Val	Ala	Val 395	Lys	Ile	Gly	Phe	Gly 400
Ala	Trp	Ser	Pro	Pro 405	Gly	Thr	Trp	Thr	Leu 410	Ala	Thr	Ser	Gly	Asn	Asn 415
Tyr	Ala	Val	Trp	Thr	Gln										
			420												

<210> 380

<211> 1644

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 380

atgaagccgc	gttgggcagc	gtggcttgct	ctcgtatgct	tgctggcggg	ctgcttcggg	60
ggagaaagcg	cgctgcccga	tcattgcggc	cggtcttcgg	ctccggacgg	aacgccggcg	120
ccggtcccgt	ccgctgcagc	cgcaggggtg	aaagcggatg	aacaaccgtc	gacgggtgtt	180
tacgagattt	tcgtgcgctc	gttctacgac	tccgacggag	acggcatcgg	ggatttgaac	240
ggcgtgacgg	agaagctcga	ctatttgaag	gagttgggag	tcggcggcat	ctggctgatg	300
ccgatcaacg	cgctgcggag	ctaccatggc	tacgatgtga	cggactacta	tgcgatcaac	360
cccgactacg	gcacgttgga	agacttgaaa	aggctgctgg	ccgaagcgca	tgcgcggggc	420
atcaaagtca	ttatggatct	tgtcgtcaac	catacagacc	gcgagcatcc	ctggttcaag	480
gaggcgcttg	ccgatccgga	cagcccttac	cgcaactggt	atacgatcaa	accgtcggac	540
gaggcggcgc	cttcggacaa	cgcggcgggg	accggcagtc	cctggcatga	tcggggcagc	600
tacaaatacc	tcgggatttt	ctgggaaggc	atgcccgatt	tgaatttcga	cgagccgaag	660
gttcgcgaag	agatgatcaa	ggtcggccgg	tactggctgg	agcagggggg	tgacggcttc	720

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cggctggatg cggccaagca tatttacggg gatttcgcct ccaccgtcca ctcggaggaa 780
atcgcgggca aaaacaaagc gtggtggcag gaattccgca gcgggttgaa cgaggccaat 840
ccgaacgctt acctcgtggg cgaaatctgg gattcggcga cgacgatcgc gcctctgctc 900
gacaaggcgc tcgattccgg attcaacttc gatttggcgg gacggattct cgacgcccgcg 960
aagggggaac gcgacccgga cgtcgccttc agcctgaagc ggggtatacgg cgcttacggc 1020
caagcgctccg gcggttcggt cacggacgcg acgtttctct ccaatcacga ccagaaccgc 1080
gccatgagcg tgctcggcgg gaacgtaaatt cacgcgaaga tggccgcggc ggtgctgctg 1140
acgctgccgg gcaatccttt cctttattac ggggaggaaa tcggcatgct cggcatgaag 1200
cccgatgaag cgatccgcga gcccatgcct tggtagccg gcccatcggg cgggcccggg 1260
cagacgtctt gggaaccgct gcggtacaac aagcccggcg ccacatcggg cgagcagcag 1320
gagcgggacg agcggtcgct gctccatcat taccgcacgc tgatcaagtg gcggaatgaa 1380
atgcccagac tgcgggacgg ggatatcgac gagtacgata ccggcaacgc gaagctcggc 1440
gcttacgtgc ggatgacgaa ggacgccgcg gtcctcgtcg tgcacaatat gagcggcgag 1500
aagcagtcgg cgaagctgac ggcgtcggag cggtagcgag cgttccggga aatcgtgcgc 1560
cagaccggca gcggcggcga aggcgttcg cttgacgggg aaacgctctc catcgctccg 1620
tacagcacgg tcatcatccg ctag 1644

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<210> 381
 <211> 547
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(29)

<400> 381
 Met Lys Pro Arg Trp Ala Ala Trp Leu Val Leu Val Cys Leu Leu Ala
 1 5 10 15
 Gly Cys Phe Gly Gly Glu Ser Ala Pro Ala Asp His Ala Ala Arg Ser
 20 25 30
 Ser Ala Pro Asp Gly Thr Pro Ala Pro Val Pro Ser Ala Ala Ala Ala
 35 40 45
 Gly Trp Lys Ala Asp Glu Gln Pro Ser Thr Val Phe Tyr Glu Ile Phe
 50 55 60
 Val Arg Ser Phe Tyr Asp Ser Asp Gly Asp Gly Ile Gly Asp Leu Asn
 65 70 75 80
 Gly Val Thr Glu Lys Leu Asp Tyr Leu Lys Glu Leu Gly Val Gly Gly
 85 90 95
 Ile Trp Leu Met Pro Ile Asn Ala Ser Pro Ser Tyr His Gly Tyr Asp
 100 105 110
 Val Thr Asp Tyr Tyr Ala Ile Asn Pro Asp Tyr Gly Thr Leu Glu Asp
 115 120 125
 Leu Lys Arg Leu Leu Ala Glu Ala His Ala Arg Gly Ile Lys Val Ile
 130 135 140
 Met Asp Leu Val Val Asn His Thr Ser Arg Glu His Pro Trp Phe Lys
 145 150 155 160
 Glu Ala Leu Ala Asp Pro Asp Ser Pro Tyr Arg Asn Trp Tyr Thr Ile
 165 170 175
 Lys Pro Ser Asp Glu Ala Ala Pro Ser Asp Asn Ala Ala Gly Thr Gly
 180 185 190
 Ser Pro Trp His Asp Ala Gly Ser Tyr Lys Tyr Leu Gly Ile Phe Trp
 195 200 205
 Glu Gly Met Pro Asp Leu Asn Phe Asp Glu Pro Lys Val Arg Glu Glu
 210 215 220
 Met Ile Lys Val Gly Arg Tyr Trp Leu Glu Gln Gly Val Asp Gly Phe
 225 230 235 240
 Arg Leu Asp Ala Ala Lys His Ile Tyr Gly Asp Phe Ala Ser Thr Val
 245 250 255
 His Ser Glu Glu Ile Ala Gly Lys Asn Lys Ala Trp Trp Gln Glu Phe
 260 265 270
 Arg Ser Gly Leu Asn Glu Ala Asn Pro Asn Ala Tyr Leu Val Gly Glu
 275 280 285
 Ile Trp Asp Ser Ala Thr Thr Ile Ala Pro Leu Leu Asp Lys Ala Leu
 290 295 300
 Asp Ser Gly Phe Asn Phe Asp Leu Ala Gly Arg Ile Leu Asp Ala Ala

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305      310      315      320
Lys Gly Glu Arg Asp Pro Asp Val Ala Phe Ser Leu Lys Arg Val Tyr
      325      330      335
Gly Ala Tyr Gly Gln Ala Ser Gly Gly Ser Phe Thr Asp Ala Thr Phe
      340      345      350
Leu Ser Asn His Asp Gln Asn Arg Ala Met Ser Val Leu Gly Gly Asn
      355      360      365
Val Asn His Ala Lys Met Ala Ala Val Leu Leu Thr Leu Pro Gly
      370      375      380
Asn Pro Phe Leu Tyr Tyr Gly Glu Glu Ile Gly Met Leu Gly Met Lys
      385      390      395      400
Pro Asp Glu Ala Ile Arg Glu Pro Met Pro Trp Tyr Ala Gly Pro Ser
      405      410      415
Gly Gly Pro Gly Gln Thr Ser Trp Glu Pro Leu Arg Tyr Asn Lys Pro
      420      425      430
Gly Ala Thr Ser Val Glu Gln Gln Glu Arg Asp Glu Arg Ser Leu Leu
      435      440      445
His His Tyr Arg Thr Leu Ile Lys Trp Arg Asn Glu Met Pro Glu Leu
      450      455      460
Arg Asp Gly Asp Ile Asp Glu Tyr Asp Thr Gly Asn Ala Lys Leu Gly
      465      470      475      480
Ala Tyr Val Arg Met Thr Lys Asp Ala Arg Val Leu Val Val His Asn
      485      490      495      499
Met Ser Gly Glu Lys Gln Ser Ala Lys Leu Thr Ala Ser Glu Arg Tyr
      500      505      510
Gly Ala Phe Arg Glu Ile Val Arg Gln Thr Gly Ser Gly Glu Gly
      515      520      525
Ala Ser Leu Asp Gly Glu Thr Leu Ser Ile Ala Pro Tyr Ser Thr Val
      530      535      540
Ile Ile Arg
545

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 <211> 4176
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

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accgacgccc aactggccgc cgagccgacc cggcacgccc tcaccaccga gcagttctac      180
ttcgtgctcc cggaccgctt cgccaacggc gaccggtcca acgacaccgc cggcatcccc      240
ggcgggccgc tcgaccacgg gtacgacccc gacgacgagg gcttctacca gggcggcgac      300
ctcgcgggcc tgatcgagaa cctggactac atcgagggcc tgggcaccac cgccctctgg      360
atggcgcccc tcttcgagaa caagccggtc cagggcgagg gggaccacgc ctcggcgggc      420
taccacggct actggatcac cgacttcacc cgggtggacc cgcacttcgg caccaacgag      480
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ggtgccgagg acgggggggt ccccgaactc gacaccgcct ccttcccgtc cacgccgggtg      720
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cacaaccgcg gcgactctc cttcgccggc gagagtccc tgtacggcga cttccacggc      840
ctggacgacc tgttcaccgc ccggcccag gtcgtcgagg gcatggtgga gatctaccgc      900
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atggagtctt ggaccgcgtg gtccgcccgc atgaaggagc acgccgcgtc cctcgccac      1020
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gtcaccgagg gcggcctgaa cgccaccctg gacttcgcct tccaggaggc cgcccgtctc      1140
ttcgctcgcg agggcggtc cgcgaagcgc ctgtccgccc tcttcggcga ggaccacgc      1200
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ttcggttcca cgacgccgt ctacctggag gaccgcgaac tgggcaccga ccgacccac      1500
gcggaggacg cccacgacac cgaccatccc ctgtaccggg cgatcgccgc cctctccgc      1560
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1001827087_1.txt

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gccgccaaca acgcgaggga cggacgcacc gtcgccgtgc ccaccgacac ccccgccgcc 1740
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 <211> 1391
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(35)

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 Ala Gln Ala Ala Pro Pro Pro Pro Thr Asp Ala Glu Leu Ala Ala Glu
 35 40 45
 Pro Thr Arg His Ala Leu Thr Thr Glu Gln Phe Tyr Phe Val Leu Pro
 50 55 60
 Asp Arg Phe Ala Asn Gly Asp Pro Ser Asn Asp Thr Ala Gly Ile Pro
 65 70 75 80
 Gly Gly Pro Leu Asp His Gly Tyr Asp Pro Asp Asp Glu Gly Phe Tyr
 85 90 95

1001827087_1.txt

Gln	Gly	Gly	Asp	Leu	Arg	Gly	Leu	Ile	Glu	Asn	Leu	Asp	Tyr	Ile	Glu
Gly	Leu	Gly	Thr	Thr	Ala	Leu	Trp	Met	Ala	Pro	Val	Phe	Glu	Asn	Lys
Pro	Val	Gln	Gly	Glu	Gly	Asp	His	Ala	Ser	Ala	Gly	Tyr	His	Gly	Tyr
Trp	Ile	Thr	Asp	Phe	Thr	Arg	Val	Asp	Pro	His	Phe	Gly	Thr	Asn	Glu
145	Glu	Leu	Ala	Glu	Leu	Val	Gly	Ala	Ala	His	Glu	Arg	Asp	Ile	Lys
	Phe	Phe	Asp	Val	Ile	Thr	Asn	His	Thr	Ala	Asp	Val	Ile	His	Tyr
	Glu	Gly	Ala	Tyr	Asp	Tyr	Leu	Ser	Lys	Gly	Ala	Phe	Pro	Tyr	Leu
	Thr	Glu	Gly	Arg	Pro	Phe	Asp	Asp	Leu	Asp	His	Ala	Gly	Ala	Glu
	Gly	Gly	Phe	Pro	Glu	Leu	Asp	Thr	Ala	Ser	Phe	Pro	Tyr	Thr	Pro
	Thr	Ser	Asp	Ala	Glu	Ala	Asp	Leu	Lys	Val	Pro	Asp	Trp	Leu	Asn
	Pro	Thr	Leu	Tyr	His	Asn	Arg	Gly	Asp	Ser	Ser	Phe	Ala	Gly	Glu
	Ser	Leu	Tyr	Gly	Asp	Phe	His	Gly	Leu	Asp	Asp	Leu	Phe	Thr	Ala
	Pro	Glu	Val	Val	Glu	Gly	Met	Val	Glu	Ile	Tyr	Arg	Thr	Trp	Val
	Asp	Phe	Gly	Ile	Asp	Gly	Phe	Arg	Ile	Asp	Thr	Val	Lys	His	Val
	Met	Glu	Phe	Trp	Thr	Arg	Trp	Ser	Ala	Ala	Met	Lys	Glu	His	Ala
	Ser	Leu	Gly	His	Glu	Asp	Phe	Phe	Met	Phe	Gly	Glu	Val	Tyr	Ser
	Asp	Val	Ala	Glu	Lys	Ala	Arg	Tyr	Val	Thr	Glu	Gly	Gly	Leu	Asn
	Thr	Leu	Asp	Phe	Ala	Phe	Gln	Glu	Ala	Ala	Arg	Ser	Phe	Ala	Ser
	Gly	Gly	Ser	Ala	Lys	Arg	Leu	Ser	Ala	Leu	Phe	Gly	Glu	Asp	His
	Tyr	Thr	Thr	Ala	Asp	Thr	Asn	Ala	Gln	Gln	Ser	Val	Thr	Phe	Leu
	Asn	His	Asp	Met	Gly	Arg	Ile	Gly	His	Phe	Leu	Arg	Ala	Asp	Asn
	Asp	Ala	Asp	Glu	Ala	Glu	Leu	Leu	Glu	Arg	Ser	Arg	Leu	Ala	His
	Leu	Met	Phe	Leu	Ser	Arg	Gly	Asn	Pro	Val	Ile	Tyr	Tyr	Gly	Asp
	Gln	Gly	Phe	Thr	Gly	Ser	Gly	Gly	Asp	Lys	Ala	Ala	Arg	Ala	Thr
	Phe	Gly	Ser	Thr	Thr	Pro	Leu	Tyr	Leu	Glu	Asp	Pro	Gln	Leu	Gly
	Asp	Arg	Thr	His	Ala	Glu	Asp	Ala	His	Asp	Thr	Asp	His	Pro	Leu
	Arg	Ala	Ile	Ala	Ala	Leu	Ser	Ala	Leu	Thr	Arg	Glu	His	Pro	Ala
	Arg	Asp	Gly	Ala	Gln	Thr	Glu	Arg	Tyr	Ala	Glu	Asp	Gly	Ala	Gly
	Arg	Ala	His	Thr	Arg	Thr	Asp	Ala	Asp	Glu	Arg	Val	Glu	Tyr	Leu
	Ala	Ala	Asn	Asn	Ala	Glu	Asp	Gly	Arg	Thr	Val	Ala	Val	Pro	Thr
	Thr	Pro	Gly	Ala	Val	Phe	Glu	Val	Leu	Tyr	Gly	Gly	Val	Gly	Asp
	Ser	Ala	Thr	Pro	Val	Thr	Ala	Asp	Ala	Asp	Gly	Val	Val	Glu	Leu
	Val	Pro	Ala	Leu	Ser	Thr	Ala	Val	Leu	Arg	Ala	Asp	Arg	Pro	Leu
	Asp	Pro	Gln	Ala	Ala	Pro	Ala	Val	Glu	Leu	Thr	Ala	Pro	Asp	Ala
	625					630									

1001827087_1.txt

Ala	His	Gly	Val	Val	Glu	Leu	Ser	Ala	Ala	Val	Thr	Gly	Gly	Ala	Leu
Asn	Arg	Val	Ala	Phe	Ala	Ala	Gln	Val	Gly	Asp	Gly	Asp	Trp	Glu	Val
Leu	Gly	Val	Ser	Asp	Ala	Pro	Pro	His	Arg	Val	Thr	His	Thr	Val	Pro
Ala	Gly	Thr	Ser	Ala	Gly	Thr	Pro	Leu	Arg	Tyr	Arg	Ala	Val	Ala	Val
Asp	Ser	Ser	Gly	Ala	Thr	Ala	Ala	Thr	Gly	Ala	Glu	Thr	Thr	Ala	Gly
Ala	Pro	Pro	Pro	Glu	Arg	Ala	Pro	Arg	Ala	Val	Ser	Arg	Glu	Trp	Ala
Val	Val	His	Tyr	His	Arg	Pro	Asp	Gly	Asn	Tyr	Asp	Asp	Trp	Ser	Leu
Trp	Ala	Trp	Gly	Asp	Leu	His	Glu	Ser	Glu	Ser	Phe	Glu	Asp	Trp	Pro
His	Gly	Arg	Pro	Phe	Val	Gly	Arg	Asp	Ala	Phe	Gly	Ala	Phe	Ala	Arg
Val	Arg	Leu	Ala	Pro	Asp	Ala	Ser	Glu	Leu	Gly	Tyr	Leu	Ile	Val	Asp
Gly	Asp	Gly	Thr	Lys	Asp	Gly	Asp	Leu	Asp	Arg	Val	Ile	Asp	Leu	Ser
Glu	Thr	Gly	Glu	Val	Trp	Ile	Glu	Gln	Gly	Glu	Pro	Glu	Ala	Leu	Thr
Glu	Ala	Pro	Glu	Gly	Val	Tyr	Pro	Ala	Asp	Glu	Ser	Arg	Ala	Val	
Leu	Arg	Tyr	His	Arg	Pro	Asp	Gly	Asp	Tyr	Asp	Gly	Trp	Gly	Val	His
Val	Trp	Thr	Gly	Ala	Ala	Glu	Pro	Thr	Asp	Trp	Ser	Arg	Pro	Val	Pro
Ala	Ser	Glu	Val	Asp	Ser	Phe	Gly	Ala	Val	Tyr	Glu	Ile	Pro	Leu	Ala
Glu	Gly	Ala	Thr	Ser	Leu	Ser	Tyr	Ile	Ile	His	Arg	Gly	Asp	Glu	Lys
Asp	Leu	Pro	Ala	Asp	Gln	Ser	Leu	Asp	Leu	Arg	Ala	Val	Gly	Tyr	Glu
Val	Trp	Ile	Ile	Ser	Gly	Glu	Glu	Thr	Tyr	Leu	Leu	Pro	Ser	Gly	Gly
Ala	Ala	Pro	Asp	Leu	Asn	Pro	Asp	Arg	Ala	Arg	Ala	His	Trp	Ile	Asp
Ala	Glu	Thr	Val	Val	Trp	Pro	Gly	Val	Ala	Pro	Pro	Gly	Gly	Ser	Arg
Gln	Leu	Ile	His	His	Pro	Glu	Gly	Ala	Ile	Ala	Leu	Glu	Asp	Gly	Ala
Leu	Thr	Asp	Glu	Gly	His	Trp	Ile	Thr	Thr	Pro	Pro	Ala	Gly	Glu	Arg
Asp	Pro	Arg	Pro	Gly	Leu	Thr	Gly	Leu	Glu	Pro	Ser	Asp	Met	Gly	Arg
Trp	Phe	Ala	Glu	Arg	Gly	Gln	Pro	Gly	Tyr	Arg	Ala	Arg	Gln	Leu	Ala
Asp	His	Val	Trp	Gln	Ala	Cys	Ala	Ile	Arg	Ala	Asp	Glu	Leu	Arg	Thr
Leu	Pro	Ala	Ala	Leu	Arg	Thr	Asp	Ile	Glu	Asp	Ser	Phe	Arg	Val	Asp
Thr	Leu	Val	Glu	Thr	Asp	Val	Arg	Pro	Ala	Asp	Ala	Gly	Leu	Thr	Glu
Lys	Ala	Leu	His	Arg	Leu	Asp	Asp	Gly	Arg	Leu	Ile	Glu	Ser	Val	Leu
Met	Arg	Tyr	Pro	Ala	Arg	Gly	Pro	Arg	Arg	Trp	Arg	Ala	Thr	Ala	Cys
Ile	Ser	Ser	Gln	Ala	Gly	Cys	Ala	Val	Gly	Cys	Pro	Phe	Cys	Ala	Thr
Gly	Glu	Leu	Gly	Phe	Met	Arg	Asp	Leu	Glu	Val	Ser	Glu	Ile	Val	Asp
Gln	Ala	Arg	Tyr	Trp	Arg	Arg	Arg	Leu	Ala	Ala	Glu	Gly	Arg	Arg	Leu
Thr	Asn	Leu	Val	Phe	Met	Gly	Met	Gly	Glu	Pro	Leu	Leu	Asn	Pro	Asp

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Arg Val Leu Ala Ala Ala Glu Ala Leu Ser Asp Ala Arg Arg Phe Gly
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 1205 1210 1215
 Met Glu Arg Leu Ile Ala Leu Arg Pro Gln Tyr Thr Leu Ala Val Ser
 1220 1225 1230
 Leu His Ala Ala Arg Pro Asp Leu Arg Asp Val Leu Val Pro Leu Asn
 1235 1240 1245
 Arg Arg Tyr Pro Val Gly Asp Val Val Ala Ala Ala Ser Ala Tyr Ala
 1250 1255 1260
 Arg Leu Thr Gly Arg Arg Val Ser Tyr Glu Tyr Val Met Ile Asp Ser
 1265 1270 1275 1280
 Val Asn Asp Thr Pro Arg Asp Ala Ala Glu Leu Thr Gln Leu Leu Gly
 1285 1290 1295
 Gly Arg His Ala His Val Asn Leu Ile Pro Met Asn Pro Val Ala His
 1300 1305 1310
 Thr Pro Trp Gln Ala Ser Pro Thr Ser Arg Ile Ala Glu Phe Ala Ser
 1315 1320 1325
 Leu Leu Arg Ser Gly Gly Ile Arg Thr Thr Val Arg Arg Asn Arg Gly
 1330 1335 1340
 Leu Glu Ile Gly Ala Ala Cys Gly Gln Leu Ala Ala Asp Ser Ala Gly
 1345 1350 1355 1360
 Ala Pro Pro Pro Ala Ala Val Ala Arg Arg Arg Glu Leu Leu Val Leu
 1365 1370 1375
 Asn Ser Ala Ala Leu Ala Ala Arg Pro Ala Val Ser Arg Ser
 1380 1385 1390

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 <211> 1434
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 ttgcacgcat tcaactggcg ctatgccacc gtcgaggcac gtgcaaacga gattgccgcg 180
 ctcggttatc gcgccgtgct cgctgccccg ccactgcgct cgcagggcaa cgaatggtgg 240
 gccgcctacc agccgcagga ctaccgcgtc atcgagcacc cgctcggcga caccgacgcg 300
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 aaggcgctcg gtgtcagcgg tttccgcata gacgccgcca agcacatgcc gaacacgcac 720
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 <211> 477
 <212> PRT
 <213> Unknown

<220>
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<221> SIGNAL
<222> (1)...(37)

<400> 385

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Pro Pro Arg Arg Thr Trp Trp Ile Val Ala Leu Leu Leu Val Ser
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Ala Ser Ala Gln Ala Asp Ala Val Leu His Ala Phe Asn Trp Arg Tyr
 35      40      45
Ala Thr Val Glu Ala Arg Ala Asn Glu Ile Ala Ala Leu Gly Tyr Arg
 50      55      60
Ala Val Leu Val Ala Pro Pro Leu Arg Ser Gln Gly Asn Glu Trp Trp
 65      70      75      80
Ala Arg Tyr Gln Pro Gln Asp Tyr Arg Val Ile Glu His Pro Leu Gly
 85      90      95
Asp Thr Asp Ala Phe Val Arg Met Glu Ser Ala Leu Arg Gln Ala Gly
100      105      110
Val Arg Leu Tyr Ala Asp Ile Val Leu Asn His Met Ala Asn Glu Ser
115      120      125
Gly Trp Arg Ser Asp Leu Asn Tyr Pro Gly Thr Leu Val Leu Asn Gln
130      135      140
Tyr Ala Ala Asn Pro Ser Arg Tyr Ala Arg Leu Arg Leu Phe Gly Asp
145      150      155      160
Leu Arg Tyr Asn Phe Leu Ser Gln His Asp Phe Gly Pro Ala Asn Cys
165      170      175
Ile Thr Asp Tyr Gly Ser Val Trp Gln Val Gln Thr Trp Arg Leu Cys
180      185      190
Gly Ala Ala Pro Asp Val Gly Leu Pro Asp Leu Val Asp Asn Asp Trp
195      200      205
Val Val Ser Gln Gln Gln Gln Tyr Leu Arg Ser Leu Lys Ala Leu Gly
210      215      220
Val Ser Gly Phe Arg Ile Asp Ala Ala Lys His Met Pro Asn Thr His
225      230      235      240
Met Asn Arg Val Leu Thr Ser Asn Ile Lys Gln Gly Val His Val Phe
245      250      255
Gly Glu Ile Ile Thr Asp Gly Gly Val Gly Asp Thr Ser Tyr Asp Arg
260      265      270
Phe Leu Ala Pro Tyr Leu Ala Gln Thr Asp His Gly Ala Tyr Asp Phe
275      280      285
Pro Leu Phe His Gln Ile Arg Asn Ala Phe Arg Pro Gly Gly Ser Met
290      295      300
Asn Leu Leu Val Asp Pro Gly Ala Tyr Gly Gln Ala Leu Pro Arg His
305      310      315      320
Arg Ala Ile Thr Phe Thr Val Thr His Asp Ile Pro Asn Asn Gly Ile
325      330      335
Phe Arg Tyr Leu Ile Leu Asp Arg Thr Asp Glu Thr Leu Ala Tyr Ala
340      345      350
Tyr Val Leu Gly Arg Asp Gly Gly Val Pro Leu Ile Tyr Ser Asp Asn
355      360      365
Asn Glu Ser Gly Asp Asn Arg Trp Val Asn Ala Tyr Arg Arg Ser Asp
370      375      380
Leu Ala Gln Met Ile Arg Phe His Asn Ser Val Gln Gly Ser Asp Met
385      390      395      400
Gln Val Leu Ser Tyr Gly Ser Cys His Leu Leu Phe Arg Arg Gly Asn
405      410      415
Arg Gly Ile Val Gly Ile Asn Lys Cys Gly Ser Ser Val Asn Ala Thr
420      425      430
Val Asn Met His Gly Ser Val Leu Trp Trp Tyr Thr Asp Tyr Arg Asp
435      440      445
Val Leu Gly Ser Gly His Val Val Asn Ile Gly Ser Gly Ser Tyr Thr
450      455      460
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<210> 386

<211> 1458

<212> DNA
<213> Unknown

<220>
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gatatcgctc gtgagtgcga aacgtactta ggcccaaacg gctttgctgc aattcaagtt      180
tcgccgccga acgagcatgc ggttgttacc gggcgcccgt ggtacgaacg ttaccaaccg      240
gtgtcgtata ttttaaatc acgcagcggc aaccgcgaag aatttattaa catggttaag      300
gcttgtaaaa aagcagggtg ggatatttat gtcgatgcag tgattaacca tatgaccgga      360
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tatccggatt attcttacga tgactttcac cattgcggtc gcaacggtaa tgatcagatc      480
agtaattatg gtgacagata tgaagtgcaa aactgtgcgc tggttgggtc tgcagattta      540
aacatcgaaa aagattatgt gcgcgaccgt atcgccagat atttaaataa attagtcgat      600
attggtgtgg ctggatttcg catagatgcc gccaaacata tgtcgactaa ctcgttaaat      660
gaaattatta aacgtgtaaa aggcacacgt tatatttatc aggaagtgat tgatcaaggc      720
ggtagccga ttcaggccaa agagtatttt aataatggtg atgtgacaga gtttaaataa      780
tcggtggata tgtcgcgtgt gatgttaaac ggaataactaa cctggttaaa tgataaggcc      840
cagtttggtg aaggctggaa ctacatgcca tcagataaag cgggtggtgtt tattgataac      900
catgataacc aacgcggtca tggcggcggc ggacatattt taaactataa agacggacgt      960
aattatgagt tggccagtgt actgatgctt gcatggcctt acggttacc acaggtgatg     1020
tcgagctttg cgtttaccga tccaaactcg tcaccgccgg ctgatatgta aggtaatacg     1080
aaagatacga attgcctttc tcgtgcagag gggccggcgg ctcaggatgg ttgggtgtgt     1140
gaacacgggt ggccgatgat tcgtgaaatg gttaaattca gaaatgtgac cagcagtaag     1200
ttttatttat cgaactgggt gagtaacggg aataaccaa ttgctttttc gcgcgggtgat     1260
ttaggatttg ttgtaatcaa ccacgaagag caggtgttgc gtcaacgatt gcaaaccggg     1320
ttaactgcag gtgtctactg tgatgtttta agtcaaaatt gtgaacgcac aattactggt     1380
aatgacaatg gtgaagctga aatcatcggt acgggtcaaa atgcagcggc catccatgcc     1440
cagagtaaat tgaagtag                                     1458
```

<210> 387
<211> 485
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(25)

```
<400> 387
Met Lys Asn Ser Asn Ala Leu Lys Asn Ile Leu Leu Ser Ser Val Ile
 1          5          10          15
Leu Thr Ser Gly Leu Leu Ala Gln Ala Asn Pro Arg Thr Val Met Val
          20          25          30
His Leu Phe Glu Trp Lys Trp Asn Asp Ile Ala Arg Glu Cys Glu Thr
          35          40          45
Tyr Leu Gly Pro Asn Gly Phe Ala Ala Ile Gln Val Ser Pro Pro Asn
          50          55          60
Glu His Ala Val Val Thr Gly Arg Pro Trp Tyr Glu Arg Tyr Gln Pro
65          70          75          80
Val Ser Tyr Ile Leu Asn Ser Arg Ser Gly Asn Arg Glu Glu Phe Ile
          85          90          95
Asn Met Val Lys Ala Cys Lys Lys Ala Gly Val Asp Ile Tyr Val Asp
          100          105          110
Ala Val Ile Asn His Met Thr Gly Thr Leu Ala His Gly Glu Glu Arg
          115          120          125
Val Gly Ile Ser Gly Ser Arg Phe Gly Arg Phe Ala Tyr Pro Asp Tyr
          130          135          140
Ser Tyr Asp Asp Phe His His Cys Gly Arg Asn Gly Asn Asp Gln Ile
145          150          155          160
Ser Asn Tyr Gly Asp Arg Tyr Glu Val Gln Asn Cys Ala Leu Val Gly
          165          170          175
Leu Ala Asp Leu Asn Ile Glu Lys Asp Tyr Val Arg Asp Arg Ile Ala
```

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[illegible]

<210>	388
<211>	1662
<212>	DNA
<213>	Unknown

<220>
<223> obtained from an environmental sample

<400>	388						
atgagccaca	tcctgctgtgc	cgccgtattg	gcgggcggtcc	tgctgccggtt	tcccgcactg		60
gccgatcagg	ccggcaagag	cccggccggg	gtgcgcctacc	acggcggcga	cgaaatcatc		120
ctccagggct	tccactggaa	cgtcgtccgc	gaagcgccca	acgactggta	caacatcctc		180
cgccaacagg	cctcgcacgat	cgcggccgac	ggcttctcgg	caatctggat	gccggtgccc		240
tggcgctgact	tctccagctg	gagcgcagcgc	ggcaagtctg	gcggcggcga	aggctacttc		300
tggcacgatt	tcaacaagaa	cgggcgctat	ggcagcgatg	cccagctgcg	ccaggccgcc		360
ggcgcgctcg	gcggtgccgg	ggtgaagggtg	ctctacgacg	tggtgcccaa	ccacatgaac		420
cgcggtctacc	cgaacaagga	aatcaacctg	ccggccggcc	agggttcttg	gcgcaacgac		480
tgcgccgacc	cgggcaacta	ccccaacgac	tgcgacgacg	gcgatcgttt	cgtcggcggc		540
gatgccgacc	tcaataccgg	ccatccgcag	gtctatggca	tgtttccgcga	cgagttcgcc		600
aacctgcgca	gccagtacgg	cgccggcggc	ttcctgtctg	atttctctcg	cgtcttcgcc		660
ccggagcggg	tcaacagctg	gatgaccgac	agtgcgcgaca	acagcttctg	cgctcggcgag		720
ctgtggaaaag	gcccttctga	atatcccagc	tgggactggc	gcaacacggc	gagctggcag		780
cagatcatca	aggactggtc	cgatcgggcc	aagtgcgccg	tgttcgactt	cgccctcaag		840
gagcgcatgc	agaacggctc	gatcgcgcac	tggagaagaag	gccttaacgg	caaccccgat		900
ccgcgcttgg	gcgaggtggc	agtgaccttt	gtcgacaacc	acgacacggg	ctactcgccc		960
gggcagaacg	gtgggcagca	tcacttgccc	ctcgaggacg	ggctgatctcg	tcaggcctac		1020
gcgtacattc	tgaccagccc	cggcacgcgc	gtggtgtact	ggtcgcacat	gtacgattgg		1080
ggctaccgcg	acttcatccg	ccagctgata	caggtgcgtc	gcgccgctgg	cgctgcgtgcc		1140

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```

gattcggcga tcagcttcca cagcggctac agcggcctcg tcgccaccgt caccggcagt 1200
cagcagaccc tgggtggtgc gctcaattcc aacctgagca atcccggcca ggtcgccagc 1260
ggcagcttca gcgaagcagt caataccagt aatggccagg tgcgggtctg gcggaccggc 1320
gccggcagtg gtggcgggtga caatggcggc ggtgagcccg gtgccctggt cagcgtgaac 1380
ttccgctgcg acaacggtgt gacgcaaccc ggcgacagcg tctacgcggt gggcaacgctc 1440
agccagctcg gcaactggag cccggcctct gcggtgcggc tgaccgatac cagcggctac 1500
ccgacctgga agggccgcat ctcgctgccg gccggccaga acgtggaatg gaagtgcctg 1560
atccgcaacg aggccaatgc aacgcaggtg cggcaatggc agggcggggc caacaacagc 1620
gtgaagccca ccgaagggtc caacacggcg gggcggttct ag 1662

```

<210> 389
 <211> 553
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(21)

<400> 389
 Met Ser His Ile Leu Arg Ala Ala Val Leu Ala Ala Val Leu Leu Pro
 1 5 10 15
 Phe Pro Ala Leu Ala Asp Gln Ala Gly Lys Ser Pro Ala Gly Val Arg
 20 25 30
 Tyr His Gly Gly Asp Glu Ile Ile Leu Gln Gly Phe His Trp Asn Val
 35 40 45
 Val Arg Glu Ala Pro Asn Asp Trp Tyr Asn Ile Leu Arg Gln Gln Ala
 50 55 60
 Ser Thr Ile Ala Ala Asp Gly Phe Ser Ala Ile Trp Met Pro Val Pro
 65 70 75 80
 Trp Arg Asp Phe Ser Ser Trp Ser Asp Gly Gly Lys Ser Gly Gly Gly
 85 90 95
 Glu Gly Tyr Phe Trp His Asp Phe Asn Lys Asn Gly Arg Tyr Gly Ser
 100 105 110
 Asp Ala Gln Leu Arg Gln Ala Ala Gly Ala Leu Gly Gly Ala Gly Val
 115 120 125
 Lys Val Leu Tyr Asp Val Val Pro Asn His Met Asn Arg Gly Tyr Pro
 130 135 140
 Asn Lys Glu Ile Asn Leu Pro Ala Gly Gln Gly Phe Trp Arg Asn Asp
 145 150 155 160
 Cys Ala Asp Pro Gly Asn Tyr Pro Asn Asp Cys Asp Asp Gly Asp Arg
 165 170 175
 Phe Val Gly Gly Asp Ala Asp Leu Asn Thr Gly His Pro Gln Val Tyr
 180 185 190
 Gly Met Phe Arg Asp Glu Phe Ala Asn Leu Arg Ser Gln Tyr Gly Ala
 195 200 205
 Gly Gly Phe Arg Phe Asp Phe Val Arg Gly Phe Ala Pro Glu Arg Val
 210 215 220
 Asn Ser Trp Met Thr Asp Ser Ala Asp Asn Ser Phe Cys Val Gly Glu
 225 230 235 240
 Leu Trp Lys Gly Pro Ser Glu Tyr Pro Ser Trp Asp Trp Arg Asn Thr
 245 250 255
 Ala Ser Trp Gln Gln Ile Ile Lys Asp Trp Ser Asp Arg Ala Lys Cys
 260 265 270
 Pro Val Phe Asp Phe Ala Leu Lys Glu Arg Met Gln Asn Gly Ser Ile
 275 280 285
 Ala Asp Trp Lys Asn Gly Leu Asn Gly Asn Pro Asp Pro Arg Trp Arg
 290 295 300
 Glu Val Ala Val Thr Phe Val Asp Asn His Asp Thr Gly Tyr Ser Pro
 305 310 315 320
 Gly Gln Asn Gly Gly Gln His His Trp Pro Leu Gln Asp Gly Leu Ile
 325 330 335
 Arg Gln Ala Tyr Ala Tyr Ile Leu Thr Ser Pro Gly Thr Pro Val Val
 340 345 350
 Tyr Trp Ser His Met Tyr Asp Trp Gly Tyr Arg Asp Phe Ile Arg Gln
 355 360 365

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Leu Ile Gln Val Arg Arg Ala Ala Gly Val Arg Ala Asp Ser Ala Ile
 370 375 380
Ser Phe His Ser Gly Tyr Ser Gly Leu Val Ala Thr Val Thr Gly Ser
 385 390 395 400
Gln Gln Thr Leu Val Val Ala Leu Asn Ser Asn Leu Ser Asn Pro Gly
 405 410 415
Gln Val Ala Ser Gly Ser Phe Ser Glu Ala Val Asn Thr Ser Asn Gly
 420 425 430
Gln Val Arg Val Trp Arg Thr Gly Ala Gly Ser Gly Gly Asp Asn
 435 440 445
Gly Gly Gly Glu Pro Gly Ala Leu Val Ser Val Asn Phe Arg Cys Asp
 450 455 460
Asn Gly Val Thr Gln Pro Gly Asp Ser Val Tyr Ala Val Gly Asn Val
 465 470 475 480
Ser Gln Leu Gly Asn Trp Ser Pro Ala Ser Ala Val Arg Leu Thr Asp
 485 490 495
Thr Ser Gly Tyr Pro Thr Trp Lys Gly Arg Ile Ser Leu Pro Ala Gly
 500 505 510
Gln Asn Val Glu Trp Lys Cys Leu Ile Arg Asn Glu Ala Asn Ala Thr
 515 520 525
Gln Val Arg Gln Trp Gln Gly Gly Ala Asn Asn Ser Val Lys Pro Thr
 530 535 540
Glu Gly Ala Asn Thr Ala Gly Arg Phe
 545 550

```

<210> 390
 <211> 1497
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

```

<400> 390
atgaaccgtc actcgacggg tgcctcgacg caccgcgacg agcctcgttt tgcccgcgcg 60
ctggcggagt gcatccacac acgtctggct tccgccatcc gcagcgccgc ccgggcccctg 120
cccggcatcg ccgcgcttgc agccttcgcc ggcagcacga cgcgggcccg ggccgacgtc 180
atctgacag ccttcaactg gcgctacgac actgtcgaag cacgcgccgc cgagatccag 240
tcgctgggct acaaggcggg gctgggtggc ccgcccgtga agtcggaggg ctcggcctgg 300
tgggcgcgct accagccgca ggactaccgc acgatcgacc atccgctggg caacaagcag 360
agcttccagc gcatgtcgaa cgcgctgcgc gcacgcggca tccgcgtcta cgccgacatc 420
atcctcaacc acatggccaa tgaggcggcc cagcgcggcg acctcaacta tcccggccag 480
cgcattctca acgcctacgc tgccaacacc agctactgga acaaccagcg cctggtttggc 540
gacctgcgct acaacttcct gtcgcagtgg gacttcggcc cggccaactg catcagcaac 600
tacaacgacg tctggcaggt gcagaactgg cgcctctgcg gcagcccagg cgacgcgggc 660
ctgcccgacc tgctcggcag cagctatgtg gtcgggcagc agcgcagcta cctgacggca 720
ctcaaggggc tcggcgtcac cggcttccgc atcgatgccg ccaagcacat gccgctctcg 780
cacatcaacg cgggtgctgac acccgagatc aagcaggggc tgcatgtgtt cggcgagggtg 840
atcaccaacg gcggcgccgg cgacagcgag tacaacggct tcctcgcccc ctacttgaac 900
ggcaccgacc acgcggccta cgactttccg ctgttcggca gcatccgcaa cgccttccgc 960
ttcggcggca gcatgaacgc gctgggtcaac ccgctggccg tcggccaggc cctgccccaac 1020
agccgggctg tgacgttcac cgtgaccac gacatcccga acaacggcgg ctccgctac 1080
ctgatcctcg accccaccga cgagaccctg gcctatgcct atgtgctggg ccgcgacggc 1140
ggcagcccgc tgctgtactc ggacaacaac gagagcggcg acaaccgctg ggtcaacgcc 1200
taccgccgca gtgacctgc cgcatgatc cgcttcaca acgcccga gggcagcgac 1260
atgcagggtc tgtcctacag cgactgccac ctgctgttcc gccgcggcaa ccgcggcatc 1320
gtcggtatca acaaatgcgg cagcacggtc aacaccacca tcaacatgaa caacagcgtg 1380
ctctggtggt acaccaatta ccgtgatgtt ctggacgcga acagcgtcgt caacatcggc 1440
agcagcagct acaccttcag cctgccggcg cgcaggcgca ggatgtggtt gaggtag 1497

```

<210> 391
 <211> 498
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(58)

<400> 391

Met	Asn	Arg	His	Ser	Thr	Gly	Ala	Ser	Thr	His	Pro	Gln	Gln	Pro	Arg
1				5					10					15	
Phe	Ala	Arg	Arg	Leu	Ala	Glu	Cys	Ile	His	Thr	Arg	Leu	Ala	Ser	Ala
			20					25					30		
Ile	Arg	Ser	Ala	Ala	Arg	Ala	Leu	Pro	Gly	Ile	Ala	Ala	Leu	Ala	Ala
		35					40					45			
Phe	Ala	Gly	Ser	Thr	Thr	Pro	Ala	Arg	Ala	Asp	Val	Ile	Leu	His	Ala
	50					55				60					
Phe	Asn	Trp	Arg	Tyr	Asp	Thr	Val	Glu	Ala	Arg	Ala	Ala	Glu	Ile	Gln
65					70					75					80
Ser	Leu	Gly	Tyr	Lys	Ala	Val	Leu	Val	Ala	Pro	Pro	Leu	Lys	Ser	Glu
				85					90					95	
Gly	Ser	Ala	Trp	Trp	Ala	Arg	Tyr	Gln	Pro	Gln	Asp	Tyr	Arg	Thr	Ile
			100					105					110		
Asp	His	Pro	Leu	Gly	Asn	Lys	Gln	Ser	Phe	Gln	Arg	Met	Ser	Asn	Ala
		115					120					125			
Leu	Arg	Ala	Arg	Gly	Ile	Arg	Val	Tyr	Ala	Asp	Ile	Ile	Leu	Asn	His
	130					135					140				
Met	Ala	Asn	Glu	Ala	Ala	Gln	Arg	Gly	Asp	Leu	Asn	Tyr	Pro	Gly	Gln
145				150						155					160
Arg	Ile	Leu	Asn	Ala	Tyr	Ala	Ala	Asn	Thr	Ser	Tyr	Trp	Asn	Asn	Gln
				165					170					175	
Arg	Leu	Phe	Gly	Asp	Leu	Arg	Tyr	Asn	Phe	Leu	Ser	Gln	Trp	Asp	Phe
			180					185					190		
Gly	Pro	Ala	Asn	Cys	Ile	Ser	Asn	Tyr	Asn	Asp	Val	Trp	Gln	Val	Gln
		195					200					205			
Asn	Trp	Arg	Leu	Cys	Gly	Ser	Pro	Gly	Asp	Ala	Gly	Leu	Pro	Asp	Leu
	210					215					220				
Leu	Gly	Ser	Ser	Tyr	Val	Val	Gly	Gln	Gln	Arg	Ser	Tyr	Leu	Thr	Ala
225					230					235					240
Leu	Lys	Gly	Leu	Gly	Val	Thr	Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His
				245					250					255	
Met	Pro	Leu	Ser	His	Ile	Asn	Ala	Val	Leu	Thr	Pro	Glu	Ile	Lys	Gln
			260					265					270		
Gly	Val	His	Val	Phe	Gly	Glu	Val	Ile	Thr	Asn	Gly	Gly	Ala	Gly	Asp
		275					280					285			
Ser	Glu	Tyr	Asn	Gly	Phe	Leu	Ala	Pro	Tyr	Leu	Asn	Gly	Thr	Asp	His
	290					295					300				
Ala	Ala	Tyr	Asp	Phe	Pro	Leu	Phe	Gly	Ser	Ile	Arg	Asn	Ala	Phe	Arg
305					310					315					320
Phe	Gly	Gly	Ser	Met	Asn	Ala	Leu	Val	Asn	Pro	Leu	Ala	Val	Gly	Gln
				325					330					335	
Ala	Leu	Pro	Asn	Ser	Arg	Ala	Val	Thr	Phe	Thr	Val	Thr	His	Asp	Ile
			340					345					350		
Pro	Asn	Asn	Gly	Gly	Phe	Arg	Tyr	Leu	Ile	Leu	Asp	Pro	Thr	Asp	Glu
		355					360					365			
Thr	Leu	Ala	Tyr	Ala	Tyr	Val	Leu	Gly	Arg	Asp	Gly	Gly	Ser	Pro	Leu
	370					375					380				
Leu	Tyr	Ser	Asp	Asn	Asn	Glu	Ser	Gly	Asp	Asn	Arg	Trp	Val	Asn	Ala
385				390						395					400
Tyr	Arg	Arg	Ser	Asp	Leu	Ala	Ala	Met	Ile	Arg	Phe	His	Asn	Ala	Ala
				405					410					415	
Gln	Gly	Ser	Asp	Met	Gln	Val	Leu	Ser	Tyr	Ser	Asp	Cys	His	Leu	Leu
			420					425					430		
Phe	Arg	Arg	Gly	Asn	Arg	Gly	Ile	Val	Gly	Ile	Asn	Lys	Cys	Gly	Ser
		435					440					445			
Thr	Val	Asn	Thr	Thr	Ile	Asn	Met	Asn	Asn	Ser	Val	Leu	Trp	Trp	Tyr
	450					455					460				
Thr	Asn	Tyr	Arg	Asp	Val	Leu	Asp	Ala	Asn	Ser	Val	Val	Asn	Ile	Gly
465					470					475					480
Ser	Ser	Ser	Tyr	Thr	Phe	Ser	Leu	Pro	Ala	Arg	Arg	Ala	Arg	Met	Trp
				485					490					495	
Leu	Arg														

<210> 392
 <211> 2100
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 392
 atgccttcag ggctcaagca tggcatcaat atccacgatg acaattctgt cacattcgtc 60
 ctatatgctc aggacaagga caaggccatg catgagtatt gctacctcgt aggtgattgg 120
 aacaactggg agcgactgc cgagagtctg atgaagaggg atacggaggc cggttgctgg 180
 tggctcaagg tcggaggatt tgatccggca aaggagtata gatataattt ccgtctcggc 240
 aacccgcctg catcagacat ctgtgtcagc gatccatata caagaatcgt ctatgatcag 300
 tggaatgacc agtatcttga aggtgttcct gcattccctg agggcgcgaa ggctctcgtt 360
 tctgcattcc agataagcaa accggcctac tcatggaagg ttacggactt caagatagag 420
 gacaggaagg atctcgtgat ctatgagatg ctgttcgcg atttctcagc gacaaaggat 480
 atcgcaggag cggctcgccg gtttgatcac atcaagaacc tcggtgtgaa tgcagtcagg 540
 tcatagccta ttcaggaatt tgacggcaac aacagttggg gatataacct taaccactat 600
 ttcgcccttg acaaggctta tggaaagcgt gacgagtaca agcagttcat tgatctttgt 660
 catgaaaacg gcattgctgt gatcgtggat gtcgtctata accacacctt cggttctcat 720
 acatgggcaa agatgtgggt ggattccgca aacaacagaa ctgcttccaa caatccttgg 780
 tacaatgaat tcccagacaca tccgtacaat gtgggccatg acttcaatca tgagaatgag 840
 atggttgacc agcatgtgaa ggaaagtctt gaattccttc tcagggagta caaggtggac 900
 ggcttcctgt tcgacctcac caagggttcc acccagaaga agacagacc ggatgtttgcg 960
 gcatggggac gttacgacca gagccgtgtg gatatacctca aggactatgc ggaccatatac 1020
 tggctcggta agaatgatgc tgtagtattc ttcgagcatc tttctgactg ggacgaagag 1080
 gaagtctctg caaatcatgg aatccagctt tggagaaatg taaataaaga atatagttcg 1140
 gctgtcaccg gcggaacagg tgatttcaag aatatgcact ctacctcacc gttcggcgga 1200
 tatgtgggat atatggagag ccacgatgag gacggtatat gctatggcgc ttctgacgat 1260
 gtatcttcga tcacatgggg tgtcatcgga cttggaaatg attgggacaa cgacaaggag 1320
 atgacggcag agaacggact gttcgttgta aagaatgtta cggttgccgc caatgacaag 1380
 ttcaagatcc gcaaggccaa ggaatggaat gacaactaca actacggtgc tgccaatgat 1440
 aacttcaaac tgacagtcgg tcagggatac aagatgacca atggaaacag ttcaaaggac 1500
 atggggcgtcc ctggttcagg caagtatgac atctggttca gccatgatat tgccaccgta 1560
 tggcttatgc ctgccggcca gaagccggat tatactccgt cgacttccgt atcatctgag 1620
 gtatcttcga ctgtagccat gcgtcgtgca ggtgcatctg cgccttctt cctcacagt 1680
 cctggacctc agatgatctg gcagttcggg gagatcggat atgatctttc gatcaactat 1740
 ccttccggca ctgaagatga caggacaagc gaaaagccgg taaagaccgc tgaatacatg 1800
 aagaatcctg cacgcaagtc tctctacgac acctatgcag gtctgttgaa gttccgccgc 1860
 gagaaccctc gcttctttga ttctgatgag aagttcagat ggactccttc cggtgagatc 1920
 aagaagatca cctgctctgt ggacggcaag accttccatg tagtgggtaa cttctcgcag 1980
 atggacaagt catattcagt tccgtcaggc agttgggaag attacttcaa tggcggagaa 2040
 tctgtttcgg gaaccattac cctcaagcag ggtgagttca ggctcctgac cagcttctga 2100

<210> 393
 <211> 699
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 393
 Met Pro Ser Gly Leu Lys His Gly Ile Asn Ile His Asp Asp Asn Ser
 1 5 10 15
 Val Thr Phe Val Leu Tyr Ala Gln Asp Lys Asp Lys Ala Met His Glu
 20 25 30
 Tyr Cys Tyr Leu Val Gly Asp Trp Asn Asn Trp Glu Arg Thr Ala Glu
 35 40 45
 Ser Ser Met Lys Arg Asp Thr Glu Ala Gly Cys Trp Trp Leu Lys Val
 50 55 60
 Gly Gly Phe Asp Pro Ala Lys Glu Tyr Arg Tyr Asn Phe Arg Leu Gly
 65 70 75 80
 Asn Pro Pro Ala Ser Asp Ile Cys Val Ser Asp Pro Tyr Thr Arg Ile
 85 90 95
 Val Tyr Asp Gln Trp Asn Asp Gln Tyr Leu Glu Gly Val Pro Ala Phe

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Pro	Glu	Gly	100	Lys	Ala	Leu	Val	105	Ser	Ala	Phe	Gln	Ile	110	Ser	Lys	Pro
Ala	Tyr	Ser	115	Trp	Lys	Val	Thr	120	Asp	Phe	Lys	Ile	Glu	125	Asp	Arg	Lys
Leu	Val	Ile	130	Tyr	Glu	Met	Leu	135	Phe	Arg	Asp	Phe	Ser	140	Ala	Thr	Lys
Ile	Ala	Gly	145	Ala	Val	150	Gly	155	Asp	His	Ile	Lys	Asn	160	Leu	Gly	Val
Asn	Ala	Val	165	Gln	Leu	Met	Pro	170	Gln	Glu	Phe	Asp	Gly	175	Asn	Asn	Ser
Trp	Gly	Tyr	180	Asn	Pro	Asn	His	185	Phe	Ala	Leu	Asp	Lys	190	Ala	Tyr	Gly
Thr	Leu	Asp	195	Glu	Tyr	Lys	Gln	200	Ile	Asp	Leu	Cys	His	205	Glu	Asn	Gly
Ile	Ala	Val	210	Ile	Val	Asp	Val	215	Tyr	Asn	His	Thr	Phe	220	Gly	Ser	His
Thr	Trp	Ala	225	Lys	Met	230	Trp	235	Asp	Ser	Ala	Asn	Asn	240	Arg	Thr	Ala
Asn	Asn	Pro	245	Trp	Tyr	Asn	Glu	250	Phe	Pro	Thr	His	Pro	255	Tyr	Asn	Val
His	Asp	Phe	260	Asn	His	Glu	Asn	265	Glu	Met	Val	Asp	Gln	270	His	Val	Lys
Ser	Leu	Glu	275	Phe	Leu	Leu	Arg	280	Glu	Tyr	Lys	Val	Asp	285	Gly	Phe	Arg
Asp	Leu	Thr	290	Lys	Gly	Phe	Thr	295	Gln	Lys	Lys	Thr	Asp	300	Pro	Asp	Val
Ala	Trp	Gly	305	Arg	Tyr	Asp	Gln	310	Ser	Arg	Val	Asp	Ile	315	Leu	Lys	Asp
Ala	Asp	His	325	Ile	Trp	Ser	Val	330	Lys	Asn	Asp	Ala	Val	335	Val	Ile	Phe
His	Leu	Ser	340	Asp	Trp	Asp	Glu	345	Glu	Val	Leu	Ala	Asn	350	His	Gly	Ile
Gln	Leu	Trp	355	Arg	Asn	Val	Asn	360	Lys	Glu	Tyr	Ser	Ser	365	Ala	Val	Thr
Gly	Thr	Gly	370	Asp	Phe	Lys	Asn	375	Met	His	Ser	Thr	Ser	380	Pro	Phe	Gly
Tyr	Val	Gly	385	Tyr	Met	390	Glu	395	Ser	His	Asp	Glu	Arg	400	Ile	Cys	Tyr
Ala	Ser	Asp	405	Val	Ser	Ser	Ile	410	Thr	Trp	Gly	Val	Ile	415	Gly	Leu	Gly
Asn	Asp	Trp	420	Asp	Asn	Asp	Lys	425	Met	Thr	Ala	Glu	Asn	430	Gly	Leu	Phe
Val	Val	Lys	435	Asn	Val	Thr	Val	440	Ala	Asn	Asp	Lys	Phe	445	Lys	Ile	Arg
Lys	Ala	Lys	450	Glu	Trp	Asn	Asp	455	Asn	Tyr	Asn	Tyr	Gly	460	Ala	Ala	Asn
Asn	Phe	Lys	465	Leu	Thr	Val	Gly	470	Gln	Gly	Tyr	Lys	Met	475	Thr	Asn	Gly
Ser	Ser	Lys	485	Met	Gly	Val	Pro	490	Val	Ala	Gly	Lys	Tyr	495	Asp	Ile	Trp
Phe	Ser	His	500	Asp	Ile	Ala	Thr	505	Trp	Leu	Met	Pro	Ala	510	Gly	Gln	Lys
Pro	Asp	Tyr	515	Thr	Pro	Ser	Thr	520	Val	Ser	Ser	Ser	Glu	525	Asp	Ala	Leu
Val	Ala	Met	530	Arg	Arg	Ala	Gly	535	Ala	Ser	Ala	Ala	Phe	540	Phe	Leu	Thr
Pro	Gly	Pro	545	Lys	Met	Ile	Trp	550	Gln	Phe	Gly	Glu	Ile	555	Gly	Tyr	Asp
Ser	Ile	Asn	565	Tyr	Pro	Ser	Gly	570	Thr	Glu	Asp	Asp	Arg	575	Thr	Ser	Glu
Pro	Val	Lys	580	Thr	Ala	Glu	Tyr	585	Met	Lys	Asn	Pro	Ala	590	Arg	Lys	Ser
Tyr	Asp	Thr	595	Tyr	Ala	Gly	Leu	600	Leu	Lys	Phe	Arg	Arg	605	Glu	Asn	Pro
Phe	Phe	Asp	610	Ser	Asp	Ala	Lys	615	Phe	Glu	Trp	Thr	Pro	620	Ser	Gly	Glu
Lys	Lys	Ile	625	Thr	Cys	Ser	Val	630	Asp	Gly	Lys	Thr	Phe	635	His	Val	Val

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Asn Phe Ser Gln 645 Met Asp Lys Ser Tyr 650 Ser Val Pro Ser Gly 655
660
Lys Asp Tyr Phe Asn Gly Gly Glu Ser Val Ser Gly Thr Ile Thr Leu
675
Lys Gln Gly Glu Phe Arg Leu Leu Thr Ser Phe
690 680 685 695

<210> 394
<211> 1347
<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 394
atgcggacat cagtgtttgtg cggagtggtc ctgatccaac aggcccgctcg tttcgcttcg 60
agaaaatcag ttcatatcca cgccagcagc acctgcagtc gggtgcggcg ctggcacatc 120
gtcgccctcg tgctcgccag tacctggctg gcgaccgagc cggcgcaagc ggccacggaa 180
cagaatcaag ccagcagcgc gatactgttc cagggttttc actggcactc cgccaacggt 240
agctggtacg gcaatctgca gggcaaggct ggcgatttga aagacttggg catcactcac 300
gtctggttcc cgccgccgtc ggacgctgcg tcaactggaag ggtatctgcc gcgccagctc 360
aatgtgctga acagcagtta cggcagtgag acggcgctac gcaatgccat cagcgcgctg 420
accggccaag gcatcaagag cgtggccgat gtcgtgatca atcaccgcgt tggcaccagc 480
aacggggccg acttcaccaa tccgagctgg gattgccgcg ctgtggtcaa caatgacgaa 540
tggagcggcg gttgcggcaa cggcgattcc ggtgacggct atggcgctgc ccgcgatctg 600
gatcactcgc aaggctttgt tcagaacgat ctgaaaacct ggctgtcgtc gcgactcaaa 660
ggcgttggtt tttccggtat tcgttacgac tactcgaaag gctacggttc cggttacgcc 720
ggcctgtatc acgacgcaat gaatccgaac ttctgcgtcg gtgagatctg gaccgatctg 780
gattacaaca atgtcgatgc gcacgcgccag ctgttgatga attatgtcga cggcaacaat 840
ggcaaatgcg gcgccttcga tttcaccagc aaaggctctg tcaatcaggc actgtgggcc 900
aacgaatatt ggcgactcgc cgccctgggat ggcaaaccgg ctggtggcat cggttggtgg 960
gcgcagaaga tggtagcgtt cgtcgacaac catgacaccg gtccgctcga gagttgcagt 1020
gccggccaga accactggcc ggtgccgtgc ggcaaagtga tgcagggtta tgcctacatc 1080
ctgagtcatc ccggtattcc gaccgtgtat taccgcacg tctatgactg gaacctgcgc 1140
agtgccatca agaccctgat cagcctgcgc aaggagaagg gcatcacctc gaccagcagc 1200
gtcgcgatcc agcgcgccga tcaaggcctc tatgccgcca tcatcaacgg caatctggcg 1260
atgaagatcg gcccgaacag ctggagtccc ggcagcggct ggacgctgcg catgtctggc 1320
gaacagtacg caatttgac ccgttga 1347

<210> 395
<211> 448
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(57)

<400> 395
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Arg Phe Ala Ser Arg Lys Ser Val His Ile His Ala Ser Ser Thr Cys
20 25 30
Ser Arg Leu Arg Arg Trp His Ile Val Ala Ser Leu Leu Ala Ser Thr
35 40 45
Trp Leu Ala Thr Glu Pro Ala Gln Ala Ala Thr Glu Gln Asn Gln Ala
50 55 60
Ser Ser Ala Ile Leu Phe Gln Gly Phe His Trp His Ser Ala Asn Gly
65 70 75 80
Ser Trp Tyr Gly Asn Leu Gln Gly Lys Ala Gly Asp Leu Lys Asp Leu
85 90 95
Gly Ile Thr His Val Trp Phe Pro Pro Ser Asp Ala Ala Ser Leu
100 105 110
Glu Gly Tyr Leu Pro Arg Gln Leu Asn Val Leu Asn Ser Ser Tyr Gly

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115      120      125
Ser Glu Thr Ala Leu Arg Asn Ala Ile Ser Ala Leu Thr Gly Gln Gly
130      135      140
Ile Lys Ser Val Ala Asp Val Val Ile Asn His Arg Val Gly Thr Ser
145      150      155
Asn Trp Ala Asp Phe Thr Asn Pro Ser Trp Asp Cys Arg Ala Val Val
165
Asn Asn Asp Glu Trp Ser Gly Arg Cys Gly Asn Gly Asp Ser Gly Asp
180      185      190
Gly Tyr Gly Ala Ala Arg Asp Leu Asp His Ser Gln Gly Phe Val Gln
195      200      205
Asn Asp Leu Lys Thr Trp Leu Ser Ser Arg Leu Lys Gly Val Gly Phe
210      215      220
Ser Gly Ile Arg Tyr Asp Tyr Ser Lys Gly Tyr Gly Ser Gly Tyr Ala
225      230      235
Gly Leu Tyr His Asp Ala Met Asn Pro Asn Phe Cys Val Gly Glu Ile
245      250      255
Trp Thr Asp Leu Asp Tyr Asn Asn Val Asp Ala His Arg Gln Leu Leu
260      265      270
Met Asn Tyr Val Asp Gly Asn Asn Gly Lys Cys Gly Ala Phe Asp Phe
275      280      285
Thr Ser Lys Gly Leu Leu Asn Gln Ala Leu Trp Ala Asn Glu Tyr Trp
290      295      300
Arg Leu Arg Ala Trp Asp Gly Lys Pro Ala Gly Gly Ile Gly Trp Trp
305      310      315
Ala Gln Lys Met Val Thr Phe Val Asp Asn His Asp Thr Gly Pro Ser
325      330      335
Gln Ser Cys Ser Ala Gly Gln Asn His Trp Pro Val Pro Cys Gly Lys
340      345      350
Val Met Gln Gly Tyr Ala Tyr Ile Leu Ser His Pro Gly Ile Pro Thr
355      360      365
Val Tyr Tyr Pro His Val Tyr Asp Trp Asn Leu Arg Ser Ala Ile Lys
370      375      380
Thr Leu Ile Ser Leu Arg Lys Glu Lys Gly Ile Thr Ser Thr Ser Ser
385      390      395
Val Ala Ile Gln Arg Ala Asp Gln Gly Leu Tyr Ala Ala Ile Ile Asn
400      405      410
Gly Asn Leu Ala Met Lys Ile Gly Pro Asn Ser Trp Ser Pro Gly Ser
420      425      430
Gly Trp Thr Leu Arg Met Ser Gly Glu Gln Tyr Ala Ile Trp Thr Arg
435      440      445

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<210> 396

<211> 1644

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 396

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aaaaacatga ccgaacaagc accttttgat tggaaagcag ccaatgtgta ttttttactc      120
accgatcgct ttgcaaacgg agaccctagc aatgacaacc tcatcaaaaag agatttgcca      180
acaggggttt tacgtgggtt tatgggagga gatttaaaag gcatcaccca aaaaatagaa      240
gaagggttatt ttaccaactt aggcattcaat gccatttgga tgactccgat tgtagaacia      300
attcacggtg cagtggatga aggcaccgga aacacctatg gttttcatgg ctattggacc      360
aaagattgga cagctattga ccctaattat ggcaccaaag ccgattttaa agaattggta      420
gaaaaagccc atgccaaagg cattcgaatt gtattagatg cagtaatcaa ccacactggg      480
ccgggttaccg aattagatcc ggtgttccca gatagtggg ttagaaccaa acctaaatgt      540
acttacgaca cgtataaaaa ttatatcgat tgtacgttag tagaaaatct tcccacatc      600
aaaaccgaaa gcaatgaaga agtagaattg ccaccaatgt tagtggaata gtggaaaaaa      660
gaaggcagat atgaagaaga aataagagaa ttagatgctt tttttgcaa aaccggttat      720
ccaagagcac caaaatacta catcatgaaa tggttggcag attacatcac cgactttggg      780
attgatgggtt acagagtggg tacggttaaa cacaccacag aagatgtttg ggcagatttt      840
gcaaaagtgt gtaaagatgc ttttacagct tataaaactg caaatcctaa aaaagtgtta      900
gacgataacg acttcctttt ggttggaaga gtatatggtt atggtttgca tggcaaaaaa      960
atctatgatt ttggagataa aaaagtgaat tattttgaaa acggattcaa ctttttaatc      1020

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aatttttgatt	ttaaaggaga	tgccaacaaa	gggtatgaag	aattgtttag	ctattacgat	1080
aaagttttgc	aaaacgaaat	gggcaactat	agtgtgatga	attacttaac	ttctcatgac	1140
gatggttggc	cttttgacaa	aaaaagagaa	cgtacttttg	aagcagctac	caagttactg	1200
ctttctcctg	gagtggcaca	agtatattat	ggagatgaaa	ctgcaagatc	tttagaaatt	1260
gaaggcacc	aaggagatgc	cactttgcca	agttttatga	attgggaaga	ggttcaaagt	1320
aatgctaaaa	ccaaagcatt	attagttcat	tatcaaaagt	tagggtcatt	tagagcaaac	1380
cacccggcag	ttggagcagg	taaacacgaa	aaaattagcg	atgcacctta	tactttttcc	1440
agaacttatc	aaaaaggtga	tttcaaagat	gtcatagtaa	ttactttaga	cgcttcaaaa	1500
ggtgccaaaa	ccatccaagt	aaataatgca	tttgagagg	gcagtaaagt	tagagatgct	1560
tattcaggaa	tgattggaga	agtaaaaaat	ggtaaagtag	agatccattc	taacgaaact	1620
atcgtgttgt	tagaaaaaat	gtaa				1644

<210> 397
 <211> 547
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(19)

<400> 397

Met	Lys	Lys	Thr	Ile	Leu	Leu	Thr	Thr	Ile	Cys	Ile	Leu	Leu	Trp	Asn
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Cys	Gln	Ser	Lys	Lys	Asn	Met	Thr	Glu	Gln	Ala	Pro	Phe	Asp	Trp	Lys
			20					25					30		
Ala	Ala	Asn	Val	Tyr	Phe	Leu	Leu	Thr	Asp	Arg	Phe	Ala	Asn	Gly	Asp
		35					40					45			
Pro	Ser	Asn	Asp	Asn	Leu	Ile	Lys	Arg	Asp	Leu	Pro	Thr	Gly	Val	Leu
	50					55					60				
Arg	Gly	Phe	Met	Gly	Gly	Asp	Leu	Lys	Gly	Ile	Thr	Gln	Lys	Ile	Glu
65				70					75					80	
Glu	Gly	Tyr	Phe	Thr	Asn	Leu	Gly	Ile	Asn	Ala	Ile	Trp	Met	Thr	Pro
				85				90						95	
Ile	Val	Glu	Gln	Ile	His	Gly	Ala	Val	Asp	Glu	Gly	Thr	Gly	Asn	Thr
			100					105					110		
Tyr	Gly	Phe	His	Gly	Tyr	Trp	Thr	Lys	Asp	Trp	Thr	Ala	Ile	Asp	Pro
		115					120					125			
Asn	Tyr	Gly	Thr	Lys	Ala	Asp	Leu	Lys	Glu	Leu	Val	Glu	Lys	Ala	His
	130					135					140				
Ala	Lys	Gly	Ile	Arg	Ile	Val	Leu	Asp	Ala	Val	Ile	Asn	His	Thr	Gly
145					150					155					160
Pro	Val	Thr	Glu	Leu	Asp	Pro	Val	Phe	Pro	Asp	Ser	Trp	Val	Arg	Thr
				165					170					175	
Lys	Pro	Lys	Cys	Thr	Tyr	Asp	Thr	Tyr	Lys	Asn	Tyr	Ile	Asp	Cys	Thr
			180					185					190		
Leu	Val	Glu	Asn	Leu	Pro	Asp	Ile	Lys	Thr	Glu	Ser	Asn	Glu	Glu	Val
		195					200					205			
Glu	Leu	Pro	Pro	Met	Leu	Val	Glu	Lys	Trp	Lys	Lys	Glu	Gly	Arg	Tyr
	210					215					220				
Glu	Glu	Glu	Ile	Arg	Glu	Leu	Asp	Ala	Phe	Phe	Ala	Lys	Thr	Gly	Tyr
225					230					235					240
Pro	Arg	Ala	Pro	Lys	Tyr	Tyr	Ile	Met	Lys	Trp	Leu	Ala	Asp	Tyr	Ile
				245					250					255	
Thr	Asp	Phe	Gly	Ile	Asp	Gly	Tyr	Arg	Val	Asp	Thr	Val	Lys	His	Thr
			260					265					270		
Thr	Glu	Asp	Val	Trp	Ala	Asp	Phe	Ala	Lys	Val	Cys	Lys	Asp	Ala	Phe
		275					280					285			
Thr	Ala	Tyr	Lys	Thr	Ala	Asn	Pro	Lys	Lys	Val	Leu	Asp	Asp	Asn	Asp
	290					295					300				
Phe	Phe	Leu	Val	Gly	Glu	Val	Tyr	Gly	Tyr	Gly	Leu	His	Gly	Lys	Gln
305					310					315					320
Ile	Tyr	Asp	Phe	Gly	Asp	Lys	Lys	Val	Asn	Tyr	Phe	Glu	Asn	Gly	Phe
				325					330					335	
Asn	Phe	Leu	Ile	Asn	Phe	Asp	Phe	Lys	Gly	Asp	Ala	Asn	Lys	Gly	Tyr
			340					345					350		

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Glu Glu Leu Phe Ser Tyr Tyr Asp Lys Val Leu Gln Asn Glu Met Gly
 355 360 365
 Asn Tyr Ser Val Met Asn Tyr Thr Ser His Asp Asp Gly Trp Pro
 370 375 380
 Phe Asp Lys Lys Arg Glu Arg Thr Phe Glu Ala Ala Thr Lys Leu Leu
 385 390 395 400
 Leu Ser Pro Gly Val Ala Gln Val Tyr Tyr Gly Asp Glu Thr Ala Arg
 405 410 415
 Ser Leu Glu Ile Glu Gly Thr Gln Gly Asp Ala Thr Leu Arg Ser Phe
 420 425 430
 Met Asn Trp Glu Glu Val Gln Ser Asn Ala Lys Thr Lys Ala Leu Leu
 435 440 445
 Val His Tyr Gln Lys Leu Gly Ser Phe Arg Ala Asn His Pro Ala Val
 450 455 460
 Gly Ala Gly Lys His Glu Lys Ile Ser Asp Ala Pro Tyr Thr Phe Ser
 465 470 475 480
 Arg Thr Tyr Gln Lys Gly Asp Phe Lys Asp Val Ile Val Ile Thr Leu
 485 490 495
 Asp Ala Ser Lys Gly Ala Lys Thr Ile Gln Val Asn Asn Ala Phe Ala
 500 505 510
 Glu Gly Ser Lys Val Arg Asp Ala Tyr Ser Gly Met Ile Gly Glu Val
 515 520 525
 Lys Asn Gly Lys Val Glu Ile His Ser Asn Glu Thr Ile Val Leu Leu
 530 535 540
 Glu Lys Met
 545

<210> 398
 <211> 2040
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 398
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 gtaggtgact ggaacaactg gacacgcgag acagagggag ccatgtacta cgatggaacc 120
 aagggctgct ggtggatcac tcttgacggc ttcgacgctg acaaggagta ccgcttcacg 180
 tatcgtctcg gcaattcctc aggtactgac acctatgtca gcgacccata cacagagatc 240
 gtatatgacc agtggaaatga ccagtatatt cagggtgttc ctgcattccc tgaaggagca 300
 aaggccctcg tttcggcatt ccagatcaat agacctgaat acagctggaa gcacaaggac 360
 ttcaaggtag aggacaagaa cgacctcggt atctacgaga tgctcttccg cgacttcgct 420
 acatctcaga acattgccgg tgcgatgacg cagctcgact atatcgagaa cctcgggtgtc 480
 actgcagtcc agctcatgcc gatccaggag ttcgacggca acctcagctg gggatataac 540
 cctaaccact ggttcgccct cgacaagtac tacggtacac gtgagcagta caaggagtcc 600
 atcgacgagt gccacgctcg cggaatggct gtgatcatcg atgtcgtata taaccacgcc 660
 acaggttcgc atccttgggc gaagatgttg tgggacgggtg acaggactgc cgcgaacaac 720
 ccttggttca acgagtacgc aaagcaccgg tacaacgtat atcatgacat gaatcatgag 780
 aatgcgatgg tcaaggagca tgtgaagaag agcctcgagt atctcctgac ggagtacgac 840
 gtggacggat tccgtttcga cctcacgaaa ggcttcaccc agaaggatac cggcggagac 900
 aacggcgacg tggcagcgtg gggacgttat gaccagagcc gtgtggacat cctcaaggga 960
 tatgccgacc acatctgggtc tgtgaacagt gatgctgtag tcatcttcga gcatctttcc 1020
 gactgggatg aggagaaagt tctcgctgag catggcatcc agctttggag aaatgtgaac 1080
 ggagaatatc gttcagcggg gaccggcgga acaggaacct tcacgaatat gtattccaac 1140
 gctccgttcg gcggattcgt gggatatatg gagagtcacg atgaggagcg tatctgttac 1200
 ggtgccgggtg ctgatgcata ttctgtgaca tggggtgtga tcggtctcgg tgacgactgg 1260
 aacaatgaca aggtaatgtc taaggacggg gtcttcttct cagtgaagaa tgtgaccgtg 1320
 actgcgaacg accgtttcaa gatccgcaag gcaggcgagt ggaacgacgc atttaactac 1380
 ggtgcatctg gcgacaactt caagctcacc gtgggtcagg gctacaagat gaccaacggt 1440
 aacagctcga aggatattga tgtaccggct gcaggaacct acgatattcta cttcagcctc 1500
 gagactgaga cagtattggc catggctgcc ggtcagcgcc ctgccgatcc ttcgacaggc 1560
 ggccggagctt ccgaggatgc ccttacgggt gcaatgcgtc gtgcagggtgc ttctgcagca 1620
 ttcttcctca ctgttcctgg tccgaagatg atctggcagt tcggtgagat cggatacgac 1680
 tattctatcg agtacaacga ccgcacaggt gagaagcctg tcgtgaccga ccagtacatg 1740
 gcagtccttc cccgcaaggc tctctatgac acctatgcgg ctctcctgaa gttccgctgt 1800
 gacaacccct gcttcttcga caaggatgcg gaactccttc aggcgagatc 1860
 aagaagatca cctgctctgt ggacggaaag accttccatg tgggtgggtaa cttcgcgaag 1920

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tcttcaaaga catacaccgt tccttcaggc agctggaagg actacttcaa taacggagca 1980
tcggtttcag gaaccatcac cctcaagcag ggcgagttca gactcctgac cagcttctaa 2040

<210> 399
<211> 679
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

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Met Thr Phe Val Phe Tyr Asp Gln Asp Thr Ala Gly Lys Ser His Lys
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Tyr Cys Tyr Ile Val Gly Asp Trp Asn Asn Trp Thr Arg Glu Thr Glu
20 25 30
Gly Ala Met Tyr Tyr Asp Gly Thr Lys Gly Cys Trp Trp Ile Thr Leu
35 40 45
Asp Gly Phe Asp Ala Asp Lys Glu Tyr Arg Phe Gln Tyr Arg Leu Gly
50 55 60
Asn Ser Ser Gly Thr Asp Thr Tyr Val Ser Asp Pro Tyr Thr Glu Ile
65 70 75 80
Val Tyr Asp Gln Trp Asn Asp Gln Tyr Ile Gln Gly Val Pro Ala Phe
85 90 95
Pro Glu Gly Ala Lys Ala Leu Val Ser Ala Phe Gln Ile Asn Arg Pro
100 105 110
Glu Tyr Ser Trp Lys His Lys Asp Phe Lys Val Glu Asp Lys Asn Asp
115 120 125
Leu Val Ile Tyr Glu Met Leu Phe Arg Asp Phe Ala Thr Ser Gln Asn
130 135 140
Ile Ala Gly Ala Met Thr Gln Leu Asp Tyr Ile Glu Asn Leu Gly Val
145 150 155 160
Thr Ala Val Gln Leu Met Pro Ile Gln Glu Phe Asp Gly Asn Leu Ser
165 170 175
Trp Gly Tyr Asn Pro Asn His Trp Phe Ala Leu Asp Lys Tyr Tyr Gly
180 185 190
Thr Arg Glu Gln Tyr Lys Glu Phe Ile Asp Glu Cys His Ala Arg Gly
195 200 205
Met Ala Val Ile Ile Asp Val Val Tyr Asn His Ala Thr Gly Ser His
210 215 220
Pro Trp Ala Lys Met Trp Trp Asp Gly Asp Arg Thr Ala Ala Asn Asn
225 230 235 240
Pro Trp Phe Asn Glu Tyr Ala Lys His Pro Tyr Asn Val Tyr His Asp
245 250 255
Met Asn His Glu Asn Ala Met Val Lys Glu His Val Lys Lys Ser Leu
260 265 270
Glu Tyr Leu Leu Thr Glu Tyr Asp Val Asp Gly Phe Arg Phe Asp Leu
275 280 285
Thr Lys Gly Phe Thr Gln Lys Asp Thr Gly Gly Asp Asn Gly Asp Val
290 295 300
Ala Ala Trp Gly Arg Tyr Asp Gln Ser Arg Val Asp Ile Leu Lys Gly
305 310 315 320
Tyr Ala Asp His Ile Trp Ser Val Asn Ser Asp Ala Val Val Ile Phe
325 330 335
Glu His Leu Ser Asp Trp Asp Glu Glu Lys Val Leu Ala Glu His Gly
340 345 350
Ile Gln Leu Trp Arg Asn Val Asn Gly Glu Tyr Arg Ser Ala Val Thr
355 360 365
Gly Gly Thr Gly Asn Phe Thr Asn Met Tyr Ser Asn Ala Pro Phe Gly
370 375 380
Gly Phe Val Gly Tyr Met Glu Ser His Asp Glu Glu Arg Ile Cys Tyr
385 390 395 400
Gly Ala Gly Ala Asp Ala Ser Ser Val Thr Trp Gly Val Ile Gly Leu
405 410 415
Gly Asp Asp Trp Asn Asn Asp Lys Val Met Ser Lys Asp Gly Val Phe
420 425 430
Phe Ser Val Lys Asn Val Thr Val Thr Ala Asn Asp Arg Phe Lys Ile
435 440 445

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Arg Lys Ala Gly Glu Trp Asn Asp Ala Phe Asn Tyr Gly Ala Ser Gly
 450 455 460
 Asp Asn Phe Lys Leu Thr Val Gly Gln Gly Tyr Lys Met Thr Asn Gly
 465 470 475 480
 Asn Ser Ser Lys Asp Met Tyr Val Pro Ala Ala Gly Thr Tyr Asp Ile
 485 490 495
 Tyr Phe Ser Leu Glu Thr Glu Thr Val Trp Leu Met Ala Ala Gly Gln
 500 505 510
 Arg Pro Ala Asp Pro Ser Thr Gly Gly Ala Ser Glu Asp Ala Leu
 515 520 525
 Thr Val Ala Met Arg Arg Ala Gly Ala Ser Ala Ala Phe Phe Leu Thr
 530 535 540
 Val Pro Gly Pro Lys Met Ile Trp Gln Phe Gly Glu Ile Gly Tyr Asp
 545 550 555 560
 Tyr Ser Ile Glu Tyr Asn Asp Arg Thr Gly Glu Lys Pro Val Val Thr
 565 570 575
 Asp Gln Tyr Met Ala Val Pro Ala Arg Lys Ala Leu Tyr Asp Thr Tyr
 580 585 590
 Ala Ala Leu Lys Phe Arg Arg Asp Asn Pro Arg Phe Phe Asp Lys
 595 600 605
 Asp Ala Lys Phe Glu Trp Thr Pro Ser Gly Glu Ile Lys Lys Ile Thr
 610 615 620
 Cys Ser Val Asp Gly Lys Thr Phe His Val Val Gly Asn Phe Ala Lys
 625 630 635 640
 Ser Ser Lys Thr Tyr Thr Val Pro Ser Gly Ser Trp Lys Asp Tyr Phe
 645 650 655
 Asn Asn Gly Ala Ser Val Ser Gly Thr Ile Thr Leu Lys Gln Gly Glu
 660 665 670
 Phe Arg Leu Leu Thr Ser Phe
 675

<210> 400
 <211> 1245
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<400> 400
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 actgagcaaa acaaagcttc gacagccatc ctgctccagg gctttcactg gaattcggcg 120
 aacgggtcct ggtacctgaa cctcgaaggc aaggccgcgg acttcggga tcttggcatc 180
 acgcacgtct ggtttcacc gccatcggac tcgggagcgc gcgaaggcta tctgcctcgg 240
 gagctttata aacttcagtc ggcctatggc agcgaggagc agctccgcaa tgccatcagc 300
 gctctgaatc gggaaggcat ccaggcgggtg gccgatatcg taatcaacca tcgcgtcggc 360
 agctggaact ggggagactt taactaccct aaatgggact gcgatagcgt cgcgaacaac 420
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 ggcttgaatt atgatgatgt caacgcgcgt cgccaggagc tgatgaatta tgtcgacggc 720
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 cgtgacaatg actactggcg cctgcgcgat gccaatggac gtccggccgg tggcatcggc 840
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 tgttcagcag gacagaacat gtggccggtt ccctgtgaca aggtgatgct gggctatgcc 960
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<210> 401
 <211> 414
 <212> PRT
 <213> Unknown

<220>

1001827087_1.txt

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(19)

<400> 401

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Met Lys Asn Phe Ile Leu Gly Met Ala Ala Met Leu Met Ala Ala Gln
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Ser Leu Ala Ala Thr Glu Gln Asn Lys Ala Ser Thr Ala Ile Leu Leu
      20      25      30
Gln Gly Phe His Trp Asn Ser Ala Asn Gly Ser Trp Tyr Leu Asn Leu
      35      40      45
Glu Gly Lys Ala Ala Asp Phe Arg Asp Leu Gly Ile Thr His Val Trp
 50      55      60
Phe Pro Pro Pro Ser Asp Ser Gly Ala Arg Glu Gly Tyr Leu Pro Arg
 65      70      75      80
Glu Leu Tyr Lys Leu Gln Ser Ala Tyr Gly Ser Glu Glu Gln Leu Arg
      85      90      95
Asn Ala Ile Ser Ala Leu Asn Arg Glu Gly Ile Gln Ala Val Ala Asp
      100      105      110
Ile Val Ile Asn His Arg Val Gly Ser Trp Asn Trp Gly Asp Phe Asn
      115      120      125
Tyr Pro Lys Trp Asp Cys Asp Ser Val Ala Asn Asn Asp Glu Trp Pro
      130      135      140
Gly Arg Cys Gly Gly Tyr Asp Ser Gly Ala Ser Tyr Gly Ala Ala Arg
      145      150      155      160
Asp Ile Asp His Ser Asn Ala Gln Val Gln Asn Asp Ile Lys Tyr Phe
      165      170      175
Leu Ser Glu Arg Leu Arg Gly Val Gly Phe Ser Gly Trp Arg Tyr Asp
      180      185      190
Tyr Ser Lys Gly Tyr Ala Pro Tyr Tyr Ala Arg Ile Tyr Asn Glu Ala
      195      200      205
Ser Arg Pro Asp Phe Cys Val Gly Glu Val Trp Pro Gly Leu Asn Tyr
      210      215      220
Asp Asp Val Asn Ala His Arg Gln Glu Leu Met Asn Tyr Val Asp Gly
      225      230      235      240
Thr Gly Gly Ser Cys Gly Ala Phe Asp Phe Thr Ser Lys Gly Leu Leu
      245      250      255
Asn Lys Val Leu Arg Asp Asn Asp Tyr Trp Arg Leu Arg Asp Ala Asn
      260      265      270
Gly Arg Pro Ala Gly Gly Ile Gly Trp Trp Pro Gln Lys Met Val Thr
      275      280      285
Phe Val Asp Asn His Asp Thr Gly Pro Ser Glu Ser Cys Ser Ala Gly
      290      295      300
Gln Asn Met Trp Pro Val Pro Cys Asp Lys Val Met Leu Gly Tyr Ala
      305      310      315      320
Tyr Ile Leu Thr His Pro Gly Val Pro Ser Leu Tyr Tyr Pro His Leu
      325      330      335
Tyr Asp Trp Gly Leu Lys Asn Ser Ile Lys Ser Leu Leu Asn Ala Arg
      340      345      350
Arg Ser Val Gly Ile Gln Ser Asn Ser Asn Ile Ser Ile Gln Lys Ala
      355      360      365
Glu Asn Gly Leu Tyr Ala Ala Ile Ile Ser Gly Arg Ser Gly Arg Leu
      370      375      380
Ala Met Lys Ile Gly Ser Gln Asp Trp Asn Pro Gly Ala Gly Trp Asp
      385      390      395      400
Leu Val Thr Trp Gly Asn Gln Tyr Ala Val Trp Ser Gln Lys
      405      410

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<210> 402

<211> 1995

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 402

1001827087_1.txt

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caaaaaacta ctttcgtgca tttattcagag tggcgctggg ctgatattgc caccgaatgc 120
gaaacatttt taggcccgca gggccttggct gctatccagg tgtcgctcc gacagagcat 180
attcagggct cggcctgggt gacccgctat caaccgggtca gctatgtggt acaaagccgc 240
agtggtaaca gagccgagtt tatctctatg gtgcaacgtt gtaaggcggc aggcgtagat 300
atttatgtcg acgctgtcat caaccatatg gccagcggca gtggtaccgg cgttgctggc 360
agcagttatg gtaacaggca gtaccccatc tacagcagcc ctgattttca taactcctgc 420
accattaaca gcagcgacta tggcaacgac cgctggcggg tgcaaaactg cgaactgggt 480
ggtttaccgg atttaaatat cagcgccagc tatgtacaaa atagcctcgc cgcttatctg 540
aacgatatgc tggcgatagg ggtaactggg tttcgtattg acgcccga gcatatggcc 600
agttcagata ttgcaggcat caaagccaaa ctacgagag caccactgat ttaccaggag 660
gtgatcgatc agggcggcga ggtgatcacc gccagcgagt acacaggcaa tggcatgggt 720
actgaattta agtattcggg gcagctgggt aatgtgttta aaaccggccg gctggccagc 780
ctgtctaact ttggtgaagc ctgggggttt ttacctggca atcaggcggg ggtttttgta 840
gataaccacg acaatcagcg cggtcatggc ggtgcaggtg atgtcgtgac ctacaaagat 900
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atagcgggcg cagtgcagtt ccgcaacaat acggcggagg aatggcgggt gacccattgg 1140
tgggataacg gcaataacca gatcgcttc ggccgtgctg ctttaggttt tgcgctatc 1200
aataaaagaa gttacaacct gagcacatca ttacaaaccg gtatgacagc aggcagttac 1260
tgcaatgtgc tcaaaggcag catcagcacc gataaaacca gttgttcagg cgaagtgatc 1320
cagataggca gcgatggccg tattcaggcc aatctggcat cctgggatgc ttttgccatt 1380
catcatcagg cgaaaatttg cggcacagtg cagcctcctg caggcaactg gcagcgctact 1440
gtggtgttta ttcaggccca aactcaaact ggccaggata tgtttgtccg cgggtgggctg 1500
gatcatgcag ttgcgcaaag ccagctcggc ttaagctgca gcagtagcaa ttatttgtgc 1560
gccattccta tacggcatcg taatctgaac aacgccacta cagcaccatg gaaagctaac 1620
gatggcttgc tcgactggta tggccgcgaa gccagccaaa gcagcgccgc tgaaggcaca 1680
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gtcgatatgg attgcagcaa aaccatcaat ggctggtttg aactcaaagc ttatgtcaaa 1860
aacggccagg gctgggaagg taatatcagc cagagcggca cgccttacgc cagtcagaac 1920
catatggcgc aatgcggtca gatgaatgtg tttagctttg gtcaaaacag cgcacaaatc 1980
agcgccctgc cctag 1995

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<210> 403

<211> 664

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(19)

<400> 403

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Ala Leu Ala Ala Gln Lys Thr Thr Phe Val His Leu Phe Glu Trp Arg
20      25      30
Trp Ser Asp Ile Ala Thr Glu Cys Glu Thr Phe Leu Gly Pro Gln Gly
35      40      45
Phe Ala Ala Ile Gln Val Ser Pro Pro Thr Glu His Ile Gln Gly Ser
50      55      60
Ala Trp Trp Thr Arg Tyr Gln Pro Val Ser Tyr Val Val Gln Ser Arg
65      70      75      80
Ser Gly Asn Arg Ala Glu Phe Ile Ser Met Val Gln Arg Cys Lys Ala
85      90      95
Ala Gly Val Asp Ile Tyr Val Asp Ala Val Ile Asn His Met Ala Ser
100     105     110
Gly Ser Gly Thr Gly Val Ala Gly Ser Ser Tyr Gly Asn Arg Gln Tyr
115     120     125
Pro Ile Tyr Ser Ser Pro Asp Phe His Asn Ser Cys Thr Ile Asn Ser
130     135     140
Ser Asp Tyr Gly Asn Asp Arg Trp Arg Val Gln Asn Cys Glu Leu Val
145     150     155     160
Gly Leu Pro Asp Leu Asn Thr Ser Ala Ser Tyr Val Gln Asn Ser Leu

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Ala	Ala	Tyr	Leu	165	Asn	Asp	Met	Leu	Ala	170	Ile	Gly	Val	Thr	Gly	175	Phe	Arg
Ile	Asp	Ala	Ala	180	Lys	His	Met	Ala	185	Ser	Ser	Asp	Ile	Ala	Gly	190	Ile	Lys
Ala	Lys	Leu	Ser	Arg	Ala	Pro	215	Leu	Ile	Tyr	Gln	Glu	Val	Ile	Asp	Gln		
Gly	Gly	Glu	Val	Ile	Thr	Ala	Ser	Glu	Tyr	Thr	220	Gly	Asn	Gly	Met	Val		
Thr	Glu	Phe	Lys	Tyr	Ser	Val	Gln	Leu	Gly	Asn	235	Val	Phe	Lys	Thr	Gly		
Arg	Leu	Ala	Ser	Leu	Ser	Asn	Phe	Gly	Glu	Ala	250	Trp	Gly	Phe	Leu	Pro		
Gly	Asn	Gln	Ala	Val	Val	Phe	Val	Asp	Asn	His	265	Asp	Asn	Gln	Arg	Gly		
His	Gly	Gly	Ala	Gly	Asn	Val	Val	Thr	Tyr	Lys	280	Asp	Gly	Lys	Leu	Tyr		
Asp	Leu	Ala	Asn	Val	Phe	Met	Leu	Ala	Tyr	Pro	295	Tyr	Gly	Tyr	Thr	Gln		
Val	Met	Ser	Ser	Tyr	Asp	Phe	Thr	Asp	Ala	Asp	310	Ala	Gly	Pro	Pro	Ser		
Ala	Ala	Val	His	Asn	Asn	Gly	Ser	Leu	Asn	Cys	325	Phe	Gly	Ser	Gln	Trp		
Lys	Cys	Glu	His	Arg	Trp	Ser	Thr	Ile	Ala	Gly	340	Ala	Val	Gln	Phe	Arg		
Asn	Asn	Thr	Ala	Glu	Glu	Trp	Arg	Val	Thr	His	355	Trp	Trp	Asp	Asn	Gly		
Asn	Asn	Gln	Ile	Ala	Phe	Gly	Arg	Ala	Ala	Leu	370	Gly	Phe	Val	Ala	Ile		
Asn	Lys	Glu	Ser	Tyr	Asn	Leu	Ser	Thr	Ser	Leu	385	Gln	Thr	Gly	Met	Thr		
Ala	Gly	Ser	Tyr	Cys	Asn	Val	Leu	Lys	Gly	Ser	405	Ile	Ser	Thr	Asp	Lys		
Thr	Ser	Cys	Ser	Gly	Glu	Val	Ile	Gln	Ile	Gly	420	Ser	Asp	Gly	Arg	Ile		
Gln	Ala	Asn	Leu	Ala	Ser	Trp	Asp	Ala	Phe	Ala	435	Ile	His	His	Gln	Ala		
Lys	Ile	Gly	Gly	Thr	Val	Gln	Pro	Pro	Ala	Gly	450	Asn	Trp	Gln	Arg	Thr		
Val	Val	Phe	Ile	Gln	Ala	Gln	Thr	Gln	Thr	Gly	465	Gln	Asp	Met	Phe	Val		
Arg	Gly	Gly	Leu	Asp	His	Ala	Val	Ala	Gln	Ser	480	Gln	Leu	Gly	Leu	Ser		
Cys	Ser	Ser	Ser	Asn	Tyr	Leu	Cys	Ala	Ile	Pro	500	Ile	Arg	His	Arg	Asn		
Leu	Asn	Asn	Ala	Thr	Thr	Ala	Pro	Trp	Lys	Ala	515	Asn	Asp	Gly	Leu	Leu		
Asp	Trp	Tyr	Gly	Arg	Glu	Ala	Ser	Gln	Ser	Ser	530	Ala	Ala	Glu	Gly	Thr		
Ala	Leu	Asp	Trp	Thr	Thr	Asn	Ser	Trp	Pro	Ala	545	Ser	Trp	Gly	Thr	Val		
Arg	Thr	Val	Ala	Thr	Asp	Gly	Tyr	Gly	Val	Glu	560	Val	Leu	Asn	Ser	Phe		
Gly	Gln	His	Tyr	Trp	Met	Leu	Asp	Val	Asp	Met	575	Asp	Cys	Ser	Lys	Thr		
Ile	Asn	Gly	Trp	Phe	Glu	Leu	Lys	Ala	Tyr	Val	590	Lys	Asn	Gly	Gln	Gly		
Trp	Glu	Gly	Asn	Ile	Ser	Gln	Ser	Gly	Thr	Pro	605	Tyr	Ala	Ser	Gln	Asn		
His	Met	Ala	Gln	Cys	Gly	Gln	Met	Asn	Val	Phe	620	Ser	Phe	Gly	Gln	Asn		
Ser	Ala	Gln	Ile	Ser	Ala	Leu	Pro				635							

<210> 404
 <211> 1653
 <212> DNA
 <213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 404

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aac	aag	gagg	cc	act	gccc	gt	tga	aac	ccc	ct	ttg	ttt	ggg	a	ggg	tgcc	aa	t	ct	gtatt	tc	120
ctc	ttg	gtg	acc	gct	ttt	ca	ta	acg	gca	ac	ccc	gata	acg	a	aca	acat	cat	c	cccc	gaaa	c	180
ga	a	aaa	acc	g	gg	t	gct	gag	aa	at	ttt	t	atg	g	gga	ggt	gacc	t	gcg	ggg	cat	240
att	gat	ga	aag	g	ct	at	ttt	ttc	c	ga	act	tag	gc	a	t	ct	gga	t	gac	tccc	atc	300
ga	a	aa	atcc	a	c	gg	agc	gg	t	ag	at	ga	a	ggc	a	cc	gga	a	ca	c	ggc	360
tgg	ac	caa	aag	a	ct	gg	ac	caa	t	at	c	g	ac	ccc	a	g	ct	at	gg	c	ccc	420
ct	gt	t	gaaa	a	ag	ccc	at	gc	a	ca	agg	c	at	t	c	g	at	c	g	cc	g	480
acc	gg	ac	cg	g	ta	acc	ga	act	c	g	at	c	ct	gt	t	tg	g	c	gg	a	a	540
aa	at	gt	ac	ct	a	t	gac	ag	ct	a	g	ca	att	at	a	t	c	g	att	gt	a	600
gac	gt	g	ctta	c	cg	ag	ag	caa	a	ga	a	ccc	g	t	a	a	c	c	t	cat	t	660
aaaa	a	ga	aag	g	cc	gt	t	ac	ga	a	at	c	g	t	ga	a	c	t	g	at	c	720
gg	t	at	cc	ga	g	ag	c	ac	caa	a	a	t	at	t	a	c	a	t	c	a	t	780
tt	g	t	att	g	a	c	g	t	t	ac	c	g	t	a	a	c	a	a	c	a	a	840
g	att	t	cc	a	a	g	t	at	g	t	a	a	a	a	a	a	a	a	a	a	a	900
gt	act	t	gac	g	act	t	t	t	at	g	att	g	at	t	g	g	c	a	g	t	c	960
aaa	a	ga	at	ct	a	t	g	ac	t	t	t	g	g	a	a	c	a	a	a	a	a	1020
ct	cat	ca	act	t	t	g	att	t	caa	a	a	g	g	c	c	g	t	g	a	a	a	1080
t	ac	a	acc	agg	c	g	t	g	ca	aaa	a	a	a	a	a	a	a	a	a	a	a	1140
c	ac	g	ac	g	ac	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	1200
ct	att	g	ct	ta	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	1260
g	aa	att	cccc	g	c	act	g	t	gg	g	c	g	at	c	c	a	a	a	a	a	a	1320
c	aaa	a	a	acc	c	c	a	a	c	c	c	a	a	a	a	a	a	a	a	a	a	1380
a	aaa	a	acc	acc	c	c	g	t	g	t	c	g	a	g	a	t	a	a	a	a	a	1440
a	c	tt	c	ag	c	a	g	a	a	a	a	a	a	a	a	a	a	a	a	a	a	1500
t	g	c	cat	t	t	g	c	a	a	a	a	a	a	a	a	a	a	a	a	a	a	1560
c	ac	g	at	g	c	c	t	t	t	c	ag	g	t	a	a	a	a	a	a	a	a	1620
c	aa	a	t	gg	g	c	a	a	a	a	a	a	a	a	a	a	a	a	a	a	a	1653

<210> 405

<211> 550

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(26)

<400> 405

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Ser	Cys	Ala	Pro	Asn	Lys	Glu	Ala	Thr	Ala	Val	Glu	Thr	Pro	Phe	Val						
			20					25					30								
Trp	Glu	Gly	Ala	Asn	Leu	Tyr	Phe	Leu	Leu	Val	Asp	Arg	Phe	His	Asn						
		35					40					45									
Gly	Asn	Pro	Asp	Asn	Asp	Asn	Ile	Ile	Pro	Arg	Asn	Glu	Glu	Thr	Gly						
	50			55					60												
Val	Leu	Arg	Asn	Phe	Met	Gly	Gly	Asp	Leu	Arg	Gly	Ile	Ile	Gln	Lys						
65				70				75						80							
Ile	Asp	Glu	Gly	Tyr	Phe	Ser	Glu	Leu	Gly	Ile	Asn	Ala	Ile	Trp	Met						
			85					90					95								
Thr	Pro	Ile	Val	Glu	Gln	Ile	His	Gly	Ala	Val	Asp	Glu	Gly	Thr	Gly						
			100					105					110								
Asn	Thr	Tyr	Gly	Phe	His	Gly	Tyr	Trp	Thr	Lys	Asp	Trp	Thr	Asn	Ile						
		115					120					125									
Asp	Pro	Ser	Tyr	Gly	Thr	Arg	Glu	Asp	Leu	Lys	Glu	Leu	Val	Glu	Lys						
	130					135					140										
Ala	His	Ala	Gln	Gly	Ile	Arg	Ile	Val	Leu	Asp	Ala	Val	Ile	Asn	His						
145				150				155						160							
Thr	Gly	Pro	Val	Thr	Glu	Leu	Asp	Pro	Val	Trp	Pro	Asp	Thr	Trp	Val						
			165					170						175							

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Arg Thr Gly Pro Lys Cys Thr Tyr Asp Ser Tyr Ser Asn Tyr Ile Asp
 180 185 190
 Cys Thr Leu Val Glu Asn Leu Pro Asp Val Leu Thr Glu Ser Lys Glu
 195 200 205
 Pro Val Glu Leu Pro Glu His Leu Ile Glu Lys Trp Lys Lys Glu Gly
 210 215 220
 Arg Tyr Glu Gln Glu Ile Ala Glu Leu Asp Ala Phe Phe Ala Arg Thr
 225 230 235 240
 Gly Tyr Pro Arg Ala Pro Lys Tyr Tyr Ile Met Lys Trp Leu Ser Asp
 245 250 255
 Tyr Ile Thr Asp Phe Gly Ile Asp Gly Tyr Arg Val Asp Thr Val Lys
 260 265 270
 His Thr Thr Glu Asp Val Trp Ala Asp Phe Ser Lys Val Cys Lys Asp
 275 280 285
 Ala Phe Ala Gln Tyr Lys Lys Asp Asn Pro Asp Lys Val Leu Asp Asp
 290 295 300
 Asn Asp Phe Phe Met Ile Gly Glu Val Tyr Gly Tyr Gly Ile His Gly
 305 310 315 320
 Lys Arg Ile Tyr Asp Phe Gly Asp Lys Gln Val Asp Tyr Phe Ala His
 325 330 335
 Gly Phe Asp Asn Leu Ile Asn Phe Asp Phe Lys Gly Asp Ala Asn Lys
 340 345 350
 Pro Phe Glu Glu Leu Phe Ser Lys Tyr Asn Gln Ala Val Gln Asn Asp
 355 360 365
 Leu Lys Gly Val Ser Ile Met Asn Tyr Val Ser Ser His Asp Asp Ser
 370 375 380
 Trp Pro Phe Asp Lys Lys Arg Glu Arg Ser Phe Glu Ala Gly Thr Lys
 385 390 395 400
 Leu Leu Leu Thr Pro Gly Ile Ser Gln Val Tyr Tyr Gly Asp Glu Thr
 405 410 415
 Ala Arg Ser Leu Glu Ile Pro Gly Thr Val Gly Asp Ala Thr Leu Arg
 420 425 430
 Ser Met Met Asn Trp Asp Asp Leu Gln Asn Asn Pro Asn Thr Gln Gln
 435 440 445
 Val Leu Lys His Phe Arg Leu Leu Gly Gln Phe Arg Lys Asn His Pro
 450 455 460
 Ala Val Gly Ala Gly Val His Gln Glu Ile Ser Lys Ala Pro Phe Tyr
 465 470 475 480
 Thr Phe Ser Arg Thr Phe Thr Lys Gly Asp Tyr Thr Asp Thr Val Ile
 485 490 495
 Ile Gly Leu Asp Leu Pro Phe Gly Lys Lys Glu Ile Gln Val Ser Ser
 500 505 510
 Ile Phe Glu Asp Gly Thr Glu Leu His Asp Ala Phe Ser Gly Ile Thr
 515 520 525
 Thr Thr Val Lys Asp Gly Lys Ala Val Ile Asp Ser Gln Met Gly Ile
 530 535 540
 Val Leu Leu Glu Arg Lys
 545 550

<210> 406

<211> 1398

<212> DNA

<213> Unknown

<220>

<223> obtained from an environmental sample

<400> 406

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ctcatcaagg	ggcggggcta	caagcagggtg	ctcatctcgc	cccctctcaa	gtcctccggc	180
aacgagtggg	gggcccgtta	ccagccccag	gatctgcggc	tcgtcgacag	cccgttgggc	240
aacaagcagg	atctcgagca	gctcatcgcc	gccatgcagg	ccgcggcat	cgccgtctat	300
gcggatgtgg	tgctcaacca	catggccaac	gagagctgga	agcgagtgga	cctcaactac	360
cccggcagcg	agctgttgca	aacctatgcc	gccaaacccg	cctacttcga	gcggcagaag	420
ctgttcggcg	atctggcgca	aaaccttctg	tcgggtcagg	acttccaccc	cgagggtgc	480
atcaccgact	ggagcgatcc	cggtcatgtc	cagtactggc	gcctgtgtgg	cggcgccggc	540
gacaaggggt	tgccggatct	cgaccccaac	aactgggtgg	tgagccagca	gcaggcctat	600

1001827087_1.txt

ctcaaggcgc	tcaaggggat	ggggatcaag	ggctttcggg	tcgatgcggt	caagcacatg	660
agcgattacc	agatcaacgc	cgtcttcacc	cccagatca	agcagggcat	gcacgtgttt	720
ggggaggtag	tcacctccgg	gggcgccggc	agcacggatt	acgagcgctt	cctcaagccc	780
tatctcgaca	acagcggcca	gggggcctat	gacttcccgc	tgtttgccct	cctgcgcggg	840
gcgctcggct	atggcggcag	catgaatcag	ctggcggatc	ccggcgccta	cggccaggcc	900
ctgccgggca	accgggcggg	caccttcgcc	atcacccacg	acatccccc	caacgacggc	960
ttccgctacc	agatcctgaa	ccagaccgac	gagaagctgg	cctatgccta	tctgctcggc	1020
cgcgatgggg	gctccccctt	ggtctattcg	gatcacggcg	agaccagga	caaggacggc	1080
ctgcgctggc	aggactacta	cctgcgcagc	gatctcaagg	ggatgatccg	cttcacaac	1140
gccgtgcagg	gtcagcccat	gcagctcatc	ggcagcggcg	actgcttcgt	gctgttcaag	1200
cgcggaagc	aggggctggt	cggggtcaac	aagtgtgact	acgagcagga	gtactggctc	1260
gacaccgcca	ggttcgagct	gaactgggat	cgcaactaca	gggacgtgct	ggatcaaaat	1320
gccgtcatca	atgtgcagag	ccagtgggta	cgggtcgcca	taccggcgcg	cagcgcccgc	1380
ctctggctgc	aggaataa					1398

<210> 407
 <211> 465
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(21)

<400> 407

Met	Lys	Asn	Thr	Ala	Gly	Ile	Trp	Val	Ile	Ala	Gly	Met	Leu	Ile	Ala
1				5					10					15	
Pro	Leu	Ala	Gln	Ala	Asp	Val	Ile	Leu	His	Ala	Phe	Asn	Trp	Lys	Tyr
			20					25					30		
Ser	Glu	Val	Thr	Ala	Lys	Ala	Asp	Leu	Ile	Lys	Gly	Ala	Gly	Tyr	Lys
		35					40					45			
Gln	Val	Leu	Ile	Ser	Pro	Pro	Leu	Lys	Ser	Ser	Gly	Asn	Glu	Trp	Trp
		50				55					60				
Ala	Arg	Tyr	Gln	Pro	Gln	Asp	Leu	Arg	Leu	Val	Asp	Ser	Pro	Leu	Gly
65					70					75				80	
Asn	Lys	Gln	Asp	Leu	Glu	Gln	Leu	Ile	Ala	Ala	Met	Gln	Ala	Arg	Gly
			85						90					95	
Ile	Ala	Val	Tyr	Ala	Asp	Val	Val	Leu	Asn	His	Met	Ala	Asn	Glu	Ser
			100					105					110		
Trp	Lys	Arg	Ser	Asp	Leu	Asn	Tyr	Pro	Gly	Ser	Glu	Leu	Leu	Gln	Thr
		115					120					125			
Tyr	Ala	Ala	Asn	Pro	Ala	Tyr	Phe	Glu	Arg	Gln	Lys	Leu	Phe	Gly	Asp
	130					135					140				
Leu	Ala	Gln	Asn	Leu	Leu	Ser	Gly	Gln	Asp	Phe	His	Pro	Glu	Gly	Cys
145					150					155					160
Ile	Thr	Asp	Trp	Ser	Asp	Pro	Gly	His	Val	Gln	Tyr	Trp	Arg	Leu	Cys
			165						170					175	
Gly	Gly	Ala	Gly	Asp	Lys	Gly	Leu	Pro	Asp	Leu	Asp	Pro	Asn	Asn	Trp
			180					185					190		
Val	Val	Ser	Gln	Gln	Gln	Ala	Tyr	Leu	Lys	Ala	Leu	Lys	Gly	Met	Gly
		195					200					205			
Ile	Lys	Gly	Phe	Arg	Val	Asp	Ala	Val	Lys	His	Met	Ser	Asp	Tyr	Gln
	210					215					220				
Ile	Asn	Ala	Val	Phe	Thr	Pro	Glu	Ile	Lys	Gln	Gly	Met	His	Val	Phe
225					230					235					240
Gly	Glu	Val	Ile	Thr	Ser	Gly	Gly	Ala	Gly	Ser	Thr	Asp	Tyr	Glu	Arg
			245						250					255	
Phe	Leu	Lys	Pro	Tyr	Leu	Asp	Asn	Ser	Gly	Gln	Gly	Ala	Tyr	Asp	Phe
			260					265					270		
Pro	Leu	Phe	Ala	Ser	Leu	Arg	Gly	Ala	Leu	Gly	Tyr	Gly	Gly	Ser	Met
		275					280					285			
Asn	Gln	Leu	Ala	Asp	Pro	Gly	Ala	Tyr	Gly	Gln	Ala	Leu	Pro	Gly	Asn
	290					295					300				
Arg	Ala	Val	Thr	Phe	Ala	Ile	Thr	His	Asp	Ile	Pro	Thr	Asn	Asp	Gly
305					310					315					320
Phe	Arg	Tyr	Gln	Ile	Leu	Asn	Gln	Thr	Asp	Glu	Lys	Leu	Ala	Tyr	Ala

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          325          330          335
Tyr Leu Leu Gly Arg Asp Gly Gly Ser Pro Leu Val Tyr Ser Asp His
      340      345      350
Gly Glu Thr Gln Asp Lys Asp Gly Leu Arg Trp Gln Asp Tyr Tyr Leu
      355      360      365
Arg Ser Asp Leu Lys Gly Met Ile Arg Phe His Asn Ala Val Gln Gly
      370      375      380
Gln Pro Met Gln Leu Ile Gly Ser Gly Asp Cys Phe Val Leu Phe Lys
      385      390      395      400
Arg Gly Lys Gln Gly Leu Val Gly Val Asn Lys Cys Asp Tyr Glu Gln
          405          410          415
Glu Tyr Trp Leu Asp Thr Ala Arg Phe Glu Leu Asn Trp Tyr Arg Asn
          420          425          430
Tyr Arg Asp Val Leu Asp Gln Asn Ala Val Ile Asn Val Gln Ser Gln
          435          440          445
Trp Val Arg Val Ala Ile Pro Ala Arg Ser Ala Arg Leu Trp Leu Gln
          450          455          460
Glu
465

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<210> 408
 <211> 1476
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

```

<400> 408
atgccgtcga tgcccccggt gcacgcttca gcgtcagccc cgctggcccc gtcccccgct      60
gcaccgcgcg cgacgtcgcg cctgcgccgt ggcctggtac gcctcgccgc caccgcactg      120
ctggccttcg gtgccatcgg cggcgcgcag gccgatgtga tcctgcatgc cttcaactgg      180
cgctatgccg acgtggaggc ccgcgcctcg cagatccagg ccgcgggcta ccgcgcggtg      240
ctggtggccc cggcgtaaa gagcgagggc agcgctggtt gggcgcgcta ccagccgcag      300
gattaccggc tgatccatca cccgctgggc gacaccaccg cgttccgcag catggtcaat      360
gcgctgaatg cgcgcgcat ccgctgttac gcggacgtca tcctaaacca catggccaac      420
gagtcggcgc aacgcagcga cctgaactac ccgggccagc gcctgctcaa cctctacgcc      480
ggcaaacacca gctactaaa cggccagcgg ctgttcggcg acctgcgcta caacttcctc      540
agcagctggg atttcggcca ggccacgtgc atcagcgact acaacgacgt gtggcagggtg      600
cagaactggc gcctgtgcgg cggcggtggt gacgccggcc tgccggacct cgtcaacggc      660
gactacgtgg tgtcgcagca gcaggcctac ctgcgcgcgc tgaaggccat gggcgtgaag      720
ggcttccgca tcgacgccgc gaagcacatg cccatcgccc acctcaaccg ggtgctgacc      780
agcgacatga agtcgggcat gcacatcttc ggcgagatca tcaactggcg cggcgcgggc      840
aacggtgagt acgaccgttt cctcgcgccg tacctgcagt acaccgacca cggcgcgtac      900
gacttccgcg tgacacgca gatccggggc gcattcggct tcggtggcag catgagcaac      960
ctggtcgacc cgcaggccta cgggcaggcg ctggcgccgg cgcgcgcggt caccctcacc      1020
gtcacgcacg acatcccga caacagcggc ttccgctacg cgctgatgga cccaccgcag      1080
gaaaccctcg cgtacgccta cgtgatgggg cgcaacggcg gctccccgct gctgtactcg      1140
gacaacaacg agagcggcga caaccgttg gtcaatgcgt ggaaccgtac cgacctaacg      1200
cagatgggtg cttccacaa tgcggtgcag ggcagcgacc aggcggtgct ggcgcacggc      1260
agctgctacc tgctgttccg ccgcggcagc ctgggcatcg tcggcatcaa caagtgcggc      1320
agcccggta acgtcacct caacatgaac aacagcgtgc tgtggtggta cgccaactac      1380
accgacgtgc tcgattcgac cagcaaggtc tatatcagca gcggaagcca taccttcacc      1440
ctgccggcgc gccgcgcgcg cctgtggctg aggtaa      1476

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<210> 409
 <211> 491
 <212> PRT
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(51)

<400> 409
 Met Pro Ser Met Pro Pro Leu His Ala Ser Ala Ser Ala Pro Leu Ala

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1	Pro	Ser	Pro	Ala	5	Pro	Arg	Ala	Thr	10	Ser	Arg	Leu	Arg	Arg	15	Gly	Leu
			20	Ala					25						30			
Val	Arg	Leu	Ala	Ala	Thr	Ala	Leu	Leu	Ala	Phe	Gly	Ala	Ile	Gly	Gly			
		35					40					45						
Ala	Gln	Ala	Asp	Val	Ile	Leu	His	Ala	Phe	Asn	Trp	Arg	Tyr	Ala	Asp			
	50					55				60								
Val	Glu	Ala	Arg	Ala	Ser	Gln	Ile	Gln	Ala	Ala	Gly	Tyr	Arg	Ala	Val			
65					70				75						80			
Leu	Val	Ala	Pro	Ala	Tyr	Lys	Ser	Glu	Gly	Ser	Ala	Trp	Trp	Ala	Arg			
			85						90					95				
Tyr	Gln	Pro	Gln	Asp	Tyr	Arg	Leu	Ile	His	His	Pro	Leu	Gly	Asp	Thr			
			100					105					110					
Thr	Ala	Phe	Arg	Ser	Met	Val	Asn	Ala	Leu	Asn	Ala	Arg	Gly	Ile	Arg			
		115					120					125						
Val	Tyr	Ala	Asp	Val	Ile	Leu	Asn	His	Met	Ala	Asn	Glu	Ser	Ala	Gln			
	130					135					140							
Arg	Ser	Asp	Leu	Asn	Tyr	Pro	Gly	Gln	Arg	Leu	Leu	Asn	Leu	Tyr	Ala			
145					150					155					160			
Gly	Asn	Thr	Ser	Tyr	Tyr	Asn	Gly	Gln	Arg	Leu	Phe	Gly	Asp	Leu	Arg			
			165					170						175				
Tyr	Asn	Phe	Leu	Ser	Ser	Trp	Asp	Phe	Gly	Gln	Ala	Thr	Cys	Ile	Ser			
		180						185					190					
Asp	Tyr	Asn	Asp	Val	Trp	Gln	Val	Gln	Asn	Trp	Arg	Leu	Cys	Gly	Gly			
		195					200					205						
Gly	Gly	Asp	Ala	Gly	Leu	Pro	Asp	Leu	Val	Asn	Gly	Asp	Tyr	Val	Val			
	210					215					220							
Ser	Gln	Gln	Gln	Ala	Tyr	Leu	Arg	Ala	Leu	Lys	Ala	Met	Gly	Val	Lys			
225					230					235					240			
Gly	Phe	Arg	Ile	Asp	Ala	Ala	Lys	His	Met	Pro	Ile	Ala	His	Leu	Asn			
			245					250					255					
Arg	Val	Leu	Thr	Ser	Asp	Ile	Lys	Ser	Gly	Met	His	Ile	Phe	Gly	Glu			
		260						265					270					
Ile	Ile	Thr	Gly	Gly	Gly	Ala	Gly	Asn	Gly	Glu	Tyr	Asp	Arg	Phe	Leu			
		275					280					285						
Ala	Pro	Tyr	Leu	Gln	Tyr	Thr	Asp	His	Gly	Ala	Tyr	Asp	Phe	Pro	Leu			
	290					295					300							
His	Thr	Gln	Ile	Arg	Gly	Ala	Phe	Gly	Phe	Gly	Gly	Ser	Met	Ser	Asn			
305					310					315					320			
Leu	Val	Asp	Pro	Gln	Ala	Tyr	Gly	Gln	Ala	Leu	Ala	Ala	Ala	Arg	Ala			
			325						330					335				
Val	Thr	Phe	Thr	Val	Thr	His	Asp	Ile	Pro	Asn	Asn	Ser	Gly	Phe	Arg			
			340					345					350					
Tyr	Ala	Leu	Met	Asp	Pro	Thr	Asp	Glu	Thr	Leu	Ala	Tyr	Ala	Tyr	Val			
		355					360					365						
Met	Gly	Arg	Asn	Gly	Gly	Ser	Pro	Leu	Leu	Tyr	Ser	Asp	Asn	Asn	Glu			
	370					375					380							
Ser	Gly	Asp	Asn	Arg	Trp	Val	Asn	Ala	Trp	Asn	Arg	Thr	Asp	Leu	Lys			
385					390					395					400			
Gln	Met	Val	Arg	Phe	His	Asn	Ala	Val	Gln	Gly	Ser	Asp	Gln	Ala	Val			
			405						410					415				
Leu	Ala	His	Gly	Ser	Cys	Tyr	Leu	Leu	Phe	Arg	Arg	Gly	Ser	Leu	Gly			
		420						425					430					
Ile	Val	Gly	Ile	Asn	Lys	Cys	Gly	Ser	Pro	Val	Asn	Val	Thr	Val	Asn			
		435					440					445						
Met	Asn	Asn	Ser	Val	Leu	Trp	Trp	Tyr	Ala	Asn	Tyr	Thr	Asp	Val	Leu			
	450					455					460							
Asp	Ser	Thr	Ser	Lys	Val	Tyr	Ile	Ser	Ser	Gly	Ser	His	Thr	Phe	Thr			
465					470					475					480			
Leu	Pro	Ala	Arg	Arg	Ala	Arg	Leu	Trp	Leu	Arg								
				485					490									

<210> 410
 <211> 1875
 <212> DNA
 <213> Unknown

<220>

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<223> Obtained from an environmental sample

<400> 410

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atgaaaaata caagcctgac cctgttagca ggctgcatgg ccctggcggt taacgcccag      60
gccgcagtcg gccaaaataa ccctgacatc atgttgcaag gcttccactg gaactccgcc      120
aaagcgggtg gctgggtacaa caccctgcaa ggcaatgtga acgaaatcgc cagcgccggc      180
ttcaacatgg tgtggctgcc gccccctcgc caggccggct cactggaagg ttacctgccg      240
gagcagtaca acaacctcaa ctccaactac ggcaccgaaa ctacgtgcg caacctgtct      300
agcgccctca aggccaacaa cgtcaaggcc attgccgaca tcgtcatcaa ccaccgtaat      360
ggctctggca gctgggtgtac cttcaccaac ccggcctggg gttttgatgc catcgtctct      420
aacgacgaag cctggggcgc agccggctcc aactgcagcg gcacccgtgg cgctgccgac      480
tctgggtgac gctatcacgc cgcccgcgac atcgaccaca gcaaaaccta tgtgcgcgac      540
agccttaaag agtggatgaa cgtccgcctc aaaggcggtg gctttgacgg ctggcgctat      600
gactacgtca aagggttcag cgggtgtctac gtcggcgaat ataacgccgc caccaacct      660
tacttcagcg tcggtgaata ctggacctcg ctctgtctaca acggcgaaga gtgctttcct      720
ggtggctctt ctccagaatc tcaccgccag gccagatca actgggtcga caagaccaat      780
ggcaacagcg cggcttttga cttaccacc aaagggtgc tcaacaaagc gctggagacc      840
tacaactaca gccacctgcg cgatgccaac ggcaaaccgg ccggggtgat cggctactgg      900
ccgtcgcgcg ccgtgacctt cgtcgacaac cagcagactg gcccagcgca agtctgtggc      960
aatgcccaga acaactggcc ggtgccgtgc gacaagggtg tgcagggtta tgcctacatc     1020
ctacccacc cggtgtgtgc ctccgtctac tacgccact acttcaactg ggggctgggc     1080
agcgaatca agaagttgat gaaactgcgc aaagacatgg ggctgcactc cgactccgcc     1140
gtgaccatcg acaaggcgca gcagggcctc tatgccgctt acatcggtgg caagggtggca     1200
gtcaaaactgg gtaatggttc ctggtcgccg tcaggcgctg gctggaccct ggcccagagc     1260
ggcactgatt gggccgtgtg gaaaaaagat gacggcaata acttcaagcg cactgtggtg     1320
ctgatctacg gcgaaactgc tgccggtcag gacatgttca tccgcggtgg tatcgaccac     1380
gcctatgccg cagccaacct cggcaagacc tgcaccagca ctaactacga gtgcgccatt     1440
ccgatcacc cacaacaacct gcgcaacgcc accaccgcac cgtggaaggc taacgacaag     1500
taccttgact ggtacgacgg tgtcgaagcc ggccagagca ctgccgctca gggctctgcc     1560
gctgactgga ccaccaacgt gtggccgtcg acctggggcg ccgtcaagac tgtggctgcc     1620
gatggtttcg gtgtagaacc gctcaacacc tacggtcagc actactggat gctggatgta     1680
cagatggatt gctccaagac cgtccaaggc atctggttcg agttcaagac cttcatcagc     1740
aacggccccg gttgggaagc caacgtcgcc cagagcggtg ccccgtagc cagcggtaac     1800
cactttggcc agtgcggtaa ggtcaacgtg ttcaagcgcg gcgtcagcgc cccggtggct     1860
atcagagact tctga                                     1875

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<210> 411

<211> 624

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(21)

<400> 411

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Phe Asn Ala Gln Ala Ala Val Ser Gln Asn Asn Pro Asp Ile Met Leu
          20          25          30
Gln Gly Phe His Trp Asn Ser Ala Lys Ala Gly Gly Trp Tyr Asn Thr
          35          40          45
Leu Gln Gly Asn Val Asn Glu Ile Ala Ser Ala Gly Phe Asn Met Val
          50          55          60
Trp Leu Pro Pro Pro Ser Gln Ala Gly Ser Leu Glu Gly Tyr Leu Pro
          65          70          75          80
Glu Gln Tyr Asn Asn Leu Asn Ser Asn Tyr Gly Thr Glu Thr Gln Leu
          85          90          95
Arg Asn Leu Leu Ser Ala Leu Lys Ala Asn Asn Val Lys Ala Ile Ala
          100          105          110
Asp Ile Val Ile Asn His Arg Asn Gly Ser Gly Ser Trp Cys Thr Phe
          115          120          125
Thr Asn Pro Ala Trp Gly Phe Asp Ala Ile Val Ser Asn Asp Glu Ala
          130          135          140
Trp Gly Ala Ala Gly Ser Asn Cys Ser Gly Thr Arg Gly Ala Ala Asp
          145          150          155          160

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Ser	Gly	Asp	Gly	Tyr	His	Ala	Ala	Arg	Asp	Ile	Asp	His	Ser	Lys	Thr
Tyr	Val	Arg	Asp	Ser	Leu	Lys	Glu	Trp	Met	Asn	Val	Arg	Leu	Lys	Gly
Val	Gly	Phe	Asp	Gly	Trp	Arg	Tyr	Asp	Tyr	Val	Lys	Gly	Phe	Ser	Gly
Val	Tyr	Val	Gly	Glu	Tyr	Asn	Ala	Ala	Thr	Asn	Pro	Tyr	Phe	Ser	Val
Gly	Glu	Tyr	Trp	Thr	Ser	Leu	Cys	Tyr	Asn	Gly	Glu	Glu	Cys	Phe	Pro
Gly	Gly	Ser	Ser	Pro	Glu	Ser	His	Arg	Gln	Ala	Gln	Ile	Asn	Trp	Val
Asp	Lys	Thr	Asn	Gly	Asn	Ser	Ala	Val	Phe	Asp	Phe	Thr	Thr	Lys	Gly
Leu	Leu	Asn	Lys	Ala	Leu	Glu	Thr	Tyr	Asn	Tyr	Ser	His	Leu	Arg	Asp
Ala	Asn	Gly	Lys	Pro	Ala	Gly	Val	Ile	Gly	Tyr	Trp	Pro	Ser	Arg	Ala
Val	Thr	Phe	Val	Asp	Asn	His	Asp	Thr	Gly	Pro	Ser	Glu	Val	Cys	Gly
Asn	Ala	Gln	Asn	Asn	Trp	Pro	Val	Pro	Cys	Asp	Lys	Val	Met	Gln	Gly
Tyr	Ala	Tyr	Ile	Leu	Thr	His	Pro	Gly	Val	Pro	Ser	Val	Tyr	Tyr	Ala
His	Tyr	Phe	Asn	Trp	Gly	Leu	Gly	Ser	Glu	Ile	Lys	Lys	Leu	Met	Lys
Leu	Arg	Lys	Asp	Met	Gly	Leu	His	Ser	Asp	Ser	Ala	Val	Thr	Ile	Asp
Lys	Ala	Gln	Gln	Gly	Leu	Tyr	Ala	Ala	Tyr	Ile	Gly	Gly	Lys	Val	Ala
Val	Lys	Leu	Gly	Asn	Gly	Ser	Trp	Ser	Pro	Ser	Gly	Ala	Gly	Trp	Thr
Leu	Ala	Gln	Ser	Gly	Thr	Asp	Trp	Ala	Val	Trp	Lys	Lys	Asp	Asp	Gly
Asn	Asn	Phe	Lys	Arg	Thr	Val	Val	Leu	Ile	Tyr	Gly	Glu	Thr	Ala	Ala
Gly	Gln	Asp	Met	Phe	Ile	Arg	Gly	Gly	Ile	Asp	His	Ala	Tyr	Ala	Ala
Ala	Asn	Leu	Gly	Lys	Thr	Cys	Thr	Ser	Thr	Asn	Tyr	Glu	Cys	Ala	Ile
Pro	Ile	Thr	His	Asn	Asn	Leu	Arg	Asn	Ala	Thr	Thr	Ala	Pro	Trp	Lys
Ala	Asn	Asp	Lys	Tyr	Leu	Asp	Trp	Tyr	Asp	Gly	Val	Glu	Ala	Gly	Gln
Ser	Thr	Ala	Ala	Gln	Gly	Ser	Ala	Ala	Asp	Trp	Thr	Thr	Asn	Val	Trp
Pro	Ser	Thr	Trp	Gly	Ala	Val	Lys	Thr	Val	Ala	Ala	Asp	Gly	Phe	Gly
Val	Glu	Pro	Leu	Asn	Thr	Tyr	Gly	Gln	His	Tyr	Trp	Met	Leu	Asp	Val
Gln	Met	Asp	Cys	Ser	Lys	Thr	Val	Gln	Gly	Ile	Trp	Phe	Glu	Phe	Lys
Thr	Phe	Ile	Ser	Asn	Gly	Pro	Gly	Trp	Glu	Ala	Asn	Val	Ala	Gln	Ser
Gly	Thr	Pro	Tyr	Ala	Ser	Gly	Asn	His	Phe	Gly	Gln	Cys	Gly	Lys	Val
Asn	Val	Phe	Lys	Arg	Gly	Val	Ser	Ala	Pro	Val	Ala	Ile	Arg	Asp	Phe

<210> 412
<211> 2088

<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

<400> 412

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ctcgggtgacg ccgtcgtctgt cgaggtcgat gcgttcgcgg acggccacga ccgcgtcgcg 180
tgccgcgtgctc ggtggcggca cgaggaagac gccgcgtggc gcgaggcgcc gatggaggcg 240
ctcgcgaacg atcgggtggcg tgcgacgttc gcggtcgatc gcctcgggcg ctggcggtat 300
gcgttcgcgcg cgtggatcga tcgctacggg acgtgggtgc acgacctcgc ccggcgcccgc 360
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gccgcgggcg cgcaagggca cgcggccgag cgactgcgcg aggtcgcgcg gctgctcggc 480
ggcgaggccg cggagagcgc gaagcgcgcg gtcgcgtcga gtgcggacac gtgcgcgtg 540
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gtcgagagg gagcggcgcc aggcacgccc gcgcgaacgc cgccgagtgc gtcggcgacc 720
ggcgcgttgc gccacggccg gttccgcgac ctcgagccga tgctcgacta cgtcgcgcgg 780
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ggtccgaaca acgcgctcgc ggcgggaccc gaggacgtcg gcagcccggt ggcgatcggg 900
ctgcgcggcg cgggcggcga gggcgggcat cgcgccgtgc acccgagct cggcacgctc 960
gaggacttcc gctggctcgt ggagcgcgcg cgagcgcgcg gcacgcgat cgcgctcgac 1020
gtcgcgttcc agtgcgcgc ggaccacccg tacgtgcacg agcacccgga gtggttccgt 1080
cgccggccgg acggcagcat ccagtacgcc gagaacccgc cgaagaagta ccaggacatc 1140
taccggttcg acttcgagtc cgagcactgg cccgcgtgt gggacgaact cgcggcggtg 1200
ttcgagttct gggtcgagca gggcgtgacg atcttccgcg tcgacaaccc gcacacgaag 1260
gcgttcccgt tctgggagta cgcgatcgcg cgcgtgaaga gtcgctgccc gcaggcgatc 1320
ttcctgtccg aggcattcac gcggccgaag gtcatgcacc ggctcgcgaa gctcggcttc 1380
acgcagtcgt acacgtactt cacctggcgc aacacgcggc acgagctcgt gcagtacttc 1440
accgagctcg cgcacgggcc gggagcggag tacttccggc cgaacgcgtg gccgaacacg 1500
ccggacatcc tgccggagta cctgcagtc ggcggccgcg cggcattcat ggtgcgcgcg 1560
gtgctcgcgt cgctgctgtc gtcgaactgg ggcgtgtacg gaccggcatt cgagctgctc 1620
gagcacgagc cgcgttccgc cggcagcgag gagtacctcg actccgagaa gtaccagctg 1680
cggcggtggg atctcgaccg gccggacagc ctgcgcgagt tcgtcgcctg gctgaaccgc 1740
atccggcgcg agtcgccggc gctgcagcag atggcgacgc tcgcgttcca cgacgccgac 1800
aacgagcagc tgctgtgctg gtcgaagacc gccggcgacg acgcgatcgt cgtcgtcgtg 1860
aacctcgacc cgcaccacgc gcaggccggc tggatcacgc tgccgctcga gcgactcggg 1920
ctcgagccgg agcgcgcgta ccaggtgcac gacctgctcg gcggcgagc cttcctctgg 1980
agcgggccgc ggaactacgt cgcgctcgcg ccggacgcgc tgccggcgca cgtgttccgc 2040
gtgcggcgac atgcgcgcgc cgagcaggac ttcgactact tcctctga 2088

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<210> 413
 <211> 695
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 413
Met Asp Ala Arg Thr Gly Glu Arg Arg Ala Ala Leu Ala Ala Leu Pro
 1          5          10          15
Gly Arg Glu Arg Val Val Leu Glu Arg Ala Thr Pro Glu Ile Asp Gly
          20          25          30
Gly Arg Phe Ala Ala Lys Arg Val Leu Gly Asp Ala Val Val Val Glu
          35          40          45
Val Asp Ala Phe Ala Asp Gly His Asp Arg Val Ala Cys Ala Leu Arg
          50          55          60
Trp Arg His Glu Glu Asp Ala Ala Trp Arg Glu Ala Pro Met Glu Ala
65          70          75          80
Leu Ala Asn Asp Arg Trp Arg Ala Thr Phe Ala Val Asp Arg Leu Gly
          85          90          95
Arg Trp Arg Tyr Ala Phe Ala Ala Trp Ile Asp Arg Tyr Gly Thr Trp
          100          105          110
Val His Asp Leu Ala Arg Arg Pro Leu Ala Asp Ala Asp Leu Pro Ala
          115          120          125
Thr Phe Ala Ala Gly Ala Ala Leu Val Arg Glu Ala Ala Ala Gly Ala
          130          135          140
Gln Gly His Ala Ala Glu Arg Leu Arg Glu Val Ala Ala Leu Leu Gly
145          150          155          160
Gly Glu Ala Ala Glu Ser Ala Lys Arg Ala Val Ala Ser Ser Ala Asp
          165          170          175
Thr Cys Ala Leu Met Arg Glu His Asp Pro Arg Leu His Ala Ala Asp

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Tyr	Gly	Arg	180	Val	Leu	Glu	Val	Ile	185	Val	Asp	Pro	Pro	Leu	190	Ala	Arg	Cys
Ser	Ala	Trp	195	Tyr	Glu	Phe	Phe	Pro	200	Arg	Ser	Phe	Gly	Val	205	Ala	Glu	Gly
Ala	Ala	Pro	210	Gly	Thr	Pro	Ala	Arg	215	Thr	Pro	Pro	Ser	Ala	220	Ser	Ala	Thr
225	Gly	Ala	Leu	Arg	His	230	Gly	Arg	235	Asp	Leu	Glu	Pro	Met	240	Leu	Asp	245
Tyr	Val	Ala	Arg	245	Leu	Gly	Phe	Asp	250	Val	Leu	His	Phe	Pro	255	Pro	Ile	His
Pro	Ile	Gly	Arg	260	Ala	Phe	Arg	Lys	265	Gly	Pro	Asn	Asn	Ala	270	Leu	Ala	Ala
Gly	Pro	Glu	Asp	275	Val	Gly	Ser	Pro	280	Trp	Ala	Ile	Gly	Leu	285	Arg	Gly	Ala
305	Gly	Gly	Glu	Gly	Gly	His	Arg	Ala	310	Val	His	Pro	Glu	Leu	315	Gly	Thr	Leu
Glu	Asp	Phe	Arg	Trp	Leu	Val	Glu	Arg	325	Ala	Arg	Ala	Arg	Gly	330	Ile	Glu	335
Ile	Ala	Leu	Asp	340	Val	Ala	Phe	Gln	345	Cys	Ser	Pro	Asp	His	350	Pro	Tyr	Val
His	Glu	His	Pro	355	Glu	Trp	Phe	Arg	360	Arg	Arg	Pro	Asp	Gly	365	Ser	Ile	Gln
Tyr	Ala	Glu	Asn	Pro	Pro	Lys	Lys	Tyr	375	Gln	Asp	Ile	Tyr	Pro	380	Phe	Asp	385
Phe	Glu	Ser	Glu	His	Trp	Pro	Ala	Leu	390	Trp	Asp	Glu	Leu	Ala	395	Gly	Val	400
Phe	Glu	Phe	Trp	Val	Glu	Gln	Gly	Val	405	Thr	Ile	Phe	Arg	Val	410	Asp	Asn	415
Pro	His	Thr	Lys	Ala	Phe	Pro	Phe	Trp	420	Glu	Tyr	Ala	Ile	Ala	425	Arg	Val	430
Lys	Ser	Arg	Cys	Pro	Gln	Ala	Ile	Phe	435	Leu	Ser	Glu	Ala	Phe	440	Thr	Arg	445
Pro	Lys	Val	Met	His	Arg	Leu	Ala	Lys	450	Leu	Gly	Phe	Thr	Gln	455	Ser	Tyr	460
Thr	Tyr	Phe	Thr	Trp	Arg	Asn	Thr	Arg	465	His	Glu	Leu	Val	Gln	470	Tyr	Phe	475
Thr	Glu	Leu	Ala	His	Gly	Pro	Gly	Ala	480	Glu	Tyr	Phe	Arg	Pro	485	Asn	Ala	490
Trp	Pro	Asn	Thr	Pro	Asp	Ile	Leu	Pro	495	Glu	Tyr	Leu	Gln	Leu	500	Gly	Gly	505
Arg	Ala	Ala	Phe	Met	Val	Arg	Ala	Val	510	Leu	Ala	Ser	Leu	Ser	515	Ser	Ser	520
Asn	Trp	Gly	Val	Tyr	Gly	Pro	Ala	Phe	525	Glu	Leu	Leu	Glu	His	530	Glu	Pro	535
Arg	Ser	Ala	Gly	Ser	Glu	Glu	Tyr	Leu	540	Asp	Ser	Glu	Lys	Tyr	545	Gln	Leu	550
Arg	Arg	Trp	Asp	Leu	Asp	Arg	Pro	Asp	555	Ser	Leu	Ala	Glu	Phe	560	Val	Ala	565
Val	Leu	Asn	Arg	Ile	Arg	Arg	Glu	Ser	570	Pro	Ala	Leu	Gln	Gln	575	Met	Ala	580
Thr	Leu	Ala	Phe	His	Asp	Ala	Asp	Asn	585	Glu	Gln	Leu	Leu	Cys	590	Trp	Ser	595
Lys	Thr	Ala	Gly	Asp	Asp	Ala	Ile	Val	600	Val	Val	Val	Val	Asn	605	Leu	Asp	610
His	His	Ala	Gln	Ala	Gly	Trp	Ile	Thr	615	Leu	Pro	Leu	Glu	Arg	620	Leu	Gly	625
Leu	Glu	Pro	Glu	Arg	Ala	Tyr	Gln	Val	630	His	Asp	Leu	Leu	Gly	635	Gly	Gly	640
Arg	Phe	Leu	Trp	Ser	Gly	Pro	Arg	Asn	645	Tyr	Val	Ala	Leu	Ala	650	Pro	Asp	655
Ala	Leu	Pro	Ala	His	Val	Phe	Arg	Val	660	Arg	Arg	His	Ala	Arg	665	Arg	Glu	670
Gln	Asp	Phe	Asp	Tyr	Phe	Leu	680	685										

<210> 414
<211> 2262

<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

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<400> 414
atgtcgaatgc agatacgtac agacaaggct caggagatcg atcgcaatgt tgtgctcact      60
atgcctgtca aggcatatcga gactgtaacg gtcactaagg atggcctttc ttctttccct      120
gagattccga atgcagatga accggcgact cttttctata aagcccctgc aggatctgtg      180
tttgacgggt atactaaaga cttatatgct catatcggag tcattgatgg cgagtggaa      240
catgtggttt cggcatggga tgtagatgtg gctaagtgca aatggcataa gacagcccag      300
gataatgtat ggtcacttac actcgagcct tctatccgtg agtggttcgc ttccggcaca      360
actccagatga atgtgatcgg tgtagtggta agagatgcaa agggagcagg aagccttcag      420
acagaagacc tgactcttaa tgtgtatgac cctgtatatg gaacaggtga tatgcctaca      480
ccgcttccgg ctggagtaaa acatggaata aatatcattg acaacaagac cgtaactctt      540
gtgcttcttg agaaagataa gaatggtggt cactatgatg agtgctacct tatcggaact      600
gactttggca aatggagcag ggatgaaaaa tacaagatga acagggatga ggctgcccgc      660
tgctgggtgga tcactcttac cggctcttacc aggacaaagg aatataatgtt ccagtacgaa      720
ctgtcaagg gcaattcttc agttagggtg catgatccat attctgagat tgtttatgat      780
atgtacaacg atcagtggac atacgcaagc aataagatac ctaccggaaa gacagacggg      840
cttgatatctt cattccagat cagaaagtct acatatccat gggattaccc tgattttgag      900
attgaagatg aggatgacct tgtgatttat gaaatgcttg taagagactt caatacaagc      960
catagcatta gcgatgccat cgaccagctg gattatatct caggtcttgg agtgaatgct      1020
atagaactca tgcctatcca ggagtttgag gcaaacgaga gctggggata tgctcctcat      1080
tcatatttcg cattggataa atattacggt caacctacag aatataagaa attcgtcgat      1140
gcatgtcatg cgagggttat tgcagtgatt gtagatgtag tatacaatca tgctacaggt      1200
gctcatccta tggccaagat gtattggagc ggtaatgcta catctgcata caatccttgg      1260
ttcaatgtga cggcacctca tggtgacagt gtatatcatg actggaatca tagtaatgtc      1320
gatgttagaa atcatatcaa gagaagcctt gagtatctca tttctgaata taagggtggat      1380
ggattcagat ttgacctttc aaaagggtttc atccagagtg gaaatgagta tgactggaat      1440
caggacagaa caaactggat aaaggaatat tacaatacca taaaggcagc tgatccgaat      1500
gcagtgggta ttcttgagca ttgggtggat aatgagaatt atgacctctg caattatgga      1560
atgaagggtg ggaacaaggc atgtgaacag tattatcagt ccggtatggg ttactcttcc      1620
gacagtgatt tctcaggaat cagagagcct tcatggttgc ctttcgggtt atatatatca      1680
ttcatggaga gccatgatga agaaagagtc ggcttcaagc agaaagagta tggtaacggc      1740
agtgtcaaga cagatcttgc tttgagaatg cgtcgtgcag gccttaatgc agcattcttc      1800
cttactgttc caggtcctaa gatgatatgg cagttcgggt agatcggtta tgatacag      1860
atcgatcaca acggacgtac cggaaacaag cctgtcgtga ctgacgagta tctcgcaaac      1920
gagcatagga aggggcttta tgatacgtat gccggtcttt tggagttcag aaagaacaat      1980
ccgcgtttct ttgatgatga ttccaacttc cgttggggcg tttctacatc aaattggcca      2040
ggacgttata tcatgaatac gtcaaaggag ggtaagactt atgccatctt cggtaatttc      2100
ggctctggaa gtcagacaat aacgatggaa ctcccttctg aaggcccttg gtataactat      2160
tacaacagca ctgaggtgtg gaacggaaag cgactctcaa agaagggtgaa      2220
tttgtcttcc tcgtagacga taagagtctg tgcaagaatt ga      2262

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<210> 415
<211> 753
<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

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<400> 415
Met Ser Met Gln Ile Arg Thr Asp Lys Ala Gln Glu Ile Asp Arg Asn
 1           5           10           15
Val Val Leu Thr Met Pro Val Lys Ala Tyr Glu Thr Val Thr Val Thr
          20           25           30
Lys Asp Gly Leu Ser Ser Phe Pro Glu Ile Pro Asn Ala Asp Glu Pro
          35           40           45
Ala Thr Leu Phe Tyr Lys Ala Pro Ala Gly Ser Val Phe Asp Gly Tyr
          50           55           60
Thr Lys Asp Leu Tyr Ala His Ile Gly Val Ile Asp Gly Glu Trp Lys
65           70           75           80
His Val Val Ser Ala Trp Asp Val Asp Val Ala Lys Cys Lys Trp His
          85           90           95
Lys Thr Ala Gln Asp Asn Val Trp Ser Leu Thr Leu Glu Pro Ser Ile

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Arg	Glu	Trp	100	Phe	Ala	Ser	Gly	Thr	105	Thr	Pro	Val	Asn	Val	110	Ile	Gly	Val
Val	Val	Arg	115	Asp	Ala	Lys	Gly	Ala	120	Gly	Ser	Leu	Gln	Thr	125	Glu	Asp	Leu
Thr	Leu	Asn	130	Val	Tyr	Asp	Pro	Val	135	Tyr	Gly	Thr	Gly	Asp	Met	Pro	Thr	
Pro	Leu	Pro	145	Ala	Gly	Val	Lys	His	150	Gly	Ile	Asn	Ile	Ile	Asp	Asn	Lys	
Thr	Val	Thr	165	Leu	Val	Leu	Leu	Glu	170	Lys	Asp	Lys	Asn	Gly	Gly	His	Tyr	
Asp	Glu	Cys	180	Tyr	Leu	Ile	Gly	Thr	185	Asp	Phe	Gly	Lys	Trp	Ser	Arg	Asp	
Glu	Lys	Tyr	195	Lys	Met	Asn	Arg	Asp	200	Glu	Ala	Ala	Gly	Cys	Trp	Trp	Ile	
Thr	Leu	Thr	210	Gly	Leu	Thr	Arg	Thr	215	Lys	Glu	Tyr	Met	Phe	Gln	Tyr	Glu	
Leu	Val	Lys	225	Gly	Asn	Ser	Ser	Val	230	Arg	Val	His	Asp	Pro	Tyr	Ser	Glu	
Ile	Val	Tyr	245	Asp	Met	Tyr	Asn	Asp	250	Gln	Trp	Thr	Tyr	Ala	Ser	Asn	Lys	
Ile	Pro	Thr	260	Gly	Lys	Thr	Asp	Gly	265	Leu	Val	Ser	Ser	Phe	Gln	Ile	Arg	
Lys	Ser	Thr	275	Tyr	Ser	Trp	Asp	Tyr	280	Pro	Asp	Phe	Glu	Ile	Glu	Asp	Glu	
Asp	Asp	Leu	290	Val	Ile	Tyr	Glu	Met	295	Leu	Val	Arg	Asp	Phe	Asn	Thr	Ser	
His	Ser	Ile	305	Ser	Asp	Ala	Ile	Asp	310	Gln	Leu	Asp	Tyr	Ile	Ser	Gly	Leu	
Gly	Val	Asn	325	Ala	Ile	Glu	Leu	Met	330	Ile	Gln	Glu	Phe	Glu	Ala	Asn		
Glu	Ser	Trp	340	Gly	Tyr	Ala	Pro	His	345	Ser	Tyr	Phe	Ala	Leu	Asp	Lys	Tyr	
Tyr	Gly	Gln	355	Pro	Thr	Glu	Tyr	Lys	360	Lys	Phe	Val	Asp	Ala	Cys	His	Ala	
Arg	Gly	Ile	370	Ala	Val	Ile	Val	Asp	375	Val	Val	Tyr	Asn	His	Ala	Thr	Gly	
Ala	His	Pro	385	Met	Ala	Lys	Met	Tyr	390	Trp	Ser	Gly	Asn	Ala	Thr	Ser	Ala	
Ser	Asn	Pro	405	Trp	Phe	Asn	Val	Thr	410	Ala	Pro	His	Gly	Asp	Ser	Val	Tyr	
His	Asp	Trp	420	Asn	His	Ser	Asn	Val	425	Asp	Val	Arg	Asn	His	Ile	Lys	Arg	
Ser	Leu	Glu	435	Tyr	Leu	Ile	Ser	Glu	440	Tyr	Lys	Val	Asp	Gly	Phe	Arg	Phe	
Asp	Leu	Ser	450	Lys	Gly	Phe	Ile	Gln	455	Ser	Gly	Asn	Glu	Tyr	Asp	Trp	Asn	
Gln	Asp	Arg	465	Thr	Asn	Trp	Ile	Lys	470	Glu	Tyr	Asn	Thr	Ile	Lys	Ala		
Ala	Asp	Pro	485	Ala	Val	Val	Ile	Leu	490	Glu	His	Trp	Val	Asp	Asn	Glu		
Asn	Tyr	Asp	500	Leu	Cys	Asn	Tyr	Gly	505	Met	Lys	Val	Trp	Asn	Lys	Ala	Cys	
Glu	Gln	Tyr	515	Tyr	Gln	Ser	Gly	Met	520	Gly	Tyr	Ser	Ser	Asp	Ser	Asp	Phe	
Ser	Gly	Ile	530	Arg	Glu	Pro	Ser	Trp	535	Leu	Pro	Phe	Gly	Ser	Tyr	Ile	Ser	
Phe	Met	Glu	545	Ser	His	Asp	Glu	Glu	550	Arg	Val	Gly	Phe	Lys	Gln	Lys	Glu	
Tyr	Gly	Asn	565	Gly	Ser	Val	Lys	Thr	570	Asp	Leu	Ala	Leu	Arg	Met	Arg	Arg	
Ala	Gly	Leu	580	Asn	Ala	Ala	Phe	Phe	585	Leu	Thr	Val	Pro	Gly	Pro	Lys	Met	
Ile	Trp	Gln	595	Phe	Gly	Glu	Ile	Gly	600	Tyr	Asp	Ile	Ser	Ile	Asp	His	Asn	
Gly	Arg	Thr	610	Gly	Asn	Lys	Pro	Val	615	Val	Thr	Asp	Glu	Tyr	Leu	Ala	Asn	
Glu	His	Arg	625	Lys	Gly	Leu	Tyr	Asp	630	Thr	Tyr	Ala	Gly	Leu	Leu	Glu	Phe	

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				645					650					655			
Arg	Lys	Asn	Asn	Pro	Arg	Phe	Phe	Asp	Asp	Ser	Asn	Phe	Arg	Trp			
			660					665				670					
Gly	Val	Ser	Thr	Ser	Asn	Trp	Pro	Gly	Arg	Tyr	Ile	Met	Asn	Thr	Ser		
		675					680					685					
Lys	Glu	Gly	Lys	Thr	Tyr	Ala	Ile	Phe	Gly	Asn	Phe	Gly	Ser	Gly	Ser		
	690					695					700						
Gln	Thr	Ile	Thr	Met	Glu	Leu	Pro	Ser	Glu	Gly	Pro	Trp	Tyr	Asn	Tyr		
705					710					715					720		
Tyr	Asn	Ser	Thr	Glu	Val	Trp	Asn	Gly	Lys	Asn	His	Lys	Pro	Thr	Leu		
				725					730					735			
Lys	Glu	Gly	Glu	Phe	Val	Phe	Leu	Val	Asp	Asp	Lys	Ser	Leu	Cys	Lys		
			740					745					750				

Asn

<210> 416
 <211> 1344
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 416

atgaaccggg	ttaagagact	ttcgattttg	gtcgttcttt	tccttaccgc	cttcattccg	60
actgtttttg	ccggcgagca	gcctctcgcc	atctttcacg	cctttaacga	ccccttcaact	120
cttggtgaat	cctatgtctg	cgaactcgcc	gggcagggat	actcacatgt	ccagatatct	180
ccagcgcaga	agtcgaaccc	tgctcgagcc	tggtatgccc	ggtatcaacc	cgtagatttt	240
actgtcatcg	aagggatggg	cactgagagc	gatctgagga	agctcacgga	taaggccac	300
gcgtgtggaa	ttaaaggatg	cgccgatgtg	gtcttcaacc	acatgtcgag	catggacgag	360
tacaaggggc	ttgacaagtt	tccgggactt	gtcctgctg	atttccaccg	gcagtgcggc	420
atcgattatt	caaaacgaga	ttcgggtgcg	aactgttggc	tcggaggcga	cttgcccgat	480
ctggaccagt	cccggccgag	ggtacaggat	gttcagagag	cccacataag	gaagctcctt	540
tccctcggca	tagacggctt	ccgcttcgat	gcggctaacc	acatcgaccc	cattgttgtg	600
aaagactaca	tcgatctcat	cgacagggag	agcaacggca	ggacctggaa	ctacctcgag	660
gtcatcgagg	atgacggcac	tcaggccacg	gactacaact	ggatagcggc	agtgaccgat	720
ttcgtcctct	acaaggagtc	gttgaggaag	gccttcagtc	tcggcgggga	cctgcgatcg	780
ctcaagatgc	ctgtggctgt	caatgattcg	cggagtatcg	tcttcgggag	aaatcacgac	840
accgtgccgg	agaataacca	gaactgcac	gtcggctgct	acgacagccg	ggaggactcc	900
tatcttgcca	cggcatacgt	cctggcccgc	gaatcgggag	tcccgtgggt	cctcaactgg	960
gacaactacg	acgcgcccta	catcagcacc	ggcgtgaagt	tccgccagat	catgacgcag	1020
cgaggacgat	cggccatgaa	cgtgaaggag	aatgtgctgg	gcgtcatcga	cagtcctgtc	1080
gtcatgtatg	tggagcgcgg	gagtgaaggc	tttttcgtcc	tcaacaagag	cgccgaccgg	1140
ttcgatatcc	cagttctgga	tctgacactg	accaatctcg	agggatgtta	tcgggagctg	1200
agaagaaaat	tcaccgtcgc	catcgagaga	aagtacggta	agaaatttgt	caccgggtgg	1260
ggacgatggg	accggggggg	cctcgaaatc	tacggccgcg	acgctctcta	cttcatacgg	1320
gaaccctggg	agcagtgcag	gtaa				1344

<210> 417
 <211> 447
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(24)

<400> 417

Met	Asn	Arg	Val	Lys	Arg	Leu	Ser	Ile	Leu	Val	Val	Leu	Phe	Leu	Thr
1				5				10					15		
Ala	Phe	Ile	Pro	Thr	Val	Phe	Ala	Gly	Glu	Gln	Pro	Leu	Ala	Ile	Phe
		20					25					30			
His	Ala	Phe	Asn	Asp	Pro	Phe	Thr	Leu	Val	Glu	Ser	Tyr	Val	Cys	Glu
	35					40				45					
Leu	Ala	Gly	Gln	Gly	Tyr	Ser	His	Val	Gln	Ile	Ser	Pro	Ala	Gln	Lys

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50      55      60
Ser  Asn  Pro  Ala  Arg  Ala  Trp  Tyr  Ala  Arg  Tyr  Gln  Pro  Val  Asp  Phe
65  Thr  Val  Ile  Glu  Gly  Met  Gly  Thr  Glu  Ser  Asp  Leu  Arg  Lys  Leu  Thr
      85
Asp  Lys  Ala  His  Ala  Cys  Gly  Ile  Lys  Val  Ile  Ala  Asp  Val  Val  Phe
      100
Asn  His  Met  Ser  Ser  Met  Asp  Glu  Tyr  Lys  Gly  Leu  Asp  Lys  Phe  Pro
      115
Gly  Leu  Ala  Pro  Ala  Asp  Phe  His  Arg  Gln  Cys  Gly  Ile  Asp  Tyr  Ser
      130
Lys  Arg  Asp  Ser  Val  Arg  Asn  Cys  Trp  Leu  Gly  Gly  Asp  Leu  Pro  Asp
145  Leu  Asp  Gln  Ser  Arg  Pro  Arg  Val  Gln  Asp  Val  Gln  Arg  Ala  His  Ile
      165
Arg  Lys  Leu  Leu  Ser  Leu  Gly  Ile  Asp  Gly  Phe  Arg  Phe  Asp  Ala  Ala
      180
Lys  His  Ile  Asp  Pro  Ile  Val  Val  Lys  Asp  Tyr  Ile  Asp  Leu  Ile  Asp
      195
Arg  Glu  Ser  Asn  Gly  Arg  Thr  Trp  Asn  Tyr  Leu  Glu  Val  Ile  Glu  Asp
210  Asp  Gly  Thr  Gln  Ala  Thr  Asp  Tyr  Asn  Trp  Ile  Ala  Ala  Val  Thr  Asp
225  Phe  Val  Leu  Tyr  Lys  Glu  Ser  Leu  Arg  Lys  Ala  Phe  Ser  Leu  Gly  Gly
      245
Asp  Leu  Arg  Ser  Leu  Lys  Met  Pro  Val  Ala  Val  Asn  Asp  Ser  Arg  Ser
      260
Ile  Val  Phe  Gly  Arg  Asn  His  Asp  Thr  Val  Pro  Glu  Asn  Asn  Gln  Asn
      275
Cys  Ile  Val  Gly  Cys  Tyr  Asp  Ser  Arg  Glu  Asp  Ser  Tyr  Leu  Ala  Thr
      295
Ala  Tyr  Val  Leu  Ala  Arg  Glu  Ser  Gly  Val  Pro  Leu  Val  Leu  Asn  Trp
305  Asp  Asn  Tyr  Asp  Ala  Pro  Tyr  Ile  Ser  Thr  Gly  Val  Lys  Phe  Arg  Gln
      325
Ile  Met  Thr  Gln  Arg  Gly  Arg  Ser  Ala  Met  Asn  Val  Lys  Glu  Asn  Val
      340
Leu  Gly  Val  Ile  Asp  Ser  Pro  Val  Val  Met  Met  Met  Glu  Arg  Gly  Ser
      355
Glu  Gly  Phe  Phe  Val  Leu  Asn  Lys  Ser  Ala  Asp  Arg  Phe  Asp  Ile  Pro
      370
Val  Leu  Asp  Leu  Thr  Leu  Thr  Asn  Leu  Glu  Gly  Cys  Tyr  Arg  Glu  Leu
385  Arg  Arg  Lys  Phe  Thr  Val  Ala  Ile  Glu  Arg  Lys  Tyr  Gly  Lys  Lys  Phe
      405
Val  Thr  Arg  Trp  Gly  Arg  Trp  Asp  Arg  Gly  Gly  Leu  Glu  Ile  Tyr  Gly
      420
Arg  Asp  Ala  Leu  Tyr  Phe  Ile  Arg  Glu  Pro  Trp  Glu  Gln  Cys  Arg
      435
      440
      445

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<210> 418
 <211> 1455
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 418
atgaaccgctc caggcacggg cgcctcgggg cgcccacaat ctcgctccgc cacgtcgtgg      60
caatcccgcga acggcggctg gctgctcgcc tcgctgctgg ccgtgtgttt cgcaacggcg      120
cccgctgcgcg ccgatgtcat cctgcattga ttcaactggc cgtatgcgac ggtcgaagcc      180
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ccgctcggca atcgcgagtc gttcgtgcgc atgtccagtg ctctgcgcgc ccgcggcatt      360
cgcgctctacg ccgacatcgt gctgaatcac atggccaatg aagccccgca gcgaccgat      420
ctgaactacc ccggtcaacg ggtcctggat cagtatgcag gcaataccgc gtacttcgcg      480
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gccaatcatg gcaatgacat gcaggtgctc gcacacggga actgccatct gctctttcgc 1260
cgcggaatc gcggcatcgt ggcgatcaac aagtgcgggc atacggtcaa cgccacgggtg 1320
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agcgtgggtg agattggcag cgccagccac acgttcagcc ttccgcgcgc gcgcgctcgg 1440
atgtggctgc gctga                                     1455

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<210> 419
 <211> 484
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(44)

```

<400> 419
Met Asn Arg Pro Gly Thr Gly Ala Ser Gly Arg Pro Gln Ser Arg Ser
 1      5      10      15
Ala Thr Ser Trp Gln Ser Arg Asn Gly Trp Leu Leu Ala Ser Leu
 20      25      30
Leu Ala Val Cys Phe Ala Thr Ala Pro Val Arg Ala Asp Val Ile Leu
 35      40      45
His Ala Phe Asn Trp Pro Tyr Ala Thr Val Glu Ala Arg Ala Asn Glu
 50      55      60
Leu Arg Asp Leu Gly Tyr Arg Ala Val Leu Val Ala Pro Pro Val Lys
 65      70      75      80
Ser Glu Gly Asn Ala Trp Trp Ala Arg Tyr Gln Pro Gln Asp Tyr Arg
 85      90      95
Val Ile Glu His Pro Leu Gly Asn Arg Glu Ser Phe Val Arg Met Ser
 100     105     110
Ser Ala Leu Arg Ala Arg Gly Ile Arg Val Tyr Ala Asp Ile Val Leu
 115     120     125
Asn His Met Ala Asn Glu Ala Pro Gln Arg Pro Asp Leu Asn Tyr Pro
 130     135     140
Gly Gln Arg Val Leu Asp Gln Tyr Ala Gly Asn Thr Ala Tyr Phe Ala
 145     150     155     160
Gln Gln Arg Leu Tyr Gly Asp Leu Arg Tyr Asn Phe Met Ser Ala Trp
 165     170     175
Asp Phe Gly Pro Ala His Cys Ile Gly Asn Tyr His Asp Val Trp Gln
 180     185     190
Val Gln Asn Trp Arg Leu Cys Ser Gly Ala Gly Asp Ala Gly Leu Pro
 195     200     205
Asp Leu Leu Ala Ser Asp Tyr Ile Val Gly Gln Gln Arg Thr Tyr Leu
 210     215     220
Gln Ala Leu Lys Asn Leu Gly Val Ser Gly Leu Arg Ile Asp Ala Ala
 225     230     235     240
Lys His Met Pro Leu Ser His Ile Asn Arg Val Leu Thr Ala Asp Leu
 245     250     255
Lys Ala Gly Met His Val Phe Gly Glu Val Ile Thr His Gly Gly Val
 260     265     270
Gly Asp Pro Glu Tyr Asp Leu Phe Leu Arg Pro Tyr Leu Asp Gly Thr
 275     280     285
Asp His Gly Ala Tyr Asp Phe Pro Leu Phe Glu Ala Ile Arg Arg Ala
 290     295     300
Phe Gly Phe Gly Gly Ser Met Ser Thr Leu Val Asp Pro Gly Ala Val

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305      310      315      320
Gly Leu Ala Leu Pro Asn Ala Arg Ser Ile Thr Phe Thr Val Thr His
      325      330      335
Asp Ile Pro Asn Asn Gly Val Phe Arg His Leu Leu Leu Asp Ala Gly
      340      345      350
Asp Glu Thr Leu Ala Tyr Ala Tyr Ile Leu Gly Arg Asp Gly Gly Ser
      355      360      365
Pro Leu Leu Tyr Ser Asp His Asn Glu Ser Gly Asp Asn Arg Trp Val
      370      375      380
His Ala Tyr Arg Arg Asn Asp Leu Ala Ala Met Ile Arg Phe His Asn
385      390      395      400
Ala Asn His Gly Asn Asp Met Gln Val Leu Ala His Gly Asn Cys His
      405      410      415
Leu Leu Phe Arg Gly Asn Arg Gly Ile Val Ala Ile Asn Lys Cys
      420      425      430
Gly His Thr Val Asn Ala Thr Val Asn Met Asn Asn Ser Val Leu Trp
      435      440      445
Trp His Thr Pro Tyr Arg Asp Val Leu Asp Ala Gly Ser Val Val Gln
      450      455      460
Ile Gly Ser Ala Ser His Thr Phe Ser Leu Pro Pro Arg Arg Ala Arg
465      470      475      480
Met Trp Leu Arg

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<210> 420
 <211> 1308
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<400> 420
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ggtgctcacg cggcgaccga acagagcaag gccagtagcg caatcctgtt ccagggcttt      180
cactggcact ccgccaatgg cagctggtac ggcaacctgc agggcaaggc cggtgatctg      240
aaggatctcg gcatcaccca tgtctggttc ccgccgccgt cggacgccgc gtcgctcgaa      300
ggctacctgc gcgccagct taatgtgctg aactccagtt acggcaacga aaccgcgctg      360
aagaacgca ttagcgcgct gaacggccag ggcacgaaga gcgtggttga cgtcgtgatc      420
aaccatcgag tgggtacagc caactgggac gatttcacca acccgacctg ggattgccgc      480
gccgtcacca gcaatgatga atggagcggc cgctgtggca ccgccgacac cggcgatggt      540
tacgatgtcg cgcgcgatct cgaccaccgc cagaccttcc tgcagaacga tctcaagact      600
tggctgtcgt cgcgtctcaa aggtgtgggc ttcagcggca ttcgctacga ctactcgaaa      660
ggctacggac cgggttacgc cgggctttac cagcatgcga tggcgccgaa cttctgcgtc      720
ggcgagatct ggaccaatct tgattacaac aatgtcgatg cccatcgcca gctgctgatg      780
aactacgtcg atggcaacgg cggcaagtgt ggcgcgttcg acttcaccac caagggcctg      840
ctgaatcagg cgctgtcggc caacgaatac tggcgccctgc gcgcttcga tggcaaaccg      900
gccggcggtg tcggctgggt ggcacagaag atggtgacct ttgtcgacaa ccacgacacc      960
gggccgtcgc agagttagcg cagcggccaa aatcactggc cggtgccctg cggcaagggt      1020
atgcagggct atgcctacgt gctgagccat ccgggtgttc cgactgtgta ctaccgcgat      1080
gtctacgact ggaacctgct tgccgcgatc aagacgtga tccagctgcg caaggaaaaa      1140
ggcatcacct cgacctcgtc ggttgcaatc cagcgtgccg atcagggcct ttacgcggcg      1200
atcatcaaca acaacctggc gatgaagatc ggcccgaact catggagtcc tggcagcggc      1260
tggacactgc gcatttcggg cgatcaatac gcgatctgga cccgctga      1308

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<210> 421
 <211> 435
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(44)

<400> 421

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Arg	Phe	Ala	Ser 20	Asn	Lys	Pro	Val	Ile 25	Thr	Arg	Arg	Leu	Gly 30	Ala	Ala
Leu	Ala	Gly 35	Leu	Leu	Phe	Cys	Ala 40	Gly	Ala	His	Ala	Ala 45	Thr	Glu	Gln
Ser	Lys 50	Ala	Ser	Ser	Ala	Ile 55	Leu	Phe	Gln	Gly	Phe 60	His	Trp	His	Ser
Ala 65	Asn	Gly	Ser	Trp	Tyr 70	Gly	Asn	Leu	Gln	Gly 75	Lys	Ala	Gly	Asp	Leu 80
Lys	Asp	Leu	Gly	Ile 85	Thr	His	Val	Trp	Phe 90	Pro	Pro	Pro	Ser	Asp	Ala 95
Ala	Ser	Leu	Glu 100	Gly	Tyr	Leu	Pro	Arg 105	Gln	Leu	Asn	Val	Leu	Asn	Ser
Ser	Tyr	Gly 115	Asn	Glu	Thr	Ala	Leu 120	Lys	Asn	Ala	Ile	Ser 125	Ala	Leu	Asn
Gly	Gln	Gly 130	Ile	Lys	Ser	Val 135	Val	Asp	Val	Val	Ile 140	Asn	His	Arg	Val
Gly 145	Thr	Ala	Asn	Trp	Ala 150	Asp	Phe	Thr	Asn	Pro 155	Thr	Trp	Asp	Cys	Arg 160
Ala	Val	Thr	Ser	Asn 165	Asp	Glu	Trp	Ser	Gly 170	Arg	Cys	Gly	Thr	Ala 175	Asp
Thr	Gly	Asp	Gly 180	Tyr	Asp	Ala	Ala	Arg 185	Asp	Leu	Asp	His	Arg 190	Gln	Thr
Phe	Leu	Gln 195	Asn	Asp	Leu	Lys	Thr 200	Trp	Leu	Ser	Ser	Arg 205	Leu	Lys	Gly
Val	Gly 210	Phe	Ser	Gly	Ile	Arg 215	Tyr	Asp	Tyr	Ser	Lys 220	Gly	Tyr	Gly	Pro
Gly 225	Tyr	Ala	Gly	Leu	Tyr 230	His	Asp	Ala	Met	Ala 235	Pro	Asn	Phe	Cys	Val 240
Gly	Glu	Ile	Trp	Thr 245	Asn	Leu	Asp	Tyr	Asn 250	Asn	Val	Asp	Ala	His 255	Arg
Gln	Leu	Leu	Met 260	Asn	Tyr	Val	Asp	Gly 265	Asn	Gly	Gly	Lys	Cys 270	Gly	Ala
Phe	Asp	Phe 275	Thr	Thr	Lys	Gly	Leu 280	Leu	Asn	Gln	Ala	Leu 285	Ser	Ala	Asn
Glu	Tyr 290	Trp	Arg	Leu	Arg	Ala 295	Ser	Asp	Gly	Lys	Pro 300	Ala	Gly	Gly	Ile
Gly 305	Trp	Trp	Ala	Gln	Lys 310	Met	Val	Thr	Phe	Val 315	Asp	Asn	His	Asp	Thr 320
Gly	Pro	Ser	Gln	Ser 325	Cys	Gly	Ser	Gly	Gln 330	Asn	His	Trp	Pro	Val 335	Pro
Cys	Gly	Lys	Val 340	Met	Gln	Gly	Tyr	Ala 345	Tyr	Val	Leu	Ser	His 350	Pro	Gly
Val	Pro	Thr 355	Val	Tyr	Tyr	Pro	His 360	Val	Tyr	Asp	Trp	Asn 365	Leu	Arg	Ala
Ala	Ile 370	Lys	Thr	Leu	Ile	Gln 375	Leu	Arg	Lys	Glu	Lys 380	Gly	Ile	Thr	Ser
Thr 385	Ser	Ser	Val	Ala	Ile 390	Gln	Arg	Ala	Asp	Gln 395	Gly	Leu	Tyr	Ala	Ala 400
Ile	Ile	Asn	Asn	Asn 405	Leu	Ala	Met	Lys	Ile 410	Gly	Pro	Asn	Ser	Trp 415	Ser
Pro	Gly	Ser	Gly 420	Trp	Thr	Leu	Arg	Thr 425	Ser	Gly	Asp	Gln	Tyr 430	Ala	Ile
Trp	Thr	Arg 435													

<210> 422
 <211> 2751
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 422
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60
 120

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gacttttaag gactttattca gaaactcgac tacattcagg accttggatt taccgcaata 300
tggaataaccc ctatcgtgca taatcgagc ccgctggact accacggcta ccatgcctgg 360
gactttacac gcgtggatcc gcgcctggaa tcacccggag ccacctttca ggacctcatc 420
aacgctgtac atgcccgggg gatgaaaatc gtgctggata ttgtaaccaa tcaactccggc 480
cggtttgga ttaaggatth tgccgaaatc aagtataata ccgatcccaa ccgcccattg 540
ggacaagacg cacagggcaa cccccgtcag gacaacccca actgggaata cgacggactt 600
accccaaac ccgatgatgg caaaatctgg agtcgtgcc aacctggccag gttacctgcc 660
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caggtgctgc aatacttgct cgacgcctac cgtaccttta ttgaaatggg tgtggacggt 900
ttccgctggg acaccattaa gcatatgagc cgcagcgtat tcctgtgggt tctggacgaa 960
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tccgaaccct ccgggatgtc cgttatcgac ttttttgcca tggccacctt ccacttgttc 1140
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tggcggggaa ttctttccct ttattatggc acagaagtcc aatttatgct tggcgcttt 1380
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ggtgtgcatg ccctgcgctt cgtgcacgc cacttttcta ccggtatgta cctgtaccga 2700
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<210> 423
 <211> 916
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(27)

<400> 423
 Met Lys Thr Ser Tyr Ser Phe Val Tyr Thr Ile Leu Ile Val Phe Val
 1 5 10 15
 Leu Ile Pro Leu Ala Pro Val Arg Val Ser Ala Gln Val Asp Phe Arg
 20 25 30
 Glu Glu Thr Ile Tyr Phe Leu Leu Thr Thr Arg Phe Phe Asp Gly Asp
 35 40 45
 Ser Ser Asn Asn Val Pro Asn Glu Trp Ser Ser Tyr His Pro Asp Pro
 50 55 60
 Asn Ile Asn Pro Ser Ile Thr Asp Pro Gln Asp Val Thr Trp Arg Gly
 65 70 75 80
 Asp Phe Lys Gly Leu Ile Gln Lys Leu Asp Tyr Ile Gln Asp Leu Gly

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625	Val	Thr	Leu	Thr	Tyr	630	Asp	Gly	Phe	Leu	Asn	635	Ser	Gln	Gln	Gln	Val	640	Asn
				645	Tyr	Asp	Gly	Trp	Thr	Asn	Val	Ser	Thr	Thr	Pro	Met			
	Leu	His	Trp	Gly	660	Asp	Val	Trp	Lys	680	Phe	Asn	Asn	Gly	685	Asp	Val	Trp	
	Gln	Lys	Asp	Gly	675	Leu	Asn	Phe	Val	695	Thr	Leu	Ala	Ser	Pro	Asp			
	Ala	Thr	Ser	Arg	690	Asn	Asn	Tyr	Ser	710	Val	Thr	Ala	Asn	Pro	Asp	Pro	Ala	Val
	Asp	Asn	Asn	Asn	705	Pro	Val	Phe	Thr	730	Ala	Asn	Pro	Asp	Pro	Ala	Val		
	Gly	Ser	Thr	Pro	725	Leu	Thr	Tyr	745	Gln	Gly	Ser	Leu	Ala	Asn	Val	Thr		
	Ala	Gly	Glu	Pro	740	His	Trp	Gly	760	Asp	Val	Trp	Thr	Leu	Thr	Met	Thr		
	Ser	Ala	Val	Asn	755	Lys	Ser	Gly	775	Arg	Glu	Leu	Asn	Phe	Val	Phe	Asn	Asp	Gly
	Asn	Thr	Pro	Met	770	Thr	Arg	Glu	790	Asn	Asn	Gly	Gly	Gln	Asp	Tyr	Leu	Leu	Asn
	Val	Pro	Ala	Thr	785	Asp	Asn	Asn	805	Ala	Asp	Phe	Glu	825	Pro	Phe	Pro	Asn	Ala
	Asn	Asn	Thr	Trp	820	Ser	Ala	Asp	840	Pro	Phe	Asn	Pro	Thr	Thr	Ser	Ile		
	Thr	Thr	Thr	Thr	835	Gly	Asn	Tyr	855	Gly	Gly	Thr	Val	Arg	Ile	Gln	Ile	Phe	Asp
	Arg	Leu	Leu	Gly	850	Ile	Glu	Glu	870	Leu	Ser	Ala	Ala	Tyr	Pro	Gly	Ala		
	Ser	Phe	Glu	Ile	865	Val	Ser	Gly	Arg	885	Leu	Arg	His	Leu	Ser	Thr	Gly	Met	
	Val	Ser	Gly	Arg	875	Ala	Thr	Phe	Gly	900	Ala	Thr	Phe	Gly	Gly	His	Val	Leu	Thr
	Gly	Val	His	Ala	885	Arg	Ala	Thr	Phe	905	Gly	Gly	His	Val	Leu	Thr	Gly	Lys	Met
	Tyr	Leu	Tyr	Arg	900	Lys													
	Thr	Leu	Val	Lys	915														

<210> 424
 <211> 2169
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 424																			
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cgctttctcg	acggggatgc	ctccaataac	ccgaccggag	cggcctatga	ctccaccggc														240
accaatctga	agctgtatct	cggcggcgac	tggaaggggc	tgaccgataa	gctgaatgat														300
aattatttta	cggatctggg	cattacggca	ctctggatct	cccagcctgt	cgagaatatt														360
tattcggttg	tcaactattc	cggagtgaac	agcacttcct	accatggcta	ctgggcccgc														420
gacttcaaga	agaccaaccc	ctatttcggc	tcgatggccg	atttcagac	tctcgtaaat														480
accgctcatg	caaaaggcat	caagatcgtc	atcgactttg	ctccgaatca	tacctctcct														540
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ctgggaggct	acacgaatga	cacgaacgga	tttttccatc	acaacggcgg	ctccgatttc														660
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gattccatgc	tcaccaatac	agctgctgac	tacaccaggg	tcaacgatca	agtaaccttt														1080
attgataatc	atgatatgga	ccgcttcaag	accggcacgc	tgaacaaccg	ccgtctggag														1140
caggctctgg	cttcactct	gacctccgc	ggcgcttcctg	ccatttatta	cggaacagaa														1200
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1001827087_1.txt

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actcaggttt ccgtacgctt tatcgtgaac aacgccacta catcgcttgg ccaaaatgta 1920
tatctaccg gcaatgtagc cgagctcggg aactggacta caggcagcag tgcaatcggg 1980
ccgttggttca accagatcat taaagtctac cctacctggt attatgatgt cagcggttccg 2040
gccggcactg cgctcgaatt caaatttttc aagaaaagcg gttcgaccgt tacctgggag 2100
agcggctcta atcatacggt caccgcctct gcgagcggaa ccgcaactgt aaccgtggat 2160
tggcagtaa

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<210> 425
 <211> 722
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(37)

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<400> 425
Met Asp Ser Met Phe Lys Gln Val Lys Arg Leu Phe Leu Ser Leu Ala
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Leu Thr Leu Gly Leu Leu Thr Gly Ser Ala Leu Pro Leu Val Pro Val
20      25      30
Ser Gln Ala Phe Ala Asp Pro Asp Thr Ala Val Ser Asn Lys Ala Ser
35      40      45
Phe Ser Thr Asp Val Ile Tyr Gln Val Phe Thr Asp Arg Phe Leu Asp
50      55      60
Gly Asp Ala Ser Asn Asn Pro Thr Gly Ala Ala Tyr Asp Ser Thr Gly
65      70      75      80
Thr Asn Leu Lys Leu Tyr Leu Gly Gly Asp Trp Lys Gly Leu Thr Asp
85      90      95
Lys Leu Asn Asp Asn Tyr Phe Thr Asp Leu Gly Ile Thr Ala Leu Trp
100     105     110
Ile Ser Gln Pro Val Glu Asn Ile Tyr Ser Val Val Asn Tyr Ser Gly
115     120     125
Val Asn Ser Thr Ser Tyr His Gly Tyr Trp Ala Arg Asp Phe Lys Lys
130     135     140
Thr Asn Pro Tyr Phe Gly Ser Met Ala Asp Phe Gln Thr Leu Val Asn
145     150     155     160
Thr Ala His Ala Lys Gly Ile Lys Ile Val Ile Asp Phe Ala Pro Asn
165     170     175
His Thr Ser Pro Ala Met Glu Thr Asp Thr Ser Phe Ala Glu Asn Gly
180     185     190
Lys Leu Tyr Asp Asn Gly Thr Leu Leu Gly Gly Tyr Thr Asn Asp Thr
195     200     205
Asn Gly Phe Phe His His Asn Gly Gly Ser Asp Phe Ser Thr Leu Glu
210     215     220
Asn Gly Ile Tyr Lys Asn Leu Tyr Asp Leu Ala Asp Leu Asn His Asn
225     230     235     240
Asn Ser Thr Ile Asp Lys Tyr Phe Lys Asp Ala Ile Lys Val Trp Ile
245     250     255
Asp Thr Gly Val Asp Gly Ile Arg Val Asp Ala Val Lys His Met Pro
260     265     270
Asp Gly Trp Gln Lys Asn Trp Val Ser Ser Ile Tyr Asn Tyr Glu Pro
275     280     285
Val phe Ile Phe Gly Glu Trp Tyr Leu Gly Ser Ser Ala Ala Asp Ala
290     295     300
Asp Asn Thr Lys Phe Ala Asn Thr Ser Gly Met Ser Leu Leu Asp Phe

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305	Arg	Phe	Asn	Gln	Glu	310	Val	Arg	Asn	Val	Phe	315	Arg	Asn	Asp	Thr	Ser	320	Thr
				325							330						335		
Met	Tyr	Ala	Leu	Asp	Ser	Met	Leu	Thr	Asn	Thr	Ala	Ala	Asp	Tyr	Thr				
			340					345							350				
Gln	Val	Asn	Asp	Gln	Val	Thr	Phe	Ile	Asp	Asn	His	Asp	Met	Asp	Arg				
		355					360					365							
Phe	Lys	Thr	Gly	Thr	Leu	Asn	Asn	Arg	Arg	Leu	Glu	Gln	Ala	Leu	Ala				
	370					375					380								
Phe	Thr	Leu	Thr	Ser	Arg	Gly	Val	Pro	Ala	Ile	Tyr	Tyr	Gly	Thr	Glu				
385					390					395					400				
Gln	Tyr	Met	Thr	Gly	Asn	Gly	Asp	Pro	Asp	Asn	Arg	Ala	Met	Met	Thr				
			405						410					415					
Ser	Phe	Ser	Thr	Ser	Thr	Thr	Ala	Phe	Asn	Val	Ile	Ser	Lys	Leu	Ala				
			420					425					430						
Pro	Leu	Arg	Lys	Ser	Asn	Pro	Ala	Ile	Ala	Tyr	Gly	Thr	Thr	Gln	Gln				
		435					440						445						
Arg	Trp	Ile	Asn	Asn	Asp	Val	Tyr	Ile	Tyr	Glu	Arg	Lys	Phe	Gly	Asn				
	450				455						460								
Asn	Val	Ala	Val	Val	Ala	Val	Asn	Arg	Asn	Leu	Ser	Thr	Ala	Thr	Ser				
465					470					475					480				
Ile	Ser	Gly	Leu	Val	Thr	Ser	Leu	Pro	Ser	Gly	Thr	Tyr	Ser	Asp	Val				
			485					490						495					
Leu	Gly	Gly	Thr	Leu	Ser	Gly	Asn	Ser	Ile	Thr	Ala	Ser	Ser	Gly	Asn				
			500					505					510						
Val	Ala	Thr	Phe	Thr	Leu	Gly	Ala	Gly	Ala	Ala	Ala	Val	Trp	Gln	Tyr				
		515					520					525							
Thr	Glu	Thr	Ser	Ser	Ala	Thr	Pro	Val	Leu	Gly	His	Val	Gly	Pro	Met				
	530				535						540								
Met	Gly	Gln	Pro	Gly	Asn	Glu	Val	Thr	Ile	Asp	Gly	Arg	Gly	Phe	Gly				
545					550					555					560				
Ser	Ser	Ala	Gly	Thr	Val	Tyr	Phe	Gly	Thr	Thr	Ala	Val	Thr	Gly	Ser				
			565					570					575						
Asn	Ile	Ile	Ser	Trp	Glu	Asp	Thr	Gln	Ile	Lys	Val	Lys	Val	Pro	Ser				
			580					585					590						
Val	Ala	Ala	Gly	Thr	Tyr	Ala	Val	Lys	Val	Thr	Asn	Ser	Ala	Gly	Thr				
		595					600					605							
Ser	Asn	Ser	Tyr	Ser	Gly	Phe	Thr	Ile	Leu	Thr	Gly	Thr	Gln	Val	Ser				
	610					615					620								
Val	Arg	Phe	Ile	Val	Asn	Asn	Ala	Thr	Thr	Ser	Leu	Gly	Gln	Asn	Val				
625					630					635					640				
Tyr	Leu	Thr	Gly	Asn	Val	Ala	Glu	Leu	Gly	Asn	Trp	Thr	Thr	Gly	Ser				
			645					650					655						
Ser	Ala	Ile	Gly	Pro	Leu	Phe	Asn	Gln	Ile	Ile	Lys	Val	Tyr	Pro	Thr				
			660					665					670						
Trp	Tyr	Tyr	Asp	Val	Ser	Val	Pro	Ala	Gly	Thr	Ala	Leu	Glu	Phe	Lys				
		675					680					685							
Phe	Phe	Lys	Lys	Ser	Gly	Ser	Thr	Val	Thr	Trp	Glu	Ser	Gly	Ser	Asn				
	690					695					700								
His	Thr	Phe	Thr	Ala	Pro	Ala	Ser	Gly	Thr	Ala	Thr	Val	Thr	Val	Asp				
705					710					715					720				
Trp	Gln																		

<210> 426
 <211> 2139
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 426	
atgattagaa aattcacaat tgcgggtatc gttaacgaga atacgatcca gtacaaggta	60
acgcccgaaa cacttatctc cgatattgag gacataaaag tgtacgacgg cgatcgggag	120
attaagggtta atgaaatctc tactctcggg aagaaatccg ttatgggata cgtaacgctt	180
gacgagtcgc ttgatctttc gaagaattac agaataaggg tgaaaggata cggctcgaag	240
atcgctgttc ctaccgatat ttttgacagc gagtacttcg ctgaaacgta ccactatgac	300

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ggagacgcatc tcggagctgt gataaacgga gataacaaca cattaaggt ttgggctccg 360
actgctctgt aggttattct caatctcttt gaaaaaggcg acggagtaga ggcttacaag 420
agcgtttcga tgactcgcgg tgaaaaggcg gtgtggtcac atacagagga atgcgggtcat 480
ggaacgtact atacttatac tgtctcgaca tccgcagggtg ttcaggaagc tgttgaccca 540
tatgctaaat ccgcagggtgt gaacggaaac cgcggaatgg ttgttgacct ttccctgaca 600
aatcccgaag gatgggaaaa cgatcagctt gagaataaga tagagaagta ttcagatgca 660
ataatctggg aagtacacgt tcgtgacttc tcgaacaaga tcgaatcttc cgaatataag 720
ggcaagtacc ttgctgtcac ggagagaggt cttgttaacg agcacggtaa atctgtcgg 780
gttgattacc ttgtagacct tggataaaca cacgttcac tgcttcctgt atatgactat 840
gctacagtag atgaatcctc gactgagccg cagtttaact ggggatacga cccgaagaat 900
tacaacgttc ccgagggcag ctattcaaca gatccgtaca acggcgagggt tcgtattcgc 960
gagttcaagg aaatggtaaa ggcactccac gaagcgggca ttggtgttat catggacgta 1020
gtatacaatc atactcatga cgaaacagc tcgttcaata agatagttcc gtactactat 1080
tacagataca cctcctcagg cgtaactca tctgcaagcg gatgcgaaa cgataccgcc 1140
tccgagagat atatgtacgg caagttcatg gttgagttca ctgcatattg ggtaagtga 1200
tacaaccttg acggacttcg cttcgacctt atgggactcc atgacgtaga aacgatgcag 1260
gaggttgaga gtgcagttca cgccatcaat ccgaatgcga tcatttacgg cgaaggatgg 1320
acaatgggcg cgactgttga cggagcgct caggcaaatc agtcaaata ctcaaagatt 1380
acgcctacag gtgacgcgat cgggtgcagtc gctgtattca acgacgtgat cagagacgg 1440
ctcaagggaa gcgtatttga aaaaacaggc cgcggcttta tcaacggatc tgccaagaat 1500
acaataaaca aggttatgtt cggtatcaga ggcggtcagg gtgttggtca gggctggaca 1560
gttgaagacg gaatggatc caactatatg agtgctcacg ataacaatac gctgtgggat 1620
aagctccttc tctcgaatcc taacgtgagc gacgagaaga gaaacgatat gaataacctt 1680
ggcggccgaa tcatcatgat ctctcgcgga actccgttct ggcaggcggg agaggaaatg 1740
ctccgcacaa agggaggaga cgagaacagc tacaagtcca gtgatgagg aaacaatatc 1800
aactggtccg tacttgcgta cggtagcgcg gagtatgaaa caatgcagta ttacaagggt 1860
ctcatcgaga tgcgtaaggc atacggatta ttctcagata tgacaaccga gatcactcac 1920
accgactccg gcaacggaat cttacaata agtatcaaag cacagaacgg tgaagaagca 1980
ctcgtaatca ttaatccgaa cgagtttgag tacacgcatg agcttgacgg cgagtggaaa 2040
ctcatatgtg acgctgaccg tgcaggtgca gaggttctcg aggctgagag cggaagcggt 2100
aaggttggcg cgatcagtat taaggtttac gtaaagtga 2139

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<210> 427

<211> 712

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 427

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Met Ile Arg Lys Phe Thr Ile Ala Gly Ile Val Asn Glu Asn Thr Ile
1      5      10      15
Gln Tyr Lys Val Thr Pro Glu Thr Leu Ile Ser Asp Ile Glu Asp Ile
20     25     30
Lys Val Tyr Asp Gly Asp Arg Glu Ile Lys Val Asn Glu Ile Ser Thr
35     40     45
Leu Gly Lys Lys Ser Val Met Gly Tyr Val Thr Leu Asp Glu Ser Leu
50     55     60
Asp Leu Ser Lys Asn Tyr Arg Ile Glu Val Lys Gly Tyr Gly Ser Lys
65     70     75     80
Ile Ala Val Pro Thr Asp Ile Phe Asp Ser Glu Tyr Phe Ala Glu Thr
85     90     95
Tyr His Tyr Asp Gly Asp Asp Leu Gly Ala Val Ile Asn Gly Asp Thr
100    105    110
Thr Thr Phe Lys Val Trp Ala Pro Thr Ala Ser Glu Val Ile Leu Asn
115    120    125
Leu Phe Glu Lys Gly Asp Gly Val Glu Ala Tyr Lys Ser Val Ser Met
130    135    140
Thr Arg Gly Glu Lys Gly Val Trp Ser His Thr Glu Glu Cys Gly His
145    150    155    160
Gly Thr Tyr Tyr Thr Tyr Thr Val Ser Thr Ser Ala Gly Val Gln Glu
165    170    175
Ala Val Asp Pro Tyr Ala Lys Ser Ala Gly Val Asn Gly Asn Arg Gly
180    185    190
Met Val Val Asp Leu Ser Leu Thr Asn Pro Glu Gly Trp Glu Asn Asp
195    200    205
Gln Leu Glu Asn Lys Ile Glu Lys Tyr Ser Asp Ala Ile Ile Trp Glu

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Val 225	His 210	Val	Arg	Asp	Phe 230	Ser 215	Asn	Lys	Ile	Glu 235	Ser 220	Ser	Glu	Tyr	Lys 240
Gly	Lys	Tyr	Leu	Ala 245	Phe	Thr	Glu	Arg	Gly 250	Leu	Val	Asn	Glu	His 255	Gly
Lys	Ser	Val	Gly 260	Val	Asp	Tyr	Leu	Val 265	Asp	Leu	Gly	Ile	Thr 270	His	Val
His	Leu	Leu	Pro	Val	Tyr	Asp	Tyr 280	Ala	Thr	Val	Asp	Glu 285	Ser	Ser	Thr
Glu 290	Pro	Gln	Phe	Asn	Trp	Gly 295	Tyr	Asp	Pro	Lys	Asn	Tyr	Asn	Val	Pro
Glu 305	Gly	Ser	Tyr	Ser	Thr 310	Asp	Pro	Tyr	Asn	Gly 315	Glu	Val	Arg	Ile	Arg 320
Glu	Phe	Lys	Glu	Met 325	Val	Lys	Ala	Leu	His 330	Glu	Ala	Gly	Ile	Gly 335	Val
Ile	Met	Asp	Val 340	Val	Tyr	Asn	His	Thr 345	His	Asp	Ala	Asn	Ser 350	Ser	Phe
Asn	Lys	Ile 355	Val	Pro	Tyr	Tyr	Tyr 360	Tyr	Arg	Tyr	Thr	Ser 365	Ser	Gly	Ala
Asn	Ser 370	Ser	Ala	Ser	Gly	Cys 375	Gly	Asn	Asp	Thr	Ala 380	Ser	Glu	Arg	Tyr
Met 385	Tyr	Gly	Lys	Phe	Met 390	Val	Glu	Ser	Thr	Ala 395	Tyr	Trp	Val	Ser	Glu 400
Tyr	Asn	Leu	Asp	Gly 405	Leu	Arg	Phe	Asp	Leu 410	Met	Gly	Leu	His	Asp 415	Val
Glu	Thr	Met	Gln 420	Glu	Val	Glu	Ser	Ala 425	Val	His	Ala	Ile	Asn 430	Pro	Asn
Ala	Ile	Ile 435	Tyr	Gly	Glu	Gly	Trp 440	Thr	Met	Gly	Ala	Thr 445	Val	Asp	Gly
Ser	Ala 450	Gln	Ala	Asn	Gln	Ser 455	Asn	Ile	Ser	Lys	Ile 460	Thr	Pro	Thr	Gly
Asp 465	Ala	Ile	Gly	Ala	Val 470	Ala	Val	Phe	Asn	Asp 475	Val	Ile	Arg	Asp	Gly 480
Leu	Lys	Gly	Ser	Val 485	Phe	Glu	Lys	Thr	Gly 490	Arg	Gly	Phe	Ile	Asn 495	Gly
Ser	Ala	Lys	Asn 500	Thr	Ile	Asn	Lys	Val 505	Met	Phe	Gly	Ile	Arg 510	Gly	Gly
Gln	Gly	Val 515	Gly	Gln	Gly	Trp	Thr 520	Val	Glu	Asp	Gly	Met 525	Val	Ile	Asn
Tyr 530	Met	Ser	Ala	His	Asp	Asn 535	Asn	Thr	Leu	Trp	Asp 540	Lys	Leu	Leu	Leu
Ser 545	Asn	Pro	Asn	Val	Ser 550	Asp	Glu	Lys	Arg	Asn 555	Asp	Met	Asn	Asn	Leu 560
Gly	Ala	Ala	Ile	Ile 565	Met	Ile	Ser	Arg	Gly 570	Thr	Pro	Phe	Trp	Gln 575	Ala
Gly	Glu	Glu	Met 580	Leu	Arg	Thr	Lys	Gly 585	Gly	Asp	Glu	Asn	Ser 590	Tyr	Lys
Ser	Ser	Asp 595	Glu	Val	Asn	Asn	Ile 600	Trp	Ser	Val	Leu 605	Ala	Asp	Gly	
Thr	Arg 610	Glu	Tyr	Glu	Thr	Met 615	Gln	Tyr	Tyr	Lys	Gly 620	Leu	Ile	Glu	Met
Arg 625	Lys	Ala	Tyr	Gly	Leu 630	Phe	Ser	Asp	Met	Thr 635	Thr	Glu	Ile	Thr	His 640
Thr	Asp	Ser	Gly	Asn 645	Gly	Ile	Leu	Thr	Ile 650	Ser	Ile	Lys	Ala	Gln 655	Asn
Gly	Glu	Glu	Ala 660	Leu	Val	Ile	Ile	Asn 665	Pro	Asn	Glu	Phe	Glu 670	Tyr	Thr
His	Glu	Leu 675	Asp	Gly	Glu	Trp	Lys 680	Leu	Ile	Cys	Asp	Ala 685	Asp	Arg	Ala
Gly	Ala	Glu	Val	Leu	Glu	Ala 695	Glu	Ser	Gly	Ser	Val 700	Lys	Val	Gly	Ala
Ile 705	Ser	Ile	Lys	Val	Tyr 710	Val	Lys								

<210> 428
 <211> 1425
 <212> DNA
 <213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 428

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tatgatgaag	ttgctgctaa	agcgacagaa	ataaaaaatc	tgggttacaa	ggcggatttg	180
gtagcgccgc	cgttaaaatc	gaacgcgggt	aattgtgcct	ggtggcagcg	ttatcagccg	240
caggatatcc	gtgtaattga	tcattgcaaa	ggtaacaaac	aatcgtttgt	aaatatgata	300
aatgccctga	acgatgccaa	cccggcgcg	aaagttgatg	tatacgccga	tatcgattta	360
aaccatatgg	caaatgagcg	taacggcgcg	actgactttc	ccggccaggc	tgcatgaaat	420
agctatggca	gtaacagcag	ttactggaat	aaccagcgcc	tggttggaac	tttaaccagg	480
gggtgtgttc	gcgccagcga	ctttaaccgg	gctaactgta	tcagcaacta	taacgatgta	540
tggcaggtgc	agaactggcg	gctgtgtggt	ggtgcgggtg	atgccggttt	gccggatctc	600
aaacccaaca	gctgggtggg	gcagcaacag	cgtaactact	taactgcatt	aaaagactta	660
ggcgtaaaag	gctttagggg	tgatgcggcc	aagcatatga	ccatttgga	tattaatgaa	720
attttcacca	gcagcattaa	aaatggcatg	tacgtgtttg	gtgaaattat	taccagcggt	780
ggagccggta	ataacgaata	cgatagcttc	ttatcgcttc	acctggctta	taccgatcat	840
aaagccatag	attttcgct	gtttagcgct	attcgcagtg	cttttggttt	tggcggtagc	900
ttaagccaac	tgggttaacc	gttgtctaac	ggccaggcgc	tgcaaaacag	ccgggcagta	960
acctttacta	ttacgcatga	tattccctacc	aatgacgggt	ttcgttatct	gattatggat	1020
gctaccgacg	agtatctggc	ctatgcctac	attatgggcc	gtgatgggtg	taaaccgctg	1080
atattcagcg	acagcaccgg	cactgataac	aaccgctggg	ttaatgccta	taaagccgat	1140
catattagca	aaatgctgaa	tttcataaac	cgatgcagg	ggcagggcat	ggaaatgtta	1200
gcctgggaat	actgtgctat	ttgtttccgc	gcgcggccag	aaggcggtgt	aggtattaat	1260
aaatgcagca	gcaaccagag	ctttaacatt	aacaccaacg	gccgctttta	ctggtaccgt	1320
aattaccgcg	atgtgttaag	tggcggtaat	ctggtgtata	tcaatggcgg	cagttacaac	1380
ttcagcattc	cggcgcgta	ggccagaatg	tggtatgccg	actaa		1425

<210> 429

<211> 474

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> SIGNAL

<222> (1)...(30)

<400> 429

Met	Gln	Val	Ile	Ser	Thr	Met	Leu	Lys	Arg	Gln	Leu	Leu	Trp	Leu	Ala
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Phe	Val	Val	Ser	Ser	Ala	Val	Phe	Ser	Ser	Ala	Ala	Val	Ala	Asp	Ala
			20					25					30		
Ile	Leu	His	Ala	Phe	Asp	Trp	His	Tyr	Asp	Glu	Val	Ala	Ala	Lys	Ala
		35					40					45			
Thr	Glu	Ile	Lys	Asn	Leu	Gly	Tyr	Lys	Ala	Val	Leu	Val	Ala	Pro	Pro
	50					55					60				
Leu	Lys	Ser	Asn	Ala	Gly	Asn	Cys	Ala	Trp	Trp	Gln	Arg	Tyr	Gln	Pro
65					70				75					80	
Gln	Asp	Ile	Arg	Val	Ile	Asp	His	Cys	Lys	Gly	Asn	Lys	Gln	Ser	Phe
				85					90					95	
Val	Asn	Met	Ile	Asn	Ala	Leu	Asn	Asp	Ala	Asn	Pro	Ala	Arg	Lys	Val
			100					105					110		
Asp	Val	Tyr	Ala	Asp	Ile	Val	Leu	Asn	His	Met	Ala	Asn	Glu	Arg	Asn
		115					120					125			
Gly	Ala	Thr	Asp	Phe	Pro	Gly	Gln	Ala	Ala	Val	Asn	Ser	Tyr	Gly	Ser
	130					135					140				
Asn	Ser	Ser	Tyr	Trp	Asn	Asn	Gln	Arg	Leu	Phe	Gly	Asn	Leu	Thr	Gln
145					150					155					160
Gly	Leu	Phe	Gly	Ala	Ser	Asp	Phe	Asn	Pro	Ala	Asn	Cys	Ile	Ser	Asn
			165						170					175	
Tyr	Asn	Asp	Val	Trp	Gln	Val	Gln	Asn	Trp	Arg	Leu	Cys	Gly	Gly	Ala
			180					185					190		
Gly	Asp	Ala	Gly	Leu	Pro	Asp	Leu	Asn	Pro	Asn	Ser	Trp	Val	Val	Gln
		195					200					205			

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Gln Gln Arg Asn Tyr Leu Thr Ala Leu Lys Asp Leu Gly Val Lys Gly
 210 215 220
 Phe Arg Val Asp Ala Ala Lys His Met Thr Ile Trp His Ile Asn Glu
 225 230 235 240
 Ile Phe Thr Ser Ser Ile Lys Asn Gly Met Tyr Val Phe Gly Glu Ile
 245 250 255
 Ile Thr Ser Gly Gly Ala Gly Asn Asn Glu Tyr Asp Ser Phe Leu Ser
 260 265 270
 Pro Tyr Leu Ala Tyr Thr Asp His Lys Ala Tyr Asp Phe Pro Leu Phe
 275 280 285
 Ser Ala Ile Arg Ser Ala Phe Gly Phe Gly Gly Ser Leu Ser Gln Leu
 290 295 300
 Val Asn Pro Leu Ser Asn Gly Gln Ala Leu Gln Asn Ser Arg Ala Val
 305 310 315 320
 Thr Phe Thr Ile Thr His Asp Ile Pro Thr Asn Asp Gly Phe Arg Tyr
 325 330 335
 Leu Ile Met Asp Ala Thr Asp Glu Tyr Leu Ala Tyr Ala Tyr Ile Met
 340 345 350
 Gly Arg Asp Gly Gly Lys Pro Leu Ile Phe Ser Asp Ser Thr Gly Thr
 355 360 365
 Asp Asn Asn Arg Trp Val Asn Ala Tyr Lys Ala Asp His Ile Ser Lys
 370 375 380
 Met Leu Asn Phe His Asn Arg Met Gln Gly Gln Gly Met Glu Met Leu
 385 390 395 400
 Ala Trp Asn Asp Cys Ala Ile Leu Phe Arg Gly Gln Glu Gly Val
 405 410 415
 Val Gly Ile Asn Lys Cys Ser Ser Asn Gln Ser Phe Asn Ile Asn Thr
 420 425 430
 Asn Gly Arg Phe Tyr Trp Tyr Arg Asn Tyr Arg Asp Val Leu Ser Gly
 435 440 445
 Gly Asn Leu Val Tyr Ile Asn Gly Gly Ser Tyr Asn Phe Ser Ile Pro
 450 455 460
 Ala Arg Gln Ala Arg Met Trp Tyr Ala Asp
 465 470

<210> 430

<211> 1650

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 430

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acaatgatgc	aatatttcga	atggtacctt	ccagacgatg	gaacactatg	gacgaaaagta	180
gcaataaacg	ctcaatcttt	agcgaatctt	ggcattactg	ccctttggct	ttccccctgcc	240
tataaaaggaa	caagcagcag	tgacgttgga	tatggcgttt	atgatttata	tgaccttgga	300
gagtttaatc	aaaaaggaaac	tgtccgaaca	aaatacggga	caaaaacaca	atatatccaa	360
gcaatccaag	cggcgcatac	agcagggatg	cgagtatatg	cagatgtcgt	ctttaaccat	420
aaagccggtg	cagatggaac	agaactagtc	gatgcagtag	aagtaaattc	ttctgaccgc	480
aatcaagaaa	tatcaggaac	atatcaaattc	caagcgtgga	caaaatttga	ttttcctggt	540
cgtggaaaca	cctattctag	ttttaaatgg	cgttggtatc	atttcgatgg	aacggactgg	600
gatgagagta	gaaaactaaa	tcgtattttac	aagttccgcg	gcacgggaaa	agcatgggat	660
tgggaagtag	atacagaaaa	cggaatttat	gactatctca	tgtatgcaga	tttagatatg	720
gatcatccag	aggttgatc	cgaactaaaa	aattggggaa	agtggatatgt	aaccacaacc	780
aatatcgacg	gattccgtct	ggatgcagtg	aagcatatta	aatatagctt	tttcccgga	840
tggctatcgt	acgtacgaac	ccaaacacaa	aagcctcttt	ttgccgttgg	ggaatttttg	900
agctatgaca	ttagcaagtt	gcacaactat	attacaaaga	cgaacggctc	tatgtcccta	960
ttcgatgcc	cgtgcataa	caatttttat	atagcatcga	aatcaagcgg	ttattttgat	1020
atgcgcacat	tactcaacaa	cacattgatg	aaagatcagc	ctacatttagc	agtcacatta	1080
gtggataatc	acgatactga	gccagggcaa	tctctgcagt	catgggtcga	gccatgggtt	1140
aaaccgttag	cttacgcatt	tatcttgacc	cgccaagaag	ggtatccttg	cgtcttttat	1200
ggagattact	atggtattcc	aaaatacaac	attcctgcgc	tgaaaagcaa	acttgatccg	1260
ctgttaattg	ccagaagaga	ttatgcctat	ggaacacagc	acgactatat	tgacagtgcg	1320
gatattatcg	gttggagcgc	ggaaggagtg	gctgaaaaag	caaattcagg	actggctgca	1380
ctcattaccg	acgggcctgg	cggaagcaaa	tggtatgatg	ttggaaaaca	acacgctggc	1440

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aaaacgtttt atgatttaac cggcaatcga agtgatacag tgacaatcaa tgctgatgga 1500
 tggggagaat ttaaagtcaa tggaggggtct gtatccatat gggttccaaa aatatcaacc 1560
 acttcccaaa taacatttac tgtaataaac gccacaaccg tttgggggaca aaatgtatac 1620
 gttgtcggga atatttcgca gctgggggaac 1650

<210> 431
 <211> 550
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(33)

<400> 431
 Met Ser Leu Phe Lys Lys Ser Phe Pro Trp Ile Leu Ser Leu Leu Leu
 1 5 10 15
 Leu Phe Ser Phe Ile Ala Pro Phe Ser Ile Gln Thr Glu Lys Val Arg
 20 25 30
 Ala Gly Ser Val Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp
 35 40 45
 Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr Lys Val Ala Asn Asn Ala
 50 55 60
 Gln Ser Leu Ala Asn Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala
 65 70 75 80
 Tyr Lys Gly Thr Ser Ser Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu
 85 90 95
 Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr
 100 105 110
 Gly Thr Lys Thr Gln Tyr Ile Gln Ala Ile Gln Ala Ala His Thr Ala
 115 120 125
 Gly Met Arg Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala
 130 135 140
 Asp Gly Thr Glu Leu Val Asp Ala Val Glu Val Asn Pro Ser Asp Arg
 145 150 155 160
 Asn Gln Glu Ile Ser Gly Thr Tyr Gln Ile Gln Ala Trp Thr Lys Phe
 165 170 175
 Asp Phe Pro Gly Arg Gly Asn Thr Tyr Ser Ser Phe Lys Trp Arg Trp
 180 185 190
 Tyr His Phe Asp Gly Thr Asp Trp Asp Glu Ser Arg Lys Leu Asn Arg
 195 200 205
 Ile Tyr Lys Phe Arg Gly Thr Gly Lys Ala Trp Asp Trp Glu Val Asp
 210 215 220
 Thr Glu Asn Gly Asn Tyr Asp Tyr Leu Met Tyr Ala Asp Leu Asp Met
 225 230 235 240
 Asp His Pro Glu Val Val Ser Glu Leu Lys Asn Trp Gly Lys Trp Tyr
 245 250 255
 Val Thr Thr Thr Asn Ile Asp Gly Phe Arg Leu Asp Ala Val Lys His
 260 265 270
 Ile Lys Tyr Ser Phe Phe Pro Asp Trp Leu Ser Tyr Val Arg Thr Gln
 275 280 285
 Thr Gln Lys Pro Leu Phe Ala Val Gly Glu Phe Trp Ser Tyr Asp Ile
 290 295 300
 Ser Lys Leu His Asn Tyr Ile Thr Lys Thr Asn Gly Ser Met Ser Leu
 305 310 315 320
 Phe Asp Ala Pro Leu His Asn Asn Phe Tyr Ile Ala Ser Lys Ser Ser
 325 330 335
 Gly Tyr Phe Asp Met Arg Thr Leu Leu Asn Asn Thr Leu Met Lys Asp
 340 345 350
 Gln Pro Thr Leu Ala Val Thr Leu Val Asp Asn His Asp Thr Glu Pro
 355 360 365
 Gly Gln Ser Leu Gln Ser Trp Val Glu Pro Trp Phe Lys Pro Leu Ala
 370 375 380
 Tyr Ala Phe Ile Leu Thr Arg Gln Glu Gly Tyr Pro Cys Val Phe Tyr
 385 390 395 400
 Gly Asp Tyr Tyr Gly Ile Pro Lys Tyr Asn Ile Pro Ala Leu Lys Ser

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Lys	Leu	Asp	Pro	405	Leu	Leu	Ile	Ala	Arg	410	Asp	Tyr	Ala	Tyr	415	Gly	Thr
Gln	His	Asp	Tyr	420	Ile	Asp	Ser	Ala	425	Asp	Ile	Ile	Gly	Trp	430	Thr	Arg
Gly	Val	Ala	Glu	435	Lys	Ala	Asn	440	Ser	Gly	Leu	Ala	Ala	Leu	445	Ile	Thr
Gly	Pro	Gly	Gly	450	Ser	Lys	Trp	455	Met	Tyr	Val	Gly	Lys	Gln	460	His	Ala
465	Lys	Thr	Phe	Tyr	470	Asp	Leu	Thr	Gly	Asn	Arg	Ser	Asp	Thr	475	Val	Thr
Asn	Ala	Asp	Gly	485	Trp	Gly	Glu	Phe	Lys	490	Val	Asn	Gly	Gly	495	Ser	Val
Ile	Trp	Val	Pro	500	Lys	Ile	Ser	Thr	505	Thr	Ser	Gln	Ile	Thr	510	Phe	Thr
Asn	Asn	Ala	Thr	515	Thr	Val	Trp	Gly	520	Gln	Asn	Val	Tyr	Val	525	Val	Gly
530	Ile	Ser	Gln	Leu	Gly	Asn	535	540									
545																	

<210> 432
 <211> 1752
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 432

atgtccgaaa	gaggcgctcag	gagagccgtg	cgcaccgccc	tggctcgggtt	ggcggcgggcc	60
gccaccgccc	cggtgacgct	cggcgctccc	accgccagg	ccgcgcgggc	cggaacaag	120
gacgtcaccg	ccgtcctggt	ctcctggaac	ttcaactcca	tcgcccgcga	gtgcaccgag	180
cgactcggcc	ccgcgggcta	cggtctcgtg	caggctctcg	cgccccagga	gcacatccag	240
ggctccgcgt	ggtggaccca	gtaccagccc	gtgagttaca	acatctccgg	ccggctcggc	300
aacgagcagc	agttccgctc	gatggtcaac	acctgtaaca	acgcgggtgt	cggcgtcatc	360
gtggacgcgg	tcataacca	catgtccgcc	ggctcgggca	ccggtaccgg	cggcacctcc	420
tactccaagt	acaactaccc	cggcctctac	agcggcaacg	acttcggttc	ctgccgcgag	480
gacatccggg	gcggtgacta	caccaacgac	cgctggcggg	tccagaactg	cgagctgggtc	540
ggcctcgccg	acctggacac	cggcagcgcc	tacgtccagc	agcagatcgc	cggtacatg	600
aaccgtctgc	tggtctgggg	cgctgcgggt	ttccgggtgg	acgccgtcaa	gcacatcccg	660
gcccggcaca	tggagcagat	ccgcgcccgg	gtcaacggcg	gcgacgtctt	ctggaaccag	720
gaggtcatct	acggcgccgg	cgaggcgatc	acgcccagg	agtacctcaa	caccggctac	780
gtgcaggagt	tccgtacgc	gttcgacctc	aagcggatgt	tccagggcga	ccggatcgcc	840
aatctgcaga	acttcggcga	gtcatggggc	tacatgccct	ccaaccgctc	cggcgtcttc	900
gtcgacaacc	acgacaccga	gcgcaacggc	tccacctca	gctacaagga	caacgccgcc	960
tacaccctgg	ccaacgtctt	catgtggcc	tggtccctac	gcagcccga	cggtgactcc	1020
ggctacgagt	tcaccaactt	cgacgcccgt	ccgcccaca	acggcaacgt	gaccgcctgc	1080
caccagagcg	ggtggaagtg	ccagcacgcc	tggcaggaga	tctcctccat	ggtgggcttc	1140
cgcaacgcc	cccgcggcca	ggccgtcacc	aactggtggt	ccaacggcaa	caacgccatc	1200
gccttcggcc	ggggcaaccg	cggtacgtg	gcgatcaacc	acgagaacac	cacactgaac	1260
cggaccttcc	agacctccct	gccggccggg	gactactgca	acgtgcagaa	cggcaccacc	1320
gtcaccgtca	acggcgccgg	gcagttcacc	gcgagcctcg	gcgcccgcac	cgccctcgcc	1380
ctgcacgtcg	acgcccgcaa	ttgcgcccgc	ggcggcaccg	ggggcaacgg	cggcaccggg	1440
ggaggcgaga	acccggtgac	cggcggtgcc	tccttcggcg	tcgacgcgac	cacgcagatg	1500
ggccagaaaca	tccacgtcgc	gggcaacatc	cccgcgtgg	gcgactggaa	caccgccaac	1560
gccccccgca	tgagcgccga	cacctaccgc	gtctggcggc	tggaactgaa	cctgcccgcc	1620
ggcaccacct	tccagtacaa	gtacatccgc	agggacgcca	acggcaacgt	gacctgggaa	1680
tccgggaaca	accgcaccgc	caccgtgccc	tcctccggcc	gggtgacgct	caacgacacc	1740
tggcgcaact	ga					1752

<210> 433
 <211> 583
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<221> SIGNAL
<222> (1)...(34)

<400> 433

Met	Ser	Glu	Arg	Gly	Val	Arg	Arg	Ala	Val	Arg	Thr	Ala	Leu	Val	Gly
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Leu	Ala	Ala	Ala	Ala	Thr	Ala	Ala	Val	Thr	Leu	Gly	Ala	Pro	Thr	Ala
			20					25					30		
Gln	Ala	Ala	Pro	Ala	Gly	Asn	Lys	Asp	Val	Thr	Ala	Val	Leu	Phe	Ser
		35					40					45			
Trp	Asn	Phe	Asn	Ser	Ile	Ala	Arg	Glu	Cys	Thr	Glu	Arg	Leu	Gly	Pro
	50					55					60				
Ala	Gly	Tyr	Gly	Phe	Val	Gln	Val	Ser	Pro	Pro	Gln	Glu	His	Ile	Gln
65					70					75					80
Gly	Ser	Ala	Trp	Trp	Thr	Gln	Tyr	Gln	Pro	Val	Ser	Tyr	Asn	Ile	Ser
				85					90					95	
Gly	Arg	Leu	Gly	Asn	Glu	Gln	Gln	Phe	Arg	Ser	Met	Val	Asn	Thr	Cys
			100					105					110		
Asn	Asn	Ala	Gly	Val	Gly	Val	Ile	Val	Asp	Ala	Val	Ile	Asn	His	Met
		115					120					125			
Ser	Ala	Gly	Ser	Gly	Thr	Gly	Thr	Gly	Gly	Thr	Ser	Tyr	Ser	Lys	Tyr
	130					135					140				
Asn	Tyr	Pro	Gly	Leu	Tyr	Ser	Gly	Asn	Asp	Phe	Gly	Ser	Cys	Arg	Glu
145					150					155					160
Asp	Ile	Arg	Gly	Gly	Asp	Tyr	Thr	Asn	Asp	Arg	Trp	Arg	Val	Gln	Asn
				165					170					175	
Cys	Glu	Leu	Val	Gly	Leu	Ala	Asp	Leu	Asp	Thr	Gly	Ser	Ala	Tyr	Val
			180					185					190		
Gln	Gln	Gln	Ile	Ala	Gly	Tyr	Met	Asn	Arg	Leu	Leu	Gly	Trp	Gly	Val
		195					200					205			
Ala	Gly	Phe	Arg	Val	Asp	Ala	Val	Lys	His	Ile	Pro	Ala	Arg	His	Met
	210					215					220				
Glu	Gln	Ile	Arg	Ala	Arg	Val	Asn	Gly	Gly	Asp	Val	Phe	Trp	Asn	Gln
225					230					235					240
Glu	Val	Ile	Tyr	Gly	Ala	Gly	Glu	Ala	Ile	Thr	Pro	Glu	Glu	Tyr	Leu
				245					250					255	
Asn	Thr	Gly	His	Val	Gln	Glu	Phe	Arg	Tyr	Ala	Phe	Asp	Leu	Lys	Arg
			260					265					270		
Met	Phe	Gln	Gly	Asp	Arg	Ile	Ala	Asn	Leu	Gln	Asn	Phe	Gly	Glu	Ser
		275					280					285			
Trp	Gly	Tyr	Met	Pro	Ser	Asn	Arg	Ser	Gly	Val	Phe	Val	Asp	Asn	His
	290					295					300				
Asp	Thr	Glu	Arg	Asn	Gly	Ser	Thr	Leu	Ser	Tyr	Lys	Asp	Asn	Ala	Ala
305					310					315					320
Tyr	Thr	Leu	Ala	Asn	Val	Phe	Met	Leu	Ala	Trp	Pro	Tyr	Gly	Ser	Pro
				325					330					335	
Asp	Val	His	Ser	Gly	Tyr	Glu	Phe	Thr	Asn	Phe	Asp	Ala	Gly	Pro	Pro
			340					345					350		
Asn	Asn	Gly	Asn	Val	Thr	Ala	Cys	His	Gln	Ser	Gly	Trp	Lys	Cys	Gln
		355					360					365			
His	Ala	Trp	Gln	Glu	Ile	Ser	Ser	Met	Val	Gly	Phe	Arg	Asn	Ala	Thr
	370					375					380				
Arg	Gly	Gln	Ala	Val	Thr	Asn	Trp	Trp	Ser	Asn	Gly	Asn	Asn	Ala	Ile
385					390					395					400
Ala	Phe	Gly	Arg	Gly	Asn	Arg	Gly	Tyr	Val	Ala	Ile	Asn	His	Glu	Asn
				405					410					415	
Thr	Thr	Leu	Asn	Arg	Thr	Phe	Gln	Thr	Ser	Leu	Pro	Ala	Gly	Asp	Tyr
			420					425					430		
Cys	Asn	Val	Gln	Asn	Gly	Thr	Thr	Val	Thr	Val	Asn	Gly	Ala	Gly	Gln
		435					440					445			
Phe	Thr	Ala	Ser	Leu	Gly	Ala	Arg	Thr	Ala	Leu	Ala	Leu	His	Val	Asp
	450					455					460				
Ala	Arg	Asn	Cys	Ala	Gly	Gly	Gly	Thr	Gly	Gly	Asn	Gly	Gly	Thr	Gly
465					470					475					480
Gly	Gly	Glu	Asn	Pro	Val	Thr	Gly	Gly	Ala	Ser	Phe	Gly	Val	Asp	Ala
				485					490					495	
Thr	Thr	Gln	Met	Gly	Gln	Asn	Ile	His	Val	Ala	Gly	Asn	Ile	Pro	Ala
			500					505					510		

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Leu Gly Asp Trp Asn Thr Ala Asn Ala Pro Arg Met Ser Ala Asp Thr
 515 520 525
 Tyr Pro Val Trp Arg Leu Glu Asn Leu Pro Pro Gly Thr Thr Phe
 530 535 540
 Gln Tyr Lys Tyr Ile Arg Arg Asp Ala Asn Gly Asn Val Thr Trp Glu
 545 550 555 560
 Ser Gly Asn Asn Arg Thr Ala Thr Val Pro Ser Ser Gly Arg Val Thr
 565 570 575
 Leu Asn Asp Thr Trp Arg Asn
 580

<210> 434
 <211> 1302
 <212> DNA
 <213> Unknown
 <220>
 <223> obtained from an environmental sample

<400> 434
 atgagcaata tgtataaacg caactcaatg tttttattta tagttctttt tgttttgtta 60
 tcttttgtgc cagcagtggc agaccacccc caggaagtat cagaaggcgt tatgcttcag 120
 ggttttggct ggaacagtca agtttagagg tgcgccctcta aatggtacac tcttattgaa 180
 agtaaagcag acagcatcaa aaatctaggc gttgattttg tgtggttccc gcctgtagct 240
 cgttctgtaa gccctcaggg atacctccct ggtgactact atgatttagg cagtgcagat 300
 taccctacat tttatggcac caaagatcag ctaaaggcag ctttaaaagc tctaaacgac 360
 agaaacattg cacctgttgc cgacattgta gttaatcata gatgtgctgg caagcaagat 420
 gagaacggca tctggaatat attccactat ctttcgggca aggcacaatg ggaacaatgg 480
 gcggtctgtg aagggtcagtt tgggtggaaca ggcaatcctg acacaggaga taattttcat 540
 gctgcacctg acattgacca cacaacaag cagggtccaac aagacataag cgattggatg 600
 aactggctaa aagagctggg ctttagagct tggcgctacg actattctag aggttatgct 660
 gctaaatatg cgggtcttta tgacaaaaac acacaacctt tgttctctgt tggcgaactt 720
 tggacgaata tggccttcga aggctcctgg ctcaagccaa atcaagatgc tcacaggcaa 780
 ttacttgttg actggcttga cgagaatcct agtccagttg ccacaatatt tgactttacc 840
 accaaagggg ttttacaggt agccgtacac ggagaatacc aaagacttat tgactcaaac 900
 aataatgctc caggccttat tggatggtgg cctcaaagag cagtaacatt tattgacaat 960
 cacgacactg gctctcagca aagtcattgg ccatttgcag acgacaaagt tatgcagggt 1020
 tatgcataca tcttaacaca ccccgggatt cctgcctat tctgggagca cgtttacgac 1080
 tggattttgt atgatcaaat caagcaacta gttgatattc gcaagcaaaa taaaataaac 1140
 agtacaagct cgctaaaaat cttaagagcc gagacgggcc tctacgcagc aattattgat 1200
 gacaagggtg ttttaaaact tggttggcaa gattggtatc ctggcgaagg ctttgaacta 1260
 aaagccagcg gagagcagta tgctgtgtgg gttaaaaact ag 1302

<210> 435
 <211> 433
 <212> PRT
 <213> Unknown
 <220>
 <223> obtained from an environmental sample
 <221> SIGNAL
 <222> (1)...(27)

<400> 435
 Met Ser Asn Met Tyr Lys Arg Asn Ser Met Phe Leu Phe Ile Val Leu
 1 5 10 15
 Phe Val Leu Leu Ser Phe Val Pro Ala Val Ala Asp Pro Pro Gln Glu
 20 25 30
 Val Ser Glu Gly Val Met Leu Gln Gly Phe Gly Trp Asn Ser Gln Val
 35 40 45
 Arg Gly Ser Pro Ser Lys Trp Tyr Thr Leu Ile Glu Ser Lys Ala Asp
 50 55 60
 Ser Ile Lys Asn Leu Gly Val Asp Phe Val Trp Phe Pro Pro Val Ala
 65 70 75 80
 Arg Ser Val Ser Pro Gln Gly Tyr Leu Pro Gly Asp Tyr Tyr Asp Leu
 85 90 95
 Gly Ser Asp Asp Tyr Pro Thr Phe Tyr Gly Thr Lys Asp Gln Leu Lys

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Ala	Ala	Leu	100	Ala	Leu	Asn	Asp	105	Arg	Asn	Ile	Ala	Pro	110	Val	Ala	Asp
Ile	Val	Val	115	Asn	His	Arg	Cys	120	Gly	Lys	Gln	Asp	Glu	125	Asn	Gly	Ile
Trp	Asn	Ile	130	Phe	His	Tyr	Pro	135	Gly	Lys	Ala	Gln	Trp	140	Glu	Gln	Trp
Ala	Val	Cys	145	Glu	Gly	Gln	Phe	150	Gly	Gly	Thr	Gly	Asn	155	Pro	Asp	Thr
Asp	Asn	Phe	165	His	Ala	Ala	Pro	170	Ile	Asp	His	Thr	Asn	175	Lys	Gln	Val
Gln	Gln	Asp	180	Ile	Ser	Asp	Trp	185	Asn	Trp	Leu	Lys	Glu	190	Leu	Gly	Phe
Arg	Ala	Trp	195	Arg	Tyr	Asp	Tyr	200	Arg	Gly	Tyr	Ala	Ala	205	Lys	Tyr	Ala
Gly	Leu	Tyr	210	Asp	Lys	Asn	Thr	215	Gln	Pro	Leu	Phe	Ser	220	Val	Gly	Glu
Trp	Thr	Asn	225	Met	Ala	Phe	Glu	230	Gly	Ser	Trp	Leu	Lys	235	Pro	Asn	Gln
Ala	His	Arg	245	Gln	Leu	Leu	Val	250	Trp	Leu	Asp	Glu	Asn	255	Pro	Ser	Pro
Val	Ala	Thr	260	Ile	Phe	Asp	Phe	265	Thr	Thr	Lys	Gly	Ile	270	Leu	Gln	Val
Val	His	Gly	275	Glu	Tyr	Gln	Arg	280	Ile	Asp	Ser	Asn	Asn	285	Asn	Ala	Pro
Gly	Leu	Ile	290	Gly	Trp	Trp	Pro	295	Gln	Arg	Ala	Val	Thr	300	Phe	Ile	Asp
His	Asp	Thr	305	Gly	Ser	Gln	Gln	310	Ser	His	Trp	Pro	Phe	315	Ala	Asp	Asp
Val	Met	Gln	325	Gly	Tyr	Ala	Tyr	330	Leu	Thr	His	Pro	Gly	335	Ile	Pro	Cys
Leu	Phe	Trp	340	Glu	His	Val	Tyr	345	Trp	Asn	Leu	Tyr	Asp	350	Gln	Ile	Lys
Gln	Leu	Val	355	Asp	Ile	Arg	Lys	360	Gln	Asn	Lys	Ile	Asn	365	Ser	Thr	Ser
Leu	Lys	Ile	370	Leu	Arg	Ala	Glu	375	Thr	Gly	Leu	Tyr	Ala	380	Ala	Ile	Ile
Asp	Lys	Val	385	Val	Leu	Lys	Leu	390	Gly	Trp	Gln	Asp	Trp	395	Tyr	Pro	Gly
Gly	Phe	Glu	405	Leu	Lys	Ala	Ser	410	Glu	Gln	Tyr	Ala	Val	415	Trp	Val	Lys
Asn			420					425						430			

<210> 436
 <211> 1311
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Synthetically generated

<400> 436																				
atggccaagt	actccgagct	ggaaaagggc	ggggtcataa	tgcaggcggt	ctactgggac															60
gtgccttcag	gaggaatatg	gtgggacaca	atacggcaga	agataccgga	gtggtacgat															120
gccggaatct	ccgcaatatg	gattcccccg	gagagcaagg	gcatgggcgg	cgctatttcg															180
atgggctacg	acccctacga	cttctttgac	ctcgggtgagt	acgaccagaa	gggaacggta															240
gagacgcgct	ttggctccaa	gcaggagctc	gtgaacatga	taaacaccgc	ccacgcctat															300
ggcatgaagg	taatagccga	tatagtcatc	aaccaccgcg	ccggcggtga	cctggagtg															360
aaccccttcg	tgaacgacta	tacctggacc	gactttctca	aggctcgcgc	gggtaaatac															420
acgggccaact	acctcgactt	ccaccgaac	gagctccatg	cgggcgattc	cggaacattt															480
ggaggctatc	ccgacatatg	ccacgacaag	agctgggacc	agtactggct	ctgggccagc															540
caggagagct	acgcggcata	tctcaggagc	atcggcatcg	atgcctggcg	cttcgactac															600
gtcaagggct	atgctccctg	ggtcgtcaag	gactggctga	actggtgggg	aggctggg															660
gttgagagct	actgggacac	caacgtcgac	gctgttctca	actgggcata	ctcgagcggt															720
gccaaaggtct	ttgacttcgc	cctctactac	aagatggatg	aggcctttga	caacaaaaac															780
attccagcgc	tcgtctctgc	ccttcagaac	ggccagactg	ttgtctcccg	cgaccggttc															840
aaggccgtaa	cctttgtagc	aaaccacgac	accgatataa	tctggaacaa	gtatccagcc															900

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tacgcggttca	tcctcaccta	cgagggccag	ccgacaatat	tctaccgcga	ctacgaggag	960
tggtcaaca	aggataagct	caagaacctc	atctggatac	atgagaacct	cgccggagga	1020
agcaccgaca	tagtctacta	cgataacgat	gaactcatct	tcgtcaggaa	cggctacggg	1080
gacaagccgg	ggcttataac	ctacatcaac	ctaggctcga	gcaaggccgg	aaggtgggtt	1140
tatgtgccga	agttcgcggg	cgcgtgcata	cacgagtata	ctggtaacct	cggaggctgg	1200
gtagacaagt	acgtctactc	aagcggctgg	gtctatctcg	aagctccagc	ttacgacctt	1260
gccaacgggc	agtatggcta	ctccgtgtgg	agctactgca	gggtgggctg	a	1311

<210> 437
 <211> 436
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> synthetically generated

<400> 437

Met	Ala	Lys	Tyr	Ser	Glu	Leu	Glu	Lys	Gly	Gly	Val	Ile	Met	Gln	Ala
1				5					10					15	
Phe	Tyr	Trp	Asp	Val	Pro	Ser	Gly	Gly	Ile	Trp	Trp	Asp	Thr	Ile	Arg
			20					25					30		
Gln	Lys	Ile	Pro	Glu	Trp	Tyr	Asp	Ala	Gly	Ile	Ser	Ala	Ile	Trp	Ile
		35					40					45			
Pro	Pro	Ala	Ser	Lys	Gly	Met	Gly	Gly	Ala	Tyr	Ser	Met	Gly	Tyr	Asp
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Glu	Thr	Arg	Phe	Gly	Ser	Lys	Gln	Glu	Leu	Val	Asn	Met	Ile	Asn	Thr
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Ala	His	Ala	Tyr	Gly	Met	Lys	Val	Ile	Ala	Asp	Ile	Val	Ile	Asn	His
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Arg	Ala	Gly	Gly	Asp	Leu	Glu	Trp	Asn	Pro	Phe	Val	Asn	Asp	Tyr	Thr
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Trp	Thr	Asp	Phe	Ser	Lys	Val	Ala	Ser	Gly	Lys	Tyr	Thr	Ala	Asn	Tyr
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Gly	Gly	Tyr	Pro	Asp	Ile	Cys	His	Asp	Lys	Ser	Trp	Asp	Gln	Tyr	Trp
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Leu	Trp	Ala	Ser	Gln	Glu	Ser	Tyr	Ala	Ala	Tyr	Leu	Arg	Ser	Ile	Gly
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Ile	Asp	Ala	Trp	Arg	Phe	Asp	Tyr	Val	Lys	Gly	Tyr	Ala	Pro	Trp	Val
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Ala	Lys	Val	Phe	Asp	Phe	Ala	Leu	Tyr	Tyr	Lys	Met	Asp	Glu	Ala	Phe
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Thr	Val	Val	Ser	Arg	Asp	Pro	Phe	Lys	Ala	Val	Thr	Phe	Val	Ala	Asn
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His	Asp	Thr	Asp	Ile	Ile	Trp	Asn	Lys	Tyr	Pro	Ala	Tyr	Ala	Phe	Ile
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Trp	Leu	Asn	Lys	Asp	Lys	Leu	Lys	Asn	Leu	Ile	Trp	Ile	His	Glu	Asn
				325					330					335	
Leu	Ala	Gly	Gly	Ser	Thr	Asp	Ile	Val	Tyr	Tyr	Asp	Asn	Asp	Glu	Leu
			340					345					350		
Ile	Phe	Val	Arg	Asn	Gly	Tyr	Gly	Asp	Lys	Pro	Gly	Leu	Ile	Thr	Tyr
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Ile	Asn	Leu	Gly	Ser	Ser	Lys	Ala	Gly	Arg	Trp	Val	Tyr	Val	Pro	Lys
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Val	Asp	Lys	Tyr	Val	Tyr	Ser	Ser	Gly	Trp	Val	Tyr	Leu	Glu	Ala	Pro

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Ala Tyr Asp Pro⁴⁰⁵ Ala Asn Gly Gln Tyr⁴¹⁰ Gly Tyr Ser Val Trp⁴¹⁵ Ser Tyr
 Cys Gly Val Gly⁴²⁰
 Gly⁴³⁵

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 <211> 1539
 <212> DNA
 <213> Unknown

<220>
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 cgcccaata atcctgaact aaaatatgaa tatcagcaat tcacggaaaa tgagacggac 420
 tggttgaatg attttgccct ttttatggcc attaaggagg ctaacggggg ggtatcctgg 480
 gacaaactggc caaagggaact gcgtagccgc cagccggacg caatcgaaaa gttcaaacaa 540
 accgaagccg accgcattca acgccatgcc tttcgacagt ttttattctt ccgccagtgg 600
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 tttgtcgctt acgatagcgc agatgcttgg tcgaaccag agttgtttta tctgaatgaa 720
 gaaggcaaac caactgttgt ggcaggcgtt ccaccggatt atttttcccc aacgggtcaa 780
 ttatggggta acccgcttta caagtgggaa gtatcatcgtc agcagaattt ccgctggtgg 840
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 atcatcgctg aagatttggg ggtcatcacc cccgatgtga ttgaactgcg cgattcgttt 1080
 aactgcccc gtatgaaggt gttccagttt gcttttacaa cggacccgct cgatcccttc 1140
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 <212> PRT
 <213> Unknown

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 <222> (11)...(493)
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 35 40 45
 Leu Gly Pro Thr Gly Tyr Gly Asp Ser Pro Tyr Gln Cys Phe Ser Ala
 50 55 60
 phe Ala Gly Asn Pro Tyr Leu Val Ser Pro Ala Leu Leu Leu Glu Asn
 65 70 75 80
 Gly Leu Leu Arg Arg Ser Asp Leu Thr Asp Arg Pro Glu Phe Asn Pro
 85 90 95
 Ile Arg Ile Asp Tyr Gly Glu Ala Ile Thr Trp Lys Leu Lys Ile Leu

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Phe	Ala	Leu	130	Phe	Met	Ala	Ile	Lys	135	Glu	Ala	Asn	Gly	Gly	140	Val	Ser	Trp
Asp	Asn	Trp	145	Pro	Lys	Glu	Leu	Arg	150	Ser	Arg	Gln	Pro	Asp	155	Ala	Ile	Glu
Lys	Phe	Lys	160	Gln	Thr	Glu	Ala	Asp	165	Ile	Gln	Arg	His	Ala	170	Phe	Arg	
Gln	Phe	Leu	175	Phe	Phe	Arg	Gln	Trp	180	Leu	Asp	Leu	Lys	Ala	185	Tyr	Ala	Asn
Gln	Lys	Asn	190	Ile	Gln	Ile	Ile	Gly	195	Asp	Ile	Pro	Ile	Phe	200	Val	Ala	Tyr
Asp	Ser	Ala	205	Asp	Ala	Trp	Ser	Asn	210	Pro	Glu	Leu	Phe	Tyr	215	Leu	Asn	Glu
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Tyr	Trp	Glu	280	Val	Pro	Phe	Gly	Met	285	Pro	Thr	Ala	Glu	Ile	290	Gly	Arg	Trp
Val	Lys	Gly	295	Pro	Gly	Lys	Glu	Leu	300	Phe	Asn	Ala	Ile	Arg	305	Asp	Ala	Leu
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Val	Ile	Glu	325	Leu	Arg	Asp	Ser	Phe	330	Asn	Leu	Pro	Gly	Met	335	Lys	Val	Phe
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Ala	Leu	Gly	370	Trp	Tyr	Gln	Ser	Ala	375	Pro	Glu	Lys	Glu	Arg	380	Asp	Phe	Ile
Arg	Arg	Tyr	385	Leu	Ala	Arg	Ser	Gly	390	Glu	Asp	Ile	Ser	Trp	395	Asp	Met	Ile
Arg	Ala	Val	400	Trp	Ser	Ser	Val	Ala	405	Val	Phe	Ala	Leu	Ala	410	Pro	Met	Gln
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Ala	Ser	Gly	430	Asn	Trp	Cys	Trp	Arg	435	Leu	His	Pro	Glu	Ala	440	Phe	Asn	Pro
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 <212> DNA
 <213> Unknown

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 <212> PRT
 <213> Unknown

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 <222> (1)...(32)

<221> DOMAIN
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Cys	Asn	Gln	Tyr	Gln	Gly	Pro	Ser	Leu	Ala	Ser	Glu	Met	Ala	Ala	Leu
Ala	Glu	Glu	Gln	Ser	Glu	Ala	Gly	Phe	Thr	Met	Met	Trp	Leu	Pro	Pro
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Gly	Thr	Glu	Val	Glu	Thr	Trp	Met	Thr	Ser	Leu	Ser	Glu	Asn	Asp	Ile
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Glu	Val	Gly	Gly	Gly	Gly	Thr	Gly	Ile	Asp	Gln	Arg	Ile	Tyr	Gly	Ile
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Thr	Arg	Thr	Asp	Phe	Thr	Asn	Met	Pro	Ser	Gly	Lys	Gly	Ala	Met	Thr
Phe	Arg	His	Phe	Lys	Pro	Asn	Gly	Ile	Thr	Pro	Thr	Cys	Met	Thr	Gly
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Gly	Ser	Gly	Ser	Pro	Ser	Val	Thr	Phe	Asn	Asp	Gln	Thr	Lys	Ser	Ser
Gly	Gly	Val	Phe	Tyr	Gly	Lys	Glu	Lys	Asp	Glu	Gln	Arg	Ser	Ile	Val
Arg	Leu	Glu	Leu	Ser	Gly	Glu	Glu	Leu	Gln	Asn	Ser	Ala	Trp	Leu	Arg
Phe	Ser	Glu	Ser	Gly	Ser	Tyr	Glu	Ser	Gln	Val	Arg	Gly	Asp	Ala	Leu
Gln	Leu	Glu	Pro	Leu	Ser	Gln	Thr	Phe	Ala	Lys	Leu	Ala	Ile	Glu	Lys
Gly	Gly	Thr	Leu	Phe	Asp	Ile	Ser	His	Leu	Pro	Val	Met	Asp	Glu	Glu
Tyr	Ser	Val	Pro	Val	His	Ile	Ser	Ala	Thr	Asn	Ser	Gly	Val	Tyr	Thr
Leu	Thr	Ala	Thr	Asp	Phe	Glu	Leu	Pro	Gly	Asn	Leu	Glu	Leu	Thr	Phe
His	Asp	His	Ala	Ala	Gly	Thr	Ser	Val	Pro	Leu	Asn	Ala	Asp	Phe	Ser
Tyr	Thr	Ile	Glu	Gln	Ala	Gln	Ala	Ala	Lys	Ala	Ala	Glu	Leu	Ser	Pro
Leu	Glu	Arg	Val	Lys	Gln	Gly	Pro	Val	Val	Ala	Lys	Val	Thr	Ala	Asp
Ser	Pro	Gln	Tyr	Ser	Ile	Thr	Val	Asn	Ser	Ser	Ala	Asp	Ile	Gly	Gly
Thr	Asp	Glu	Leu	Pro	Glu	Arg	Val	Ala	Leu	Arg	Gln	Asn	Tyr	Pro	Asn
Pro	Phe	Asn	Pro	Val	Thr	Val	Ile	Thr	Tyr	Glu	Leu	Pro	Glu	Asn	Ser
Asn	Val	Thr	Leu	Glu	Val	Phe	Asp	Met	Ala	Gly	Arg	Lys	Val	Ser	Thr

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Leu Val Asn Glu Lys Val Ser Ala Gly Ser His Asp Val Ala Phe Asp
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 <211> 1668
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 cacggcgggcg acgagatcat cctccagggc ttctactgga ataccgtgcg gacctcgagc 180
 aactggtagc cgacgttggc cagcatggcg ccgacctgg ccgccgatgg cttcagtgcg 240
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<210> 443
 <211> 555
 <212> PRT
 <213> Unknown

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 <222> (1)...(27)

<221> DOMAIN
 <222> (45)...(395)
 <223> Alpha amylase, catalytic domain

<221> DOMAIN
 <222> (458)...(554)
 <223> Starch binding domain

<400> 443
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 Leu Ala Tyr Pro Trp Gly Asn Leu Val Arg Ala Ala Asp Ala Pro Gly
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Asn	Asp	Cys	160	Asp	Gly	Asp	Arg	165	Phe	Met	Gly	Gly	Asp	170	Ala	Asp	Leu
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Arg	Gly	Tyr	205	Ala	Gly	Glu	Arg	210	Val	Ala	Ser	Trp	Met	215	Ser	Asp	Ala
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Pro	Ser	Trp	235	Asp	Trp	Arg	Asn	240	Ala	Ser	Trp	Gln	Gln	245	Ile	Leu	Lys
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Gly	Asn	Pro	280	Asp	Ala	Arg	Trp	285	Ala	Glu	Val	Ala	Val	290	Thr	Phe	Val
Asn	His	Asp	295	Thr	Gly	Tyr	Ser	300	Pro	Gly	Pro	His	Gly	305	Gly	Gln	His
Trp	Pro	Leu	310	Pro	Asp	Ala	Arg	315	Lys	Gln	Ala	Tyr	Ala	320	Tyr	Ile	Leu
Ser	Ser	Pro	325	Gly	Thr	Pro	Val	330	Tyr	Trp	Pro	His	Met	335	Tyr	Asp	Trp
Gly	His	Gly	340	Asp	Phe	Ile	Arg	345	Gln	Leu	Ile	Gln	Ile	350	Arg	Ala	Ala
Gly	Val	Lys	355	Ala	Ala	Ser	Ala	360	Ile	Gln	Phe	His	Ser	365	Gly	Phe	Ser
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Gln	Ala	Leu	400	Asn	Thr	Asp	Asn	405	Ala	Ile	Arg	Ile	Trp	410	Arg	Ser	Gly
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Cys	Asp	Asn	430	Gly	Val	Thr	Gln	435	Trp	Gly	Asp	Ser	Val	440	Tyr	Ala	Leu
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Thr	Asp	Thr	460	Ser	Ala	Tyr	Pro	465	Trp	Lys	Gly	Ser	Ile	470	Ala	Leu	Pro
Ala	Gly	Gln	475	Gln	Val	Gln	Trp	480	Cys	Ile	Val	Arg	Ser	485	Glu	Ser	Asn
Pro	Thr	Gln	490	Val	Lys	Thr	Trp	495	Gln	Pro	Gly	Gly	Asn	500	Ser	Val	Thr
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<212> DNA
<213> Unknown

<220>
<223> Obtained from an environmental sample

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<210> 445
<211> 506
<212> PRT
<213> Unknown

<220>
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<222> (1)...(24)

<221> DOMAIN
<222> (45)...(409)
<223> Alpha amylase, catalytic domain

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Ser Ala Arg Arg Thr Trp His Asp Glu Ile Ile Tyr Phe Ile Phe Thr
35 40 45
Asp Arg Phe Glu Asn Gly Asp Arg Ser Asn Asp Tyr Arg Val Lys Pro
50 55 60
Asn Asp Pro Trp Ala Tyr His Gly Gly Asp Leu Gln Gly Ile Ile Asn
65 70 75 80
Arg Leu Asp Tyr Ile Gln Asn Leu Gly Ala Thr Ala Leu Trp Ile Thr
85 90 95
Pro Val Ile Asp Asn Arg Asp Gly Pro Phe Val Ala Asp Phe Gly Asn
100 105 110
Gly Arg Lys Gln Glu Ile Trp Gly Tyr His Gly Tyr Trp Phe Lys Asp
115 120 125

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Phe Tyr Lys Val Glu Glu His Leu Gly Asp Met Ala Lys Leu Arg Glu
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 Leu Val Gln Lys Ala His Gln Arg Gly Ile Lys Val Leu Leu Asp Ile
 145 150 155
 Val Val Asn His Thr Asp Tyr Asp His Pro Phe Ala Leu Gln Ala Gln
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 Glu Pro Gly Asn Lys Tyr His Ser Trp Phe Asn Gln His Gly Asp Ile
 180 185 190
 Arg Asp Trp Asn Asp Pro Trp Trp Val Glu Asn Gly Arg Ala Gly
 195 200 205
 Leu Pro Asp Leu Asn Gln Gly Asn Pro Glu Thr Ala Arg Tyr Leu Ile
 210 215 220
 Asp Ala Met Lys Phe Trp Ile Lys Glu Thr Gly Val Asp Gly Phe Arg
 225 230 235 240
 Ile Asp Thr Val Lys His Val Pro Arg Thr Phe Trp Gln Gln Phe Asn
 245 250 255
 Arg Glu Ile Arg Gln Phe Ala Gly Asp Asp Phe Leu Leu Leu Gly Glu
 260 265 270
 Ile Phe Thr Gly His Pro Glu Val Gln Val Pro Tyr Leu Arg Glu Gly
 275 280 285
 Met His Thr Ala Phe Asp Phe Pro Leu Tyr Tyr Ala Ile Lys Asp Ala
 290 295 300
 Trp Gly Gln Ser Arg Ser Met Arg Ser Leu Gly Ala Ile Phe Ala Lys
 305 310 315 320
 Asp His Leu Tyr Pro Asp Ala Asn Leu Leu Ser Pro Phe Ile Asp Asn
 325 330 335
 His Asp Val Pro Arg Phe Val His Glu Ala Asn Gly Arg Gln Arg Asp
 340 345 350
 Arg Leu Met Ala Ala Leu Gly Phe Ile Phe Ala Ile Arg Gly Met Pro
 355 360 365
 Ser Leu Tyr Tyr Gly Thr Glu Val Ala Leu Pro Gly Gly Gly Asp Pro
 370 375 380
 Asp Asn Arg Arg Asp Met Gln Phe Asn Arg Asp Pro Glu Leu Gln Ala
 385 390 395 400
 Tyr Val Lys Arg Leu Ala Glu Met Arg Lys Gln Gln Pro Ala Leu Arg
 405 410 415
 Arg Gly Arg Gln Leu Glu Met Trp Gln Asp Asp Gln Val Tyr Ala Phe
 420 425 430
 Ser Arg Leu Thr Pro Gln Ala Asn Glu Glu Val Leu Ala Phe Phe Asn
 435 440 445
 Asn Ser Phe Gln Pro Gln Gln Arg Arg Ile Ala Leu Arg Asp Glu Ser
 450 455 460
 Pro Asn Lys Gly Arg Asn Pro Arg Leu Val Asn Leu Leu Asn Ala Gln
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<210> 446

<211> 3537

<212> DNA

<213> Unknown

<220>

<223> obtained from an environmental sample

<400> 446

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 <211> 1178
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> DOMAIN
 <222> (32)...(426)
 <223> Alpha amylase, catalytic domain

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 20 25 30
 Ile Met Thr Asp Arg Phe Asn Asp Gly Asp Pro Ser Asn Asn Thr Ala

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Ser	65	Ala	Ile	Trp	Ile	Ser	70	Pro	Val	Gln	Phe	Ser	75	Val	Asn	Ala	Tyr
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Tyr	Val	Ile	Ile	Asp	Ile	Val	115	Ile	Asn	His	Met	Ala	120	Asp	Leu	Ile	Gly
Ser	130	Ser	Asp	Pro	Gly	Phe	135	Pro	Ala	Phe	Asn	Trp	140	Asn	Gly	Tyr	Asn
Arg	145	Trp	Trp	Asp	Pro	Asn	150	Arg	Arg	His	Ala	Pro	155	Pro	Phe	Asp	Asp
Ser	Arg	Phe	His	Asn	Phe	Gly	165	Glu	Ile	Asp	Asn	Tyr	170	Asp	Asp	Pro	Trp
Gln	Val	Leu	Tyr	Gly	Glu	Phe	180	Lys	Pro	Gly	Gly	Leu	185	Asp	Asp	Ile	Arg
Thr	Glu	Asp	Pro	Gln	Val	Arg	195	Gln	Asp	Leu	Ile	Arg	200	Ile	Phe	Lys	Ala
Leu	Ile	Asp	Ala	Thr	Asp	Ala	210	Asp	Gly	Phe	Arg	Val	215	Asp	Ala	Val	Lys
His	Val	Glu	Glu	Gly	Phe	Phe	225	Ala	Glu	Phe	Met	Pro	230	Ala	Ile	Tyr	Ala
His	Ala	Ala	Tyr	Arg	Gly	Lys	245	Glu	Asn	Phe	Leu	Ile	250	Phe	Gly	Glu	Ser
Ile	Asp	Gly	Asn	Asp	Val	Thr	260	Leu	Ser	Arg	Trp	Thr	265	His	Pro	Asn	Tyr
His	Phe	Asn	Ser	Met	Leu	Asn	275	Phe	Pro	Met	Tyr	Tyr	280	Lys	Met	Met	Asp
Val	Phe	Val	His	Arg	Gln	Arg	295	Thr	Ser	Leu	Leu	Thr	300	Glu	Arg	Ile	Asn
Asp	Leu	Trp	Arg	Tyr	Ser	Glu	310	Gly	Ser	Arg	Leu	Gln	315	Leu	Val	Asn	Phe
Leu	Asp	Asn	His	Asp	Thr	Trp	325	Arg	Ile	Met	Tyr	Asp	330	Gln	Asn	Leu	Asp
Gly	Asp	Ile	Trp	Arg	Ile	Arg	340	Pro	Ala	Leu	Thr	Phe	345	Leu	Tyr	Thr	Ser
Leu	Gln	Ile	Pro	Cys	Leu	Tyr	355	Tyr	Gly	Thr	Glu	Gln	360	Ala	Phe	Asn	Gly
Gly	370	Thr	Asp	Pro	Trp	Asn	375	Arg	Glu	Asn	Met	Phe	380	Asp	Gly	Gly	Phe
Trp	385	Gly	Pro	Ser	Ala	Gly	390	Asp	Arg	Phe	Asp	Thr	395	Thr	His	Pro	Leu
Gln	Phe	Ile	Arg	Lys	Leu	Asn	405	Gln	Leu	Arg	Arg	Asp	410	Tyr	Pro	Ala	Leu
Thr	Leu	Gly	Asn	Phe	Val	Gln	420	Arg	Trp	Gln	Thr	Ser	425	Ser	Gly	Pro	Gly
Ile	Tyr	Ala	Tyr	Ser	Lys	Ile	435	Leu	Gly	Asp	Gln	Glu	440	Leu	Leu	Val	Val
Leu	450	Asn	Thr	Ser	Ala	Ser	455	Gln	Asn	Cys	Thr	Pro	460	Ala	Val	Ser	Arg
Pro	465	Asn	Gly	Thr	Val	Phe	470	Val	Asn	Leu	Leu	Asn	475	Pro	Ser	Glu	Gln
Thr	Val	Ser	Gly	Gly	Thr	Leu	485	Ala	Val	Ser	Val	Leu	490	Gly	His	Asp	Gln
Lys	Ile	Phe	Ala	Glu	Pro	Arg	500	Pro	Pro	Leu	Trp	Leu	505	Gly	Asn	Val	Ser
Thr	Trp	Pro	Gly	Pro	Thr	His	515	Asn	Gln	Cys	Pro	Gly	520	Glu	Leu	Cys	Asn
Pro	530	Thr	Leu	Glu	Pro	Asp	535	Glu	Glu	Leu	Trp	Val	540	Asp	Ala	Glu	Thr
Pro	545	Ile	Arg	Pro	Gly	Gln	550	Thr	Val	Thr	Val	Tyr	555	Tyr	Arg	Thr	Ala
Ala	Ala	Thr	Trp	Thr	Ala	Arg	565	Glu	Met	Leu	Trp	Arg	570	Glu	Asn	Thr	Ala

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Gln	Gly	Ser	580	Leu	Trp	His	Val	Asn	585	Leu	Gly	Gly	Phe	Pro	590	Ala	Gly	Val
Thr	Ile	Glu	595	Tyr	Tyr	Val	Arg	600	Ala	Thr	Asp	Gly	Asp	Glu	605	Val	Gln	Tyr
Ala	Asn	Asn	610	Gly	Gly	Ser	615	Asn	Ser	Phe	Ile	Ser	Val	Val	Tyr	Glu	Ala	
625	Val	Pro	630	Ser	Leu	Ser	635	Val	Ala	Pro	Trp	Pro	Ser	Ile	Gly	Cys	640	
Pro	Val	Thr	645	Leu	Leu	Tyr	Lys	Pro	Asn	Asp	Gly	Pro	Leu	Ala	Gly	Ala		
Ser	Ala	Ile	660	Tyr	Ala	His	Leu	Gly	665	Arg	Asn	Gly	Trp	Asn	Asp	Ile	Val	
Thr	Pro	Ala	675	Pro	Ala	Leu	Glu	680	Tyr	Val	Ala	Gly	Leu	Asp	Ala	Trp	Arg	
Leu	Val	Val	690	Thr	Pro	Ser	Pro	695	Gly	Thr	Leu	Gln	Leu	Asp	Val	Val	Phe	
705	His	Asp	710	Gly	Val	Trp	Asp	715	Asn	Asn	Asp	Phe	Glu	Asn	Trp	Phe		
Val	Pro	Val	725	Gln	Asp	Cys	Asp	730	Glu	Pro	Gly	Val	Phe	Leu	Val	Thr		
Tyr	Pro	Ala	740	Glu	Ser	Pro	Tyr	745	Leu	Ala	Glu	Ser	Ser	Val	Ala	Ser	Ile	
Asp	Phe	Glu	755	Gly	Val	Ala	Asp	760	Gly	Leu	Ala	Gly	Gln	Leu	Val	Trp	Ser	
Asn	Ala	Leu	770	Thr	Gly	Ala	Thr	775	Gly	Val	Gln	Pro	Leu	Ala	Pro	Val	Trp	
785	Arg	Ile	790	Asp	Ala	Ile	Gly	795	Leu	Ala	Glu	Gly	His	Asn	Pro	Ile	Thr	
Trp	Ala	Leu	805	His	Asp	Ala	Gly	810	Ser	Val	Thr	Gln	Ala	Val	Asp	Arg		
Ala	Leu	His	820	Tyr	Ala	Ala	Gly	825	Trp	Ser	Asp	Gly	Gly	Asp	Leu	Gly		
Gly	Gly	Trp	835	Gly	Gly	Gly	Trp	840	Phe	Leu	Gln	Ser	Phe	Gly	Thr	Ala	Gly	
His	Phe	Leu	850	Ala	Asp	Ala	Gly	855	Val	Asn	Gly	Asn	Leu	Arg	Leu	Gly	Asp	
865	His	Ala	870	Trp	Gly	Leu	Trp	875	Ser	Gln	Thr	Asp	Ala	Leu	Ala	Glu	Ala	
Arg	Pro	Phe	885	Ala	His	Pro	Leu	890	Val	Gly	Gln	Arg	Phe	Glu	Val	Arg		
Leu	Gln	Asn	900	Asn	Trp	Leu	Leu	905	Gly	Gln	Gln	Gly	Val	Gly	Val	Ala		
Leu	Arg	Asp	915	Gly	Asp	Gly	Ala	920	Ser	Arg	Val	Gln	Phe	Tyr	Phe	Asn	Gly	
Gly	Asp	Glu	930	Gln	Tyr	Ser	Val	935	Ala	Asp	Arg	Asp	Gly	Ala	Arg	Pro	Ser	
945	Gly	Ile	950	His	Trp	Thr	Asp	955	Glu	Pro	Leu	Pro	Leu	Ser	Ile	Glu	Val	
Ser	Ala	Gly	965	Val	Tyr	Arg	Leu	970	Thr	Val	Asn	Gly	Val	Glu	Val	Asp	Gly	
His	Phe	Glu	980	Gly	Asp	Leu	Ser	985	Glu	Val	Arg	Val	Trp	Ser	Tyr	Asn	Gly	
Gly	Leu	Thr	995	Pro	Asp	Tyr	Asp	1000	Phe	Phe	Phe	Asp	Asp	Ile	Arg	Ile	Val	
Ala	Pro	Thr	1010	Phe	Thr	Ala	Thr	1015	Asp	Arg	Val	Asp	Arg	Val	Val	Ile	Arg	
1025	Ser	Gln	1030	Ser	Gly	Thr	Gly	1035	Asp	Ser	Asn	Gly	Asp	Gly	Ile	Pro	Asp	
Trp	Tyr	Leu	1045	Arg	Tyr	Gly	Phe	1050	Asn	Pro	Phe	Gly	Pro	Ser	Ile	Ala	His	
Leu	Asp	Ser	1060	Asp	Gly	Asp	Gly	1065	Ser	Ser	Asn	Gly	Glu	Glu	Phe	Val	Ala	
Asp	Thr	Asn	1075	Pro	Asn	Asp	Pro	1080	Ser	Ser	Tyr	Phe	Pro	Lys	Thr	Ile	Glu	
Ala	Asn	Met	1090	Gln	Gly	Ala	Thr	1095	Ile	Thr	Phe	Thr	Leu	Gly	Pro	Pro	Thr	
1105	Ser	Pro	1110	Arg	Glu	Tyr	Glu	1115	Ile	Leu	His	Cys	Pro	Asp	Met	Val	Thr	

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1125 1130 1135
 Gly Ile Trp Thr Pro Met Asn Leu Arg Arg Ser Gly Arg Ala Asp Arg
 1140 1145 1150
 Gly Ala Met Gln Ile Thr Val Thr Asn Gln Gly Pro Gln Gly Ala Tyr
 1155 1160 1165
 Arg Thr Arg Val Ser Leu Pro Thr Glu Pro
 1170 1175

<210> 448
 <211> 1575
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 448
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 ccaatggcga cgagctatga gatttttggt cgctcatttg ctgatagcaa tgggtgacgg 180
 attggagata taaaagggaat gacagaaaag ttaccttatt tgcaagacct gggaattaaa 240
 ggagtatggg taatgccgat tcatccctca cttcttacc ataaatatga tgtgacggac 300
 tataaaggcg tgcataaaga ctatggtacc cttagaggatt tcaaaaagtt tgtaaagcga 360
 gcgcatgagc atgatattaa agtagtgata gatttttgta tcaatcatac agcgagggag 420
 cacccttggt ttcaagagtc gatgaagggg cctgataatc cttatcgcga ctactatgtg 480
 tgggcgaaaa aggagatat agaagatgaa atcaataaaa aggaacggc attagactct 540
 gacaatatcg agcagtgcca tgaggcaccc ggcaatgagg agctatacta cggctacttt 600
 tggggtggtg tgccagatgt gaattttgac aatcctgaag tgaagcgtga aatatttgat 660
 gcgggaagggt tttggttgga agaagttggg gtggatggct ttcgattgga tgctgcaaga 720
 catatttttc cagacgatcg ggcagctgac aatcatgctt tttgggtaga atttcgcaat 780
 gagatgaaaa gagtaaaccg tgatgtttat ctagtgtgtg aagtatgggc tgatatggaa 840
 acagtagcac cttatttgag tggtttgccc gcacttttca attttgatat gagctataaa 900
 atcgtggagg cgggtgcagcg tcaggccaat gaagaaatag cggagtatca tgcaaagggtg 960
 atgaactatt atcaagggat caaccctaatt tatattgatg ccaccttcac taccaatcac 1020
 gatcagccac ggagcatgag tcagtttgag ggggatgtga ataaagggaa gctagctgct 1080
 tcaactatat ttacactgcc tggcgcacct tatttgactt atggtgaaga aattgggatg 1140
 ttggggccaa agcctgatcc tgaaatcaga gagccgtttt tatggaaaga aggtgagaag 1200
 gatgtttacc gcacgagttg gatagaggcg aaaaactcta atgacgaaac agtaattcca 1260
 gccaaagtac gaagaagaaga tccggaaagc ttattgaacc actacaagag atgggtggct 1320
 ctacagacata aaagtgaggc gcttagcgaa ggccgaatcg tgcttgtagc cacagataat 1380
 caacaaatgg ctgtttggga gcgagtaagt aaaaatgaaa gggtagctcg gattcataat 1440
 ctttcagata gtccctcaaac tataatcagag gaggtaagcc gtaaatatga tttggcta 1500
 aaaaagatga tatggggaag tgtacaaaac aatgagctaa gtcctagagg gtccttagtc 1560
 tttgtgatta actaa 1575

<210> 449
 <211> 524
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> DOMAIN
 <222> (47)...(442)
 <223> Alpha amylase, catalytic domain

<221> SIGNAL
 <222> (1)...(23)

<400> 449
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 Val Phe Thr Gly Cys Glu Ser Ser Lys Phe Lys Glu Glu Ser Asn Tyr
 20 25 30
 Thr Ser Ser Tyr Ser Ala Glu Trp Pro Met Ala Thr Ser Tyr Glu Ile
 35 40 45
 Phe Val Arg Ser Phe Ala Asp Ser Asn Gly Asp Gly Ile Gly Asp Ile

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Lys 65	50 Gly	Met	Thr	Glu	Lys 70	55 Leu	Pro	Tyr	Leu	Gln 75	60 Asp	Leu	Gly	Ile	Lys 80
Gly	Val	Trp	Leu	Met 85	Pro	Ile	His	Pro	Ser 90	Pro	Ser	Tyr	His	Lys 95	Tyr
Asp	Val	Thr	Asp 100	Tyr	Lys	Gly	Val	His 105	Glu	Asp	Tyr	Gly	Thr 110	Leu	Glu
Asp	Phe	Lys 115	Lys	Phe	Val	Lys	Arg 120	Ala	His	Glu	His	Asp 125	Ile	Lys	Val
Val	Ile	Asp	Phe	Val	Ile	Asn 135	His	Thr	Ala	Arg	Glu 140	His	Pro	Trp	Phe
Gln 145	Glu	Ser	Met	Lys	Gly 150	Pro	Asp	Asn	Pro	Tyr 155	Arg	Asp	Tyr	Tyr	Val 160
Trp	Ala	Lys	Lys	Glu 165	Asp	Ile	Glu	Asp	Glu 170	Ile	Asn	Lys	Lys	Glu 175	Thr
Ala	Leu	Asp	Ser 180	Asp	Asn	Ile	Glu	Gln 185	Trp	His	Glu	Ala	Pro 190	Gly	Asn
Glu	Glu	Leu 195	Tyr	Tyr	Gly	Tyr	Phe 200	Trp	Gly	Gly	Met	Pro 205	Asp	Leu	Asn
Phe	Asp 210	Asn	Pro	Glu	Val	Lys 215	Arg	Glu	Ile	Phe	Asp 220	Ala	Gly	Arg	Phe
Trp 225	Leu	Glu	Glu	Val	Gly 230	Val	Asp	Gly	Phe	Arg 235	Leu	Asp	Ala	Ala	Arg 240
His	Ile	Phe	Pro	Asp 245	Asp	Arg	Ala	Ala	Asp 250	Asn	His	Ala	Phe	Trp 255	Val
Glu	Phe	Arg	Asn 260	Glu	Met	Lys	Arg	Val 265	Asn	Pro	Asp	Val	Tyr 270	Leu	Val
Gly	Glu	Val 275	Trp	Ala	Asp	Met	Glu 280	Thr	Val	Ala	Pro	Tyr 285	Leu	Ser	Gly
Leu	Pro 290	Ala	Leu	Phe	Asn	Phe 295	Asp	Met	Ser	Tyr	Lys 300	Ile	Val	Glu	Ala
Val 305	Gln	Arg	Gln	Ala	Asn 310	Glu	Ile	Ala	Glu 315	Tyr	His	Ala	Lys	Val 320	
Met	Asn	Tyr	Tyr	Gln 325	Gly	Ile	Asn	Pro	Asn 330	Tyr	Ile	Asp	Ala	Thr 335	Phe
Ile	Thr	Asn	His 340	Asp	Gln	Pro	Arg	Ser 345	Met	Ser	Gln	Phe	Glu 350	Gly	Asp
Val	Asn	Lys 355	Gly	Lys	Leu	Ala	Ala 360	Ser	Leu	Leu	Phe	Thr 365	Leu	Pro	Gly
Ala	Pro 370	Tyr	Leu	Tyr	Tyr	Gly 375	Glu	Glu	Ile	Gly	Met 380	Leu	Gly	Pro	Lys
Pro 385	Asp	Pro	Glu	Ile	Arg 390	Glu	Pro	Phe	Leu	Trp 395	Lys	Glu	Gly	Glu	Lys 400
Asp	Val	Tyr	Arg	Thr 405	Ser	Trp	Ile	Glu	Ala 410	Lys	Asn	Ser	Asn	Asp 415	Glu
Thr	Val	Ile	Pro 420	Ala	Lys	Val	Gln	Lys 425	Glu	Asp	Pro	Glu	Ser 430	Leu	Leu
Asn	His	Tyr 435	Lys	Arg	Trp	Val	Ala	Leu	Arg	His	Lys	Ser 445	Glu	Ala	Leu
Ser	Glu 450	Gly	Arg	Ile	Val	Pro 455	Val	Ala	Thr	Asp	Asn 460	Gln	Gln	Met	Ala
Val 465	Trp	Glu	Arg	Val	Ser 470	Lys	Asn	Glu	Arg	Val 475	Leu	Val	Ile	His	Asn 480
Leu	Ser	Asp	Ser	Pro 485	Gln	Thr	Ile	Ser	Glu 490	Glu	Val	Ser	Arg	Lys 495	Tyr
Asp	Leu	Ala	Asn 500	Lys	Lys	Met	Ile	Trp 505	Gly	Ser	Val	Gln	Asn 510	Asn	Glu
Leu	Ser	Pro 515	Arg	Gly	Ser	Leu	Val 520	Phe	Val	Ile	Asn				

<210> 450
 <211> 1890
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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ctattgagtt gtggtgaggc tactgagtcc tcatccctcg ccaatcaaaa aggagaaaaa 180
gagcgctcag atgaaggctc tgatgataag aaagtaattt atcaagtgtt tacacgaaca 240
tttggttaatc agaattctac taacaagatg cacggtactg ttgaggaaaa tggcacgggt 300
aagttcaatg acttcactct agacgcactc gaaggcatca aggagcttgg cgtaactcac 360
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gatatgggtca agcgtacgca tgagcgtggg atgaagggtga tcatagactt tgtgcctaata 600
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gaggatgata ctagcaaagc atttgatgta aataatcact tttactactt accgggtcag 720
gattttcaag tgccggcagg ctatcgtccg ctggggcttt ctgccaagga gatggaagac 780
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ccatcagtga atgactgggt tgagacgatc aagttgaact atggtgtgga ttacaaaaat 900
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ctattaattg cagagattta caatcccgat gcttataagc cttacatcaa acaaggaggc 1140
ttcgattatt tatatgacaa agtacaaacc tacgacagcc tgcgcaatgt gatgcaagat 1200
agaggtaacta cgcgggggtat ttccggcatca ttggaagcct tggtagatgt ggagccacat 1260
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gatgcttttg ccggtctttc tgcactggcc attagcaact tggctagtgg tagtcccttc 1380
atgctttatt ttggtcagga agttggggaa cccgccactg aagctgctgg ctttagtgag 1440
gcggacggta gaagtacaat ttttgactac ttcggtatac ctgagcatca gaagtggatg 1500
aatgaagggtg cctttgatgg tggacagctt tcagaagatc agcagcggct tagagcattg 1560
taccaagaga ttatccgagt atctacactc agtcgattgg caaactcaga gccttttact 1620
tgcagattag accaagtgga gtcctcatcc cccgaaaaaa tacattgctt ttctagagaa 1680
aatgaggaag aagagtggta tgtggtcacc aacttttctg atcaatcgac tgaggtgagt 1740
gtgaaagacc taagtgcaga caaggctcag atgatgtttt actggagaaa agagcttagc 1800
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gccatcttgg tgttttcttt ggctaagtaa

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<210> 451
<211> 629
<212> PRT
<213> Unknown

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<220>
<223> Obtained from an environmental sample

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<221> DOMAIN
<222> (75)...(496)
<223> Alpha amylase, catalytic domain

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<221> SIGNAL
<222> (1)...(49)

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Gly Ser Leu Ala Leu Leu Met Leu Leu Ser Cys Gly Glu Ala Thr
35 40 45
Glu Ser Ser Ser Leu Ala Asn Gln Lys Gly Glu Lys Glu Arg Ser Asp
50 55 60
Glu Gly Ser Asp Asp Lys Lys Val Ile Tyr Gln Val Phe Thr Arg Thr
65 70 75 80
Phe Gly Asn Gln Asn Ser Thr Asn Lys Met His Gly Thr Val Glu Glu
85 90 95
Asn Gly Thr Gly Lys Phe Asn Asp Phe Thr Leu Asp Ala Leu Glu Gly
100 105 110
Ile Lys Glu Leu Gly Val Thr His Ile Trp Tyr Thr Gly Val Ile Ala
115 120 125
His Ala Thr Ala Thr Asp Phe Ser Glu Tyr Gly Val Ala Val Asp Asn
130 135 140

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Tyr	Tyr	Asp	Val	Asn 165	Pro	Asp	Leu	Ala	Glu 170	Asp	Val	Asp	Lys	Arg 175	Met
Asp	Glu	Phe	Glu 180	Asp	Met	Val	Lys	Arg 185	Thr	His	Glu	Arg	Gly 190	Met	Lys
Val	Ile	Ile 195	Asp	Phe	Val	Pro	Asn 200	His	Val	Ala	Arg	Ser 205	Tyr	Gln	Gly
Gln	Gln 210	Gln	Pro	Glu	Gly	Ile 215	Thr	Ala	Leu	Gly	Ala 220	Glu	Asp	Asp	Thr
Ser 225	Lys	Ala	Phe	Asp	Val 230	Asn	Asn	His	Phe	Tyr 235	Tyr	Leu	Pro	Gly	Gln 240
Asp	Phe	Gln	Val	Pro 245	Ala	Gly	Tyr	Arg	Pro 250	Leu	Gly	Leu	Ser	Ala 255	Lys
Glu	Met	Glu	Asp 260	Leu	Pro	Ala	Tyr	Glu 265	Glu	Ser	Pro	Ala	Lys 270	Ala	Thr
Gly	Asn	Asp 275	Val	Phe	Ser	Ala	Ser 280	Pro	Ser	Val	Asn	Asp 285	Trp	Phe	Glu
Thr	Ile 290	Lys	Leu	Asn	Tyr	Gly 295	Val	Asp	Tyr	Gln	Asn 300	Gly	Gly	Lys	Gln
His 305	Phe	Ser	Ala	Thr	Gln 310	Ile	Pro	Asp	Thr	Trp 315	Glu	Arg	Met	Lys	Glu 320
Ile	Leu	Leu	Phe	Trp 325	Ala	Gly	Lys	Gly	Val 330	Asp	Gly	Phe	Arg	Cys 335	Asp
Met	Ala	Glu	Met 340	Val	Pro	Val	Glu	Phe 345	Trp	Glu	Trp	Val	Ser 350	Ala	Ala
Leu	Lys	Glu 355	Glu	Tyr	Pro	Asp	Ile 360	Leu	Leu	Ile	Ala	Glu 365	Ile	Tyr	Asn
Pro	Asp 370	Ala	Tyr	Lys	Pro	Tyr 375	Ile	Lys	Gln	Gly	Gly 380	Phe	Asp	Tyr	Leu
Tyr 385	Asp	Lys	Val	Gln	Thr 390	Tyr	Asp	Ser	Leu	Arg 395	Asn	Val	Met	Gln	Asp 400
Arg	Gly	Thr	Thr	Arg 405	Gly	Ile	Ser	Ala	Ser 410	Leu	Glu	Ala	Leu	Val 415	Asp
Val	Glu	Pro	His 420	Met	Leu	Arg	Phe	Leu 425	Glu	Asn	His	Asp	Glu 430	Gln	Arg
Ile	Ala	Ser 435	Phe	Phe	Ala	Gly 440	Asp	Ala	Phe	Ala	Gly 445	Leu	Pro	Ala	
Leu	Ala	Ile	Ser	Asn	Leu	Ala 455	Ser	Gly	Ser	Pro	Phe 460	Met	Leu	Tyr	Phe
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Ala	Asp	Gly	Arg	Ser 485	Thr	Ile	Phe	Asp	Tyr 490	Phe	Gly	Ile	Pro	Glu 495	His
Gln	Lys	Trp	Met 500	Asn	Glu	Gly	Ala	Phe 505	Asp	Gly	Gly	Gln	Leu	Ser	Glu
Asp	Gln 515	Gln	Arg	Leu	Arg	Ala	Leu 520	Tyr	Gln	Glu	Ile	Ile 525	Arg	Val	Ser
Thr 530	Leu	Ser	Arg	Leu	Ala	Asn 535	Ser	Glu	Pro	Phe	Thr 540	Cys	Arg	Leu	Asp
Gln 545	Val	Glu	Ser	Ser	Ser 550	Pro	Glu	Lys	Ile	His 555	Cys	Phe	Ser	Arg	Glu 560
Asn	Glu	Glu	Glu	Glu 565	Trp	Tyr	Val	Val	Thr 570	Asn	Phe	Ser	Asp	Gln 575	Ser
Thr	Glu	Val	Ser 580	Val	Lys	Asp	Leu	Ser 585	Ala	Asp	Lys	Ala	Gln 590	Met	Met
Phe	Tyr	Trp 595	Arg	Lys	Glu	Leu	Ser 600	Glu	Leu	Pro	Glu	Glu 605	Ile	Glu	Ile
Glu	Asp 610	Gly	Lys	Leu	Lys	Trp 615	Glu	Leu	Asp	Pro	Tyr 620	Ala	Ile	Leu	Val
phe 625	Ser	Leu	Ala	Lys											

<210> 452
 <211> 1734
 <212> DNA
 <213> Unknown

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<220>

<223> Obtained from an environmental sample

<400> 452

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ctggctgcgg tcagcgcttg ccagccggca tccgagtcgg cttatcaagc cccaactgca 120
gcgcctgtcg agcagaccga agcgccctgg tgggacgatg cggttttcta ccaaactctgg 180
ccgcgtagtt ttacgatac cagcggtgat ggccatggcg attttaatgg catgaccgcc 240
aaactgcctt atttgctga gctaggcatt gatgccatct ggctaacgcc gatgtttgag 300
gcgccgtctt atcatggcta tgactttacc gagttttatc aggtggaatc ggactatggc 360
accatggccg acttcgagcg ctttttgcaa caagctgaag agcgcggtat ccgggtgatt 420
ctggatctgg tgatcaacca catttcccgt gagcaccagt ggtttaagca gtcagcccag 480
cgattgagc cgtacagcga ttactttatc tggcgggaaa gcctgccaga agccggcaag 540
ggctggggcc atgcctggtc ggataacgat atgcccgaag cggctctggc ttttgatgag 600
acccggggcg agtactacta cgccgccttt ggtgccacct agccggatct gaatctggag 660
caccgggatg tgatcgaaga aatgaaacgg atggctaaat tctggctgga taaaggggtc 720
gatggcttcc ggctcgatgc ggtgcgcttt gtgatagagc gcggcccgaa aggccaggcc 780
gacttgcccg agaccattga ttactggaaa gacttcaacg cttttgtcaa aagcgttaac 840
cccgaagcct acttagtggg cgaagcctgg gttgatattc cggctggcagc caaatactgg 900
ggtgaaggca aagggtgga tcagggcttt gattttgagt ttggttacat cgtgctggat 960
atgctgcagc aagggtggc cacggaagcc cagtttgga ccatgagcca gcagtcggg 1020
caggctgaag atggcgcttt gcatccactg gcgcagaatg tgcagcgccg cctggagtcg 1080
ggcgaccgcg tggccttctt cacgcccttt ttaaccaatc acgatcagga cagagttggc 1140
tttcaattgc tggaaagtca ggccaaagcc aaacaagcgg cggcgctgtt atttagctcg 1200
ccgggaccca agtacatcta ctacggtgaa gaaattggca tgaccagct aaaaagtggc 1260
catgatgttt acaagcgggc gccaatgttg tgggatcaag ggcacagggc cggctttacc 1320
caggcagaaa aggcttgggt agagcaaaaa gatttattca gccgccattt tgatgcctgg 1380
tggccggatt tcctgacgca gcagttggca gtcggcgatc ggacagtgga agcacaacgt 1440
gcacagccgg actccttggt gaacttgcac cgtctgctga tcgagctgca aaagcagcgg 1500
ccggaaaatgc gtttggctgg ctcgtaccag gtgcaatcgc cacagcaggg tattctgtcg 1560
gtgacgcgtg agctgggcga gcaaagcac gtgtttgtgc tgaatctggc tggtgaaatg 1620
gctgatattg ctgctgtgct ggcagcgcac agcaatctgt cagtaagtgg gggctttgat 1680
ttggatggta atttgttggc tatagatggt ttgttgattt tgcaaagtca gtag 1734

```

<210> 453

<211> 577

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample

<221> DOMAIN

<222> (58)...(440)

<223> Alpha amylase, catalytic domain

<221> SIGNAL

<222> (1)...(34)

<400> 453

```

Met Met Gln Leu Asn Pro Trp Phe Ser Thr Thr Leu Lys Ala Ala Gly
 1           5           10           15
Leu Ala Thr Ala Leu Ala Ala Val Ser Ala Cys Gln Pro Ala Ser Glu
          20           25           30
Ser Ala Tyr Gln Ala Pro Thr Ala Ala Pro Val Glu Gln Thr Glu Ala
          35           40           45
Pro Trp Trp Asp Asp Ala Val Phe Tyr Gln Ile Trp Pro Arg Ser Phe
          50           55           60
Tyr Asp Thr Ser Gly Asp Gly His Gly Asp Phe Asn Gly Met Thr Ala
65           70           75           80
Lys Leu Pro Tyr Leu Ser Glu Leu Gly Ile Asp Ala Ile Trp Leu Thr
          85           90           95
Pro Met Phe Glu Ala Pro Ser Tyr His Gly Tyr Asp Phe Thr Glu Phe
          100          105          110
Tyr Gln Val Glu Ser Asp Tyr Gly Thr Met Ala Asp Phe Glu Arg Phe
          115          120          125
Leu Gln Gln Ala Glu Glu Arg Gly Ile Arg Val Ile Leu Asp Leu Val
          130          135          140

```

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Ile 145	Asn	His	Ile	Ser	Arg 150	Glu	His	Gln	Trp	Phe 155	Lys	Gln	Ser	Ala	Gln 160
Arg	Ile	Glu	Pro	Tyr 165	Ser	Asp	Tyr	Phe	Ile 170	Trp	Arg	Glu	Ser	Leu	Pro 175
Glu	Ala	Gly	Lys 180	Gly	Trp	Gly	His	Ala 185	Trp	Ser	Asp	Asn	Asp	Met	Pro 190
Glu	Ala	Val 195	Trp	His	Phe	Asp	Glu 200	Thr	Arg	Gly	Glu	Tyr 205	Tyr	Tyr	Ala
Ala	Phe 210	Gly	Ala	Thr	Gln	Pro 215	Asp	Leu	Asn	Leu	Glu 220	His	Pro	Asp	Val
Ile 225	Glu	Glu	Met	Lys	Arg 230	Met	Ala	Lys	Phe	Trp 235	Leu	Asp	Lys	Gly	Val 240
Asp	Gly	Phe	Arg	Leu 245	Asp	Ala	Val	Arg	Phe 250	Val	Ile	Glu	Arg	Gly	Pro 255
Lys	Gly	Gln	Ala 260	Asp	Leu	Pro	Glu	Thr 265	Ile	Asp	Tyr	Trp	Lys 270	Asp	Phe
Asn	Ala	Phe 275	Val	Lys	Ser	Val	Asn 280	Pro	Glu	Ala	Tyr	Leu 285	Val	Gly	Glu
Ala	Trp 290	Val	Asp	Ile	Pro	Val 295	Ala	Ala	Lys	Tyr	Trp 300	Gly	Glu	Gly	Lys
Gly 305	Leu	Asp	Gln	Gly	Phe 310	Asp	Phe	Glu	Phe	Gly 315	Tyr	Ile	Val	Leu	Asp 320
Met	Leu	Gln	Gln	Gly 325	Leu	Ala	Thr	Glu	Ala 330	Gln	Phe	Gly	Thr	Met	Ser 335
Gln	Gln	Ser	Gly 340	Gln	Ala	Glu	Asp	Gly 345	Ala	Leu	His	Pro	Leu 350	Ala	Gln
Asn	Val	Gln 355	Arg	Arg	Leu	Glu	Ser 360	Gly	Ala	Pro	Leu	Ala 365	Phe	Phe	Thr
Pro	Phe 370	Leu	Thr	Asn	His	Asp 375	Gln	Asp	Arg	Val	Gly 380	Phe	Gln	Leu	Leu
Glu 385	Ser	Gln	Ala	Lys	Ala 390	Lys	Gln	Ala	Ala	Ala 395	Leu	Leu	Phe	Ser	Ser 400
Pro	Gly	Thr	Lys	Tyr 405	Ile	Tyr	Tyr	Gly	Glu 410	Glu	Ile	Gly	Met	Thr	Gln 415
Leu	Lys	Ser	Gly 420	His	Asp	Val	Tyr	Lys 425	Arg	Ala	Pro	Met	Leu 430	Trp	Asp
Gln	Gly	His 435	Gln	Ala	Gly	Phe	Thr 440	Gln	Ala	Glu	Lys	Ala 445	Trp	Val	Glu
Gln	Lys 450	Asp	Leu	Phe	Ser	Arg 455	His	Phe	Asp	Ala	Trp 460	Trp	Pro	Asp	Phe
Leu 465	Thr	Gln	Gln	Leu	Ala 470	Val	Gly	Asp	Arg	Thr 475	Val	Glu	Ala	Gln	Arg 480
Ala	Gln	Pro	Asp	Ser 485	Leu	Trp	Asn	Leu	His 490	Arg	Leu	Leu	Ile	Glu	Leu 495
Gln	Lys	Gln	Arg 500	Pro	Glu	Met	Arg	Leu 505	Ala	Gly	Ser	Tyr	Gln 510	Val	Gln
Ser	Pro	Gln 515	Gln	Gly	Ile	Leu	Ser 520	Val	Thr	Arg	Glu	Leu 525	Gly	Glu	Gln
Ser	Thr 530	Val	Phe	Val	Leu	Asn 535	Leu	Ala	Gly	Glu	Met 540	Ala	Asp	Ile	Ala
Ala 545	Val	Leu	Ala	Ala	His 550	Ser	Asn	Leu	Ser	Val 555	Ser	Trp	Gly	Phe	Asp 560
Leu	Asp	Gly	Asn	Leu 565	Leu	Ala	Ile	Asp	Gly 570	Leu	Leu	Ile	Leu	Gln	Ser 575
Gln															

<210> 454
 <211> 1341
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 454
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 ttgttattta tagttgcgac cgtgatactg gcagttctgg tgtcgttttc ggcagagctc

60
 120

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```

gaagcaagag tcaagcctac caatgctgtt atgctccagg gattcggctg gaactctcag 180
tcccagggca ccccaagcaa atgggtacaat ttgattggcg acagggccaa agacatagcc 240
gatctagggt taacgatggt ctgggttccg ccggtgagtc gctctgtgag tcctcagggc 300
tatcttccgg gcgactggta tgatctcgga acagagaagg acccgacgtt ttacggccat 360
aaggatagcc ttttgtacgc ttttaagcgag ctgaaaaagc acggtatttc tccgatcgcc 420
gacatagtca taaaccacag gtgcgccagc catcaggaca agaacgggat ttggaacatc 480
taccattttc cgtcaggaaa ggcgcgctgg gagcaatggg cgatctgccg cggccagttt 540
ggtggacagg gaaaccctga cacaggcgag agctaccatg cggctccaga cgttgatcat 600
acgaacgtca gggttcagcg ggatataatc gagtggatga actggctgaa atctcttggc 660
tttgaagggtt ggcgttatga ctactcaaaa ggctaccatc caagggtatac tgccctttat 720
gaccgggtcta ccaggccccgt attgtctgta ggtgagatat ggacaaacat gtccttcaac 780
ggctcttatc ttaaccctaa tcaggacgcc caccgacagc agctctgcga ctggcttgac 840
ggggccggca actatgccac cgcctttgac ttcactacga aggggattct tcaggtagcc 900
gttaaagggt agtactggcg actacgcgac aaggacggca aggcgtcggg attaataggc 960
tggtggcccg aacgcgcagt cacatttgtc gataatcacg ataccggttc ccagcaaagc 1020
cactggccat ttcccgccaa caaggttatg caggggtacg catatattct tacacatcca 1080
ggcataccct gcattttctg ggagcatgtc tacgactgga aactacgtga accgatcaaa 1140
aggcttgctc agatcaggaa gaaatatgga ttgcacagca agagcagttc taagatcgtt 1200
agagccgagc agaatcttta tgctgcgac attgacggaa aggtagcctt gaagcttggc 1260
agcgtgact ggtcaccgcc aggcaatgat tttgaacttc tcacacatgg cgaccagttt 1320
gcagtttggg gcagaaaata a 1341

```

<210> 455
 <211> 446
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(37)

<221> DOMAIN
 <222> (49)...(386)
 <223> Alpha amylase, catalytic domain

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<400> 455
Met Asp Leu Leu Glu Tyr Lys Asn Thr Ile Gln Arg Arg Gln Thr Met
 1          5          10          15
Thr Asp Arg Lys Leu Leu Phe Ile Val Ala Thr Val Ile Leu Ala Val
          20          25          30
Leu Val Ser Phe Ser Ala Glu Leu Glu Ala Arg Val Lys Pro Thr Asn
          35          40          45
Ala Val Met Leu Gln Gly Phe Gly Trp Asn Ser Gln Ser Arg Gly Thr
          50          55          60
Pro Ser Lys Trp Tyr Asn Leu Ile Gly Asp Arg Ala Lys Asp Ile Ala
          65          70          75          80
Asp Leu Gly Val Thr Met Val Trp Phe Pro Val Ser Arg Ser Val
          85          90          95
Ser Pro Gln Gly Tyr Leu Pro Gly Asp Trp Tyr Asp Leu Gly Thr Glu
          100          105          110
Lys Asp Pro Thr Phe Tyr Gly His Lys Asp Ser Leu Leu Tyr Ala Leu
          115          120          125
Ser Glu Leu Lys Lys His Gly Ile Ser Pro Ile Ala Asp Ile Val Ile
          130          135          140
Asn His Arg Cys Ala Ser His Gln Asp Lys Asn Gly Ile Trp Asn Ile
          145          150          155          160
Tyr His Phe Pro Ser Gly Lys Ala Arg Trp Glu Gln Trp Ala Ile Cys
          165          170          175
Arg Gly Gln Phe Gly Gly Gln Gly Asn Pro Asp Thr Gly Glu Ser Tyr
          180          185          190
His Ala Ala Pro Asp Val Asp His Thr Asn Val Arg Val Gln Arg Asp
          195          200          205
Ile Ile Glu Trp Met Asn Trp Leu Lys Ser Leu Gly Phe Glu Gly Trp
          210          215          220
Arg Tyr Asp Tyr Ser Lys Gly Tyr His Pro Arg Tyr Thr Ala Leu Tyr
          225          230          235          240

```

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Asp	Arg	Ser	Thr	Arg	Pro	Val	Leu	Ser	Val	Gly	Glu	Ile	Trp	Thr	Asn
				245					250					255	
Met	Ser	Phe	Asn	Gly	Ser	Tyr	Leu	Asn	Pro	Asn	Gln	Asp	Ala	His	Arg
			260					265					270		
Gln	Gln	Leu	Cys	Asp	Trp	Leu	Asp	Gly	Ala	Gly	Asn	Tyr	Ala	Thr	Ala
		275					280					285			
Phe	Asp	Phe	Thr	Thr	Lys	Gly	Ile	Leu	Gln	Val	Ala	Val	Lys	Gly	Glu
	290					295					300				
Tyr	Trp	Arg	Leu	Arg	Asp	Lys	Asp	Gly	Lys	Ala	Ser	Gly	Leu	Ile	Gly
	305				310					315					320
Trp	Trp	Pro	Glu	Arg	Ala	Val	Thr	Phe	Val	Asp	Asn	His	Asp	Thr	Gly
				325					330					335	
Ser	Gln	Gln	Ser	His	Trp	Pro	Phe	Pro	Gly	Asn	Lys	Val	Met	Gln	Gly
			340					345					350		
Tyr	Ala	Tyr	Ile	Leu	Thr	His	Pro	Gly	Ile	Pro	Cys	Ile	Phe	Trp	Glu
		355					360					365			
His	Val	Tyr	Asp	Trp	Lys	Leu	Arg	Glu	Pro	Ile	Lys	Arg	Leu	Val	Gln
	370					375						380			
Ile	Arg	Lys	Lys	Tyr	Gly	Leu	His	Ser	Lys	Ser	Ser	Leu	Lys	Ile	Val
	385				390					395					400
Arg	Ala	Glu	Gln	Asn	Leu	Tyr	Ala	Ala	Ile	Ile	Asp	Gly	Lys	Val	Ala
				405					410					415	
Leu	Lys	Leu	Gly	Ser	Ala	Asp	Trp	Ser	Pro	Pro	Gly	Asn	Asp	Phe	Glu
			420					425					430		
Leu	Leu	Thr	His	Gly	Asp	Gln	Phe	Ala	Val	Trp	Gly	Arg	Lys		
		435					440					445			

<210> 456
 <211> 1734
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 456	
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gcgcctgtcg agcagaccga agcgccctgg tgggacgatg cggttttcta ccaaactctgg	180
ccgcgtagtt ttacgatac cagcggtgat ggtcatggcg actttaatgg catgaccgcc	240
aaactgcctt atttgtcgga gcttggcatt gatgccatct ggctgacgcc aatgtttgag	300
gcgccgtctt atcatggcta tgactttacc gagttttatc aggttgaatc ggactatggc	360
accatggccg acttcgagcg ctttttgcaa caggctgaag agcgcggtat ccgggtgatt	420
ctggatttgg tgatcaacca catttcccggt gaacaccagt ggtttcagcg ctccgccgag	480
cgatttgagc cgtacagcga ttactttatc tggcgggaaa gcctgccaga aaccggcaaa	540
ggctggggtc atgcctggtc agataacgat atgcctgaag cggctctggca ttttgatgag	600
acccggggcg agtactacta cgccgccttt ggtgccaccc agccggatct gaatttgga	660
caccgggatg tgatcgaaga aatgaaacgg atggccaagt tctggctgga taagggggtc	720
gatggcttcc ggctcgatgc ggtgcgcttt gtgatagagc gagggccgaa aggccaggcc	780
gacttgcccg agaccattga ttactggaaa gacttcaatg cctttgtcaa aagcgtaaat	840
ccagaagcct acttagtggg cgaagcctgg gtcgatattc cgggtggcggc gaaatactgg	900
ggggaaggca aagggtgga tcagggtctt gatthtgaat ttggctacat cgtgctggat	960
atgctgcagc aaggcttggc cacagaagcc cagtttgga ccatgagcca gcagtccggt	1020
caggcggaag acggcgctt gcattccgtg gcgcagaatg tgcagcgccg cttggagtcg	1080
ggcgaccgcg tgaccttctt cacgcccttt ttaaccaatc acgatcagga cagagttggc	1140
tttcaattgc tggaaaacca ggccaaagcc aaacaagcgg cggcactgtt attcagctcg	1200
ccgggcagca agtacatcta ctacggtgaa gaaattggca tgactcagct caaaagtggc	1260
catgatgtct acaagcgggc gccgatgttg tgggatcaag ggcatacaggc cggctttacc	1320
caggcagaaa aggcctgggt ggagcaaaaa gatttattca gccgccattt tgatgcctgg	1380
tggccggatt tcctgacaca gcagctggca gtcggcgatc ggacagtggc agcacagcgc	1440
gctcaaccgg actccttgtg gaacttgcac cgtctgctga tcgcgatgca aaagcagcgg	1500
ccggaaaatgc gtttggcggg ctcgtatcag gtgcaatcgc cacagcaggg tattttgacg	1560
gtggtgcggg agctggataa cagccgctcg gtcctttgtac tgaatctggc tggtgaaacg	1620
gccgatatcg catcggtgat ggctgcctat ggcgagctgt cggtcacttg gggctttgac	1680
ctggacggca gcttgctggc gaccgatggt ctggtcatct tgcaatcgcg gtaa	1734

<210> 457
 <211> 577

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<212> PRT
<213> Unknown

<220>
<223> Obtained from an environmental sample

<221> SIGNAL
<222> (1)...(26)

<221> DOMAIN
<222> (58)...(440)
<223> Alpha amylase, catalytic domain

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<400> 457
Met Met Gln Leu Asn Pro Trp Phe Ser Ala Ser Leu Lys Ala Ala Gly
 1      5      10      15
Leu Ala Thr Ala Leu Ala Ala Val Ser Ala Cys Gln Pro Ala Pro Glu
      20      25      30
Ser Gly Tyr Gln Ala Pro Thr Ala Ala Pro Val Glu Gln Thr Glu Ala
      35      40      45
Pro Trp Trp Asp Asp Ala Val Phe Tyr Gln Ile Trp Pro Arg Ser Phe
      50      55      60
Tyr Asp Thr Ser Gly Asp Gly His Gly Asp Phe Asn Gly Met Thr Ala
      65      70      75      80
Lys Leu Pro Tyr Leu Ser Glu Leu Gly Ile Asp Ala Ile Trp Leu Thr
      85      90      95
Pro Met Phe Glu Ala Pro Ser Tyr His Gly Tyr Asp Phe Thr Glu Phe
      100      105      110
Tyr Gln Val Glu Ser Asp Tyr Gly Thr Met Ala Asp Phe Glu Arg Phe
      115      120      125
Leu Gln Gln Ala Glu Glu Arg Gly Ile Arg Val Ile Leu Asp Leu Val
      130      135      140
Ile Asn His Ile Ser Arg Glu His Gln Trp Phe Gln Arg Ser Ala Glu
      145      150      155      160
Arg Ile Glu Pro Tyr Ser Asp Tyr Phe Ile Trp Arg Glu Ser Leu Pro
      165      170      175
Glu Thr Gly Lys Gly Trp Gly His Ala Trp Ser Asp Asn Asp Met Pro
      180      185      190
Glu Ala Val Trp His Phe Asp Glu Thr Arg Gly Glu Tyr Tyr Tyr Ala
      195      200      205
Ala Phe Gly Ala Thr Gln Pro Asp Leu Asn Leu Glu His Pro Asp Val
      210      215      220
Ile Glu Glu Met Lys Arg Met Ala Lys Phe Trp Leu Asp Lys Gly Val
      225      230      235      240
Asp Gly Phe Arg Leu Asp Ala Val Arg Phe Val Ile Glu Arg Gly Pro
      245      250      255
Lys Gly Gln Ala Asp Leu Pro Glu Thr Ile Asp Tyr Trp Lys Asp Phe
      260      265      270
Asn Ala Phe Val Lys Ser Val Asn Pro Glu Ala Tyr Leu Val Gly Glu
      275      280      285
Ala Trp Val Asp Ile Pro Val Ala Ala Lys Tyr Trp Gly Glu Gly Lys
      290      295      300
Gly Leu Asp Gln Gly Phe Asp Phe Glu Phe Gly Tyr Ile Val Leu Asp
      305      310      315      320
Met Leu Gln Gln Gly Leu Ala Thr Glu Ala Gln Phe Gly Thr Met Ser
      325      330      335
Gln Gln Ser Gly Gln Ala Glu Asp Gly Ala Leu His Pro Leu Ala Gln
      340      345      350
Asn Val Gln Arg Arg Leu Glu Ser Gly Ala Pro Leu Ala Phe Phe Thr
      355      360      365
Pro phe Leu Thr Asn His Asp Gln Asp Arg Val Gly Phe Gln Leu Leu
      370      375      380
Glu Asn Gln Ala Lys Ala Lys Gln Ala Ala Ala Leu Leu Phe Ser Ser
      385      390      395      400
Pro Gly Ser Lys Tyr Ile Tyr Tyr Gly Glu Glu Ile Gly Met Thr Gln
      405      410      415
Leu Lys Ser Gly His Asp Val Tyr Lys Arg Ala Pro Met Leu Trp Asp
      420      425      430

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Gln Gly His Gln Ala Gly Phe Thr Gln Ala Glu Lys Ala Trp Val Glu
 435 440 445
 Gln Lys Asp Leu Phe Ser Arg His Phe Asp Ala Trp Trp Pro Asp Phe
 450 455 460
 Leu Thr Gln Gln Leu Ala Val Gly Asp Arg Thr Val Ala Ala Gln Arg
 465 470 475 480
 Ala Gln Pro Asp Ser Leu Trp Asn Leu His Arg Leu Leu Ile Ala Met
 485 490 495
 Gln Lys Gln Arg Pro Glu Met Arg Leu Ala Gly Ser Tyr Gln Val Gln
 500 505 510
 Ser Pro Gln Gln Gly Ile Leu Thr Val Val Arg Glu Leu Asp Asn Ser
 515 520 525
 Arg Ser Val Phe Val Leu Asn Leu Ala Gly Glu Thr Ala Asp Ile Ala
 530 535 540
 Ser Val Met Ala Ala Tyr Gly Glu Leu Ser Val Thr Trp Gly Phe Asp
 545 550 555 560
 Leu Asp Gly Ser Leu Leu Ala Thr Asp Gly Leu Val Ile Leu Gln Ser
 565 570 575
 Arg

<210> 458
 <211> 1698
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 458
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 acctatgact ttttgagttt gtttccgaat catcgaaact attatcaaat actcgttcgt 180
 agctttgccg acggcaatgg cgatggcatc ggtgatttta ttggcttaaa agaagccctt 240
 ccacatcttg ccgatttagg gattgaaggc ttatggctgt tgccgattca tccttcacca 300
 agttaccacg ggtatgacgt gacagacttt agagcgatca atccggatta tggtagcatg 360
 gaagattttg aagcgtttat tgcagaagca cagcgtttga acattgacgt cgtgatcgat 420
 tatgtcatca atcattcttc caatcaacac ccttggtttc aaaatttcaa acagggtatc 480
 gccccttacg atcaatttta tcgtcgtatt gataacactg atcccagatg gaacacgcgt 540
 ggttcttggg gtcaaagcat ctggcactca ctgggggatg gaaccgctta tgtgggctat 600
 tttggcgggt acatgcctga tttgaactgg tcgaatgacg ctttagttga agaaatgatc 660
 gacatcgggtc ttttctggct tgaaaaaggc gtgagtgggt tccgtttaga tgcggccttg 720
 catttgtagt ccaataacga agtgccgtca agcaccacag cgttagaaga aaccttattt 780
 atgctagagt attttgagtt tcgtttaaaa gagaaatttc ctgatacttt cattgtgggt 840
 gaaatttggg atgcttttag tatttacaac atgttttacc gtagtatgga ctcagttttt 900
 cactttgact ttggtcactt ggtcgtgaac acgattaaca ctggttctaa tcgcaactat 960
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 ccgaatctcg gcccacacgc ccacctgat aaactgcgtt tagccgcaga gatgttattg 1140
 actgtccctg gaagtccgtt tttttattat ggtgaagaac tcggcatgaa aggggaagcc 1200
 tctggtcgag cccaacgtg ggattcaacg attcgtttac cttacttgtg ggacgacgaa 1260
 cgtcaaccaa cgtggacgtt tgaccggttt gggtttgcg atacgtttta tgtcgatgtc 1320
 ccgaccttgc cagaacaaca agccgatccg ttatcactct ttaacacata ccgtacctta 1380
 ctacatttaa gacaagcgca tcctgcctta agactcggtt ctattgaagc gtttgaagaa 1440
 aatcatgcgg gcttacaagg gttttaccgt atttttgaat acagcacctt ccaagaagtg 1500
 gtcttagtgc ttcataacgt gagtgatgag actttgaacg tgactcgttt acctgaaggt 1560
 gagatgattt acctaatgac acatcttgaa gattttccag cgctcgaaca actcaatcag 1620
 acgcttgata tcggccctag aagtacgctc atgattcgtt taagtgagac agcgatggag 1680
 cgtttcttca atgaatga 1698

<210> 459
 <211> 565
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<221> DOMAIN

<222> (56)...(464)

<223> Alpha amylase, catalytic domain

<221> SIGNAL

<222> (1)...(29)

<400> 459

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Met Phe Lys Val Ser Leu Arg Ser Lys Asp Met Lys Lys Leu Ser Leu
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Ile Val Thr Ile Leu Val Leu Ala Leu Thr Leu Ser Ala Cys Gln Arg
 25      30      35      40      45
Glu Thr Glu Glu Val Gly Glu Thr Tyr Asp Phe Leu Ser Leu Phe
Pro Asn His Arg Asn Tyr Tyr Gln Ile Leu Val Arg Ser Phe Ala Asp
 50      55      60      65      70      75
Gly Asn Gly Asp Gly Ile Gly Asp Phe Ile Gly Leu Lys Glu Ala Leu
Pro His Leu Ala Asp Leu Gly Ile Glu Gly Leu Trp Leu Leu Pro Ile
 85      90      95      100      105      110
His Pro Ser Pro Ser Tyr His Gly Tyr Asp Val Thr Asp Phe Arg Ala
Ile Asn Pro Asp Tyr Gly Thr Met Glu Asp Phe Glu Ala Phe Ile Ala
 115      120      125      130      135      140
Glu Ala Gln Arg Leu Asn Ile Asp Val Val Ile Asp Tyr Val Ile Asn
His Ser Ser Asn Gln His Pro Trp Phe Gln Asn Phe Lys Gln Gly Ile
 145      150      155      160      165      170
Ala Pro Tyr Asp Gln Phe Tyr Arg Arg Ile Asp Asn Thr Asp Pro Arg
Trp Asn Thr Arg Gly Ser Trp Gly Gln Ser Ile Trp His Ser Leu Gly
 180      185      190      195      200      205
Asp Gly Thr Ala Tyr Val Gly Tyr Phe Gly Gly Tyr Met Pro Asp Leu
Asn Trp Ser Asn Asp Ala Leu Val Glu Glu Met Ile Asp Ile Gly Leu
 210      215      220      225      230      235
Phe Trp Leu Glu Lys Gly Val Ser Gly Phe Arg Leu Asp Ala Ala Leu
His Leu Leu Ala Asn Asn Glu Val Pro Ser Ser Thr Pro Ala Leu Glu
 245      250      255      260      265      270
Glu Thr Leu Phe Met Leu Glu Tyr Phe Glu Phe Arg Leu Lys Glu Lys
Phe Pro Asp Thr Phe Ile Val Gly Glu Ile Trp Asp Ala Phe Ser Ile
 275      280      285      290      295      300
Tyr Asn Met Phe Tyr Arg Ser Met Asp Ser Val Phe His Phe Asp Phe
Gly His Leu Val Val Asn Thr Ile Asn Thr Gly Ser Asn Arg Asn Tyr
 305      310      315      320      325      330
Ala Asp Gln Ile Val Arg Trp His Asn Gln Ala Gln Ala Val Asp Asp
Val Val Ile Glu Ala Pro Phe Leu Arg Asn His Asp Gln Asn Arg Leu
 340      345      350      355      360      365
Ala Ser Ala Ser Ser Pro Gly Ser Pro Asn Leu Gly Pro Asn Ala His
Pro Asp Lys Leu Arg Leu Ala Glu Met Leu Leu Thr Val Pro Gly
 370      375      380      385      390      395
Ser Pro Phe Ile Tyr Tyr Gly Glu Glu Leu Gly Met Lys Gly Glu Ala
Ser Gly Arg Ala Pro Thr Trp Asp Ser Thr Ile Arg Leu Pro Tyr Leu
 405      410      415      420      425      430
Trp Asp Asp Glu Arg Gln Pro Thr Trp Thr Phe Asp Arg Phe Gly Phe
Val Asp Thr Phe Asn Val Asp Val Pro Thr Leu Pro Glu Gln Gln Ala
 435      440      445      450      455      460
Asp Pro Leu Ser Leu Phe Asn Thr Tyr Arg Thr Leu Leu His Leu Arg
Gln Ala His Pro Ala Leu Arg Leu Gly Thr Ile Glu Ala Phe Glu Glu
 465      470      475      480

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Asn His Ala Gly Leu Gln Gly Phe Tyr Arg Ile Phe Glu Tyr Ser Thr
 485 490 495
 Phe Gln Glu Val Val Leu Val Leu His Asn Val Ser Asp Glu Thr Leu
 500 505 510
 Asn Val Thr Arg Leu Pro Glu Gly Glu Met Ile Tyr Leu Ser Ala His
 515 520 525
 Leu Glu Asp Phe Pro Ala Leu Glu Gln Leu Asn Gln Thr Leu Asp Ile
 530 535 540
 Gly Pro Arg Ser Thr Leu Met Ile Arg Leu Ser Glu Thr Ala Met Glu
 545 550 555 560
 Arg Phe Phe Asn Glu
 565

<210> 460
 <211> 1752
 <212> DNA
 <213> *Cochliobolus heterostrophus* ATCC 48331

<220>
 <221> CDS
 <222> (1)...(1752)
 <223> Exon

<400> 460
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 1 5 10 15
 ata tgg ccg gcg tct ttc aaa gat tcg aac agt gat ggc ata ggc gat 96
 Ile Trp Pro Ala Ser Phe Lys Asp Ser Asn Ser Asp Gly Ile Gly Asp
 20 25 30
 ctc aat gga gtt ctc agc aag ctc gac tac ctc cga tca ctg gaa gtc 144
 Leu Asn Gly Val Leu Ser Lys Leu Asp Tyr Leu Arg Ser Leu Glu Val
 35 40 45
 gat ctc atc tgg ttt tct cct gtc tat gat tcg ccc aac aga gac atg 192
 Asp Leu Ile Trp Phe Ser Pro Val Tyr Asp Ser Pro Asn Arg Asp Met
 50 55 60
 gga tat gat gtc tct gat tac gaa tcc att cac cca cag tat ggc acc 240
 Gly Tyr Asp Val Ser Asp Tyr Glu Ser Ile His Pro Gln Tyr Gly Thr
 65 70 75 80
 att gcc gac atg gaa aac ctc atc aaa gag gcc aag aaa cgt aac atc 288
 Ile Ala Asp Met Glu Asn Leu Ile Lys Glu Ala Lys Lys Arg Asn Ile
 85 90 95
 aaa att gtc atg gat atg gtc aca aat cat tcg tct gac caa cac cgc 336
 Lys Ile Val Met Asp Met Val Thr Asn His Ser Ser Asp Gln His Arg
 100 105 110
 tgg ttc ttg gag agt tgc aaa tct cgg tca ggg cca tat agt gac ttt 384
 Trp Phe Leu Glu Ser Cys Lys Ser Arg Ser Gly Pro Tyr Ser Asp Phe
 115 120 125
 tac atc tgg ccg gat cca aaa tac gat agc gaa ggt aac cga caa cca 432
 Tyr Ile Trp Arg Asp Pro Lys Tyr Asp Ser Glu Gly Asn Arg Gln Pro
 130 135 140
 cca aat aat tgg cgc act ggg tca cga tat gga aaa agt aca tgg aag 480
 Pro Asn Asn Trp Arg Thr Gly Ser Arg Tyr Gly Lys Ser Thr Trp Lys
 145 150 155 160
 tac gtc gaa gca aga gat cag tac tat ttc tgc ttt gga act act tat 528
 Tyr Val Glu Ala Arg Asp Gln Tyr Tyr Phe Cys Phe Gly Thr Thr Tyr
 165 170 175

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1001827087_1.txt																	
cag Gln	cct Pro	aac Asn	ttg Leu 180	aat Asn	tgg Trp	ttc Phe	tgc Cys	gac Asp 185	gat Asp	ttg Leu	aga Arg	cgc Arg	gca Ala 190	ata Ile	tac Tyr	576	
gac Asp	agc Ser	gcg Ala 195	gtg Val	aaa Lys	ttt Phe	tgg Trp	ttg Leu 200	gac Asp	aga Arg	ggg Gly	gtc Val	gat Asp 205	ggg Gly	ttc Phe	agg Arg	624	
atg Met 210	gat Asp	ttg Leu	gtc Val	ggg Gly	tta Leu	tac Tyr 215	tgg Trp	aaa Lys	gat Asp	cca Pro	acc Thr 220	ttt Phe	cct Pro	gat Asp	gca Ala	672	
aaa Lys 225	gaa Glu	gtt Val	tat Tyr	cct Pro	gga Gly 230	gag Glu	gcg Ala	ctg Leu	caa Gln	ccg Pro 235	ctt Leu	gac Asp	gga Gly	tcg Ser	cgg Arg 240	720	
tgt Cys	ttg Leu	aat Asn	ggc Gly	gaa Glu 245	gaa Glu	gtg Val	cat His	gtc Val	tgg Trp 250	tta Leu	aag Lys	gag Glu	ttg Leu	aag Lys 255	gaa Glu	768	
aga Arg	atc Ile	acg Thr	caa Gln 260	gac Asp	ttt Phe	gga Gly	cag Gln	gac Asp 265	atc Ile	gtg Val	ttg Leu	att Ile	ggg Gly 270	gag Glu	ctc Leu	816	
ccc Pro	tcg Ser	act Thr 275	gac Asp	cgg Arg	gag Glu	aca Thr	ttc Phe 280	cta Leu	cga Arg	tac Tyr	ata Ile	tct Ser 285	cct Pro	gaa Glu	tcg Ser	864	
aga Arg 290	gaa Glu	cta Leu	gag Glu	atg Met	gct Ala	ctc Leu 295	gat Asp	aca Thr	gac Asp	atc Ile	ttc Phe 300	atc Ile	gtg Val	gga Gly	aat Asn	912	
gaa Glu 305	tgg Trp	aca Thr	gat Asp	ggg Gly	ctg Leu 310	tac Tyr	gag Glu	ctc Leu	aaa Lys	aag Lys 315	cca Pro	gag Glu	cta Leu	ccg Pro	cta Leu 320	960	
ttc Phe	aaa Lys	gat Asp	gcc Ala	gtc Val 325	ctg Leu	aag Lys	acg Thr	cag Gln	agc Ser 330	ctt Leu	ctt Leu	gac Asp	gat Asp	gga Gly 335	ggc Gly	1008	
tgg Trp	cct Pro	act Thr	gca Ala 340	ttc Phe	ctc Leu	gag Glu	aat Asn	cat His 345	gat Asp	ttc Phe	gct Ala	cgc Arg	agt Ser 350	gta Val	tcc Ser	1056	
cgt Arg	ttt Phe	ggg Gly 355	cca Pro	ggg Gly	gag Glu	ggg Gly	acc Thr 360	tat Tyr	cgc Arg	gaa Glu	gct Ala	gcc Ala 365	gca Ala	agg Arg	atg Met	1104	
ctt Leu	gct Ala 370	ctc Leu	atg Met	gct Ala	gca Ala	aca Thr 375	ctt Leu	tcc Ser	gga Gly	acc Thr	ttt Phe 380	ttc Phe	atc Ile	tac Tyr	cag Gln	1152	
ggg Gly 385	caa Gln	gaa Glu	att Ile	gga Gly	atg Met 390	acc Thr	aac Asn	gtc Val	cca Pro	tca Ser 395	cat His	tgg Trp	agg Arg	cga Arg	gat Asp 400	1200	
gac Asp	ttc Phe	aaa Lys	gat Asp	aac Asn 405	gcc Ala	gac Asp	ctc Leu	gaa Glu	cac His 410	atc Ile	gaa Glu	gat Asp	att Ile	gat Asp 415	cca Pro	1248	
atc Ile	aaa Lys	gac Asp	gct Ala 420	gag Glu	gcg Ala	gtg Val	gag Glu	aaa Lys 425	tcc Ser	gtt Val	gat Asp	gcg Ala	ctt Leu 430	acc Thr	aca Thr	1296	
tgg Trp	ggg Gly	cga Arg 435	gat Asp	aat Asn	gga Gly	aga Arg	acg Thr 440	cca Pro	gtc Val	cag Gln	tgg Trp	tca Ser 445	gac Asp	aaa Lys	gat Asp	1344	

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cat His	gcc Ala 450	ggg Gly	ttt Phe	agc Ser	gac Asp	gtg Val 455	aaa Lys	ccc Pro	tgg Trp	att Ile	agg Arg 460	gtc Val	atc Ile	gac Asp	aac Asn	1392
tac Tyr 465	act Thr	tac Tyr	ata Ile	aac Asn	gtc Val 470	gaa Glu	gag Glu	caa Gln	ctt Leu	gga Gly 475	aac Asn	ccg Pro	aac Asn	agc Ser	ata Ile 480	1440
cgt Arg	tct Ser	ttc Phe	tgg Trp	gta Val 485	aag Lys	atg Met	atg Met	aag Lys	atg Met 490	agg Arg	aaa Lys	gag Glu	atg Met	tat Tyr 495	gat Asp	1488
gtg Val	tta Leu	gcc Ala	tgc Cys 500	ggg Gly	cat His	ttt Phe	cgt Arg	ctc Leu 505	gtt Val	gac Asp	agg Arg	gac Asp	aac Asn 510	ctc Leu	agt Ser	1536
gtg Val	ttc Phe	tcg Ser 515	tat Tyr	acg Thr	aaa Lys	aca Thr	agt Ser 520	gct Ala	gat Asp	gga Gly	aag Lys	acc Thr 525	gag Glu	atg Met	cta Leu	1584
gtg Val 530	gta Val	ttg Leu	aac Asn	ttt Phe	tcg Ser	agc Ser 535	gag Glu	gag Glu	gct Ala	cgg Arg	gct Ala 540	ccg Pro	agg Arg	caa Gln	atc Ile	1632
ggg Gly 545	aaa Lys	gat Asp	cta Leu	atg Met	cat His 550	ctc Leu	gtt Val	tgc Cys	acg Thr	aca Thr 555	gct Ala	gcg Ala	atg Met	tcg Ser	acg Thr 560	1680
aag Lys	act Thr	agc Ser	tta Leu	gat Asp 565	gaa Glu	gtc Val	aca Thr	ttg Leu	cag Gln 570	cca Pro	tgg Trp	gag Glu	ggt Gly	cgc Arg 575	ttg Leu	1728
tat Tyr	cga Arg	ctg Leu	gca Ala 580	cca Pro	aag Lys	ttg Leu	tag *									1752

<210> 461
 <211> 583
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

<220>
 <221> DOMAIN
 <222> (16)...(421)

<400> 461
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 Ile Trp Pro Ala Ser Phe Lys Asp Ser Asn Ser Asp Gly Ile Gly Asp
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 Leu Asn Gly Val Leu Ser Lys Leu Asp Tyr Leu Arg Ser Leu Glu Val
 35 40 45
 Asp Leu Ile Trp Phe Ser Pro Val Tyr Asp Ser Pro Asn Arg Asp Met
 50 55 60
 Gly Tyr Asp Val Ser Asp Tyr Glu Ser Ile His Pro Gln Tyr Gly Thr
 65 70 75 80
 Ile Ala Asp Met Glu Asn Leu Ile Lys Glu Ala Lys Lys Arg Asn Ile
 85 90 95
 Lys Ile Val Met Asp Met Val Thr Asn His Ser Ser Asp Gln His Arg
 100 105 110
 Trp Phe Leu Glu Ser Cys Lys Ser Arg Ser Gly Pro Tyr Ser Asp Phe
 115 120 125
 Tyr Ile Trp Arg Asp Pro Lys Tyr Asp Ser Glu Gly Asn Arg Gln Pro
 130 135 140
 Pro Asn Asn Trp Arg Thr Gly Ser Arg Tyr Gly Lys Ser Thr Trp Lys
 145 150 155 160
 Tyr Val Glu Ala Arg Asp Gln Tyr Tyr Phe Cys Phe Gly Thr Thr Tyr

1001827087_1.txt

Gln	Pro	Asn	Leu	165	Asn	Trp	Phe	Cys	Asp	170	Leu	Arg	Arg	Ala	175	Ile	Tyr
Asp	Ser	Ala	Val	180	Lys	Phe	Trp	Leu	Asp	185	Arg	Gly	Val	Asp	190	Gly	Phe
Met	Asp	Leu	Val	195	Gly	Leu	Tyr	Trp	Lys	200	Pro	Thr	Phe	Pro	Asp	Ala	
Lys	Glu	Val	Tyr	210	Pro	Gly	Glu	Ala	Leu	215	Gln	Pro	Leu	Asp	Gly	Ser	Arg
Cys	Leu	Asn	Gly	225	Glu	Glu	Val	His	Val	230	Trp	Leu	Lys	Glu	Leu	Lys	Glu
Arg	Ile	Thr	Gln	245	Asp	Phe	Gly	Gln	Asp	250	Ile	Val	Leu	Ile	Gly	Glu	Leu
Pro	Ser	Thr	Asp	260	Arg	Glu	Thr	Phe	Leu	265	Arg	Tyr	Ile	Ser	Pro	Glu	Ser
Arg	Glu	Leu	Glu	275	Met	Ala	Leu	Asp	Thr	280	Asp	Ile	Phe	Ile	Val	Gly	Asn
Glu	Trp	Thr	Asp	290	Gly	Leu	Tyr	Glu	Leu	295	Lys	Lys	Pro	Glu	Leu	Pro	Leu
Phe	Lys	Asp	Ala	305	Val	Leu	Lys	Thr	Gln	310	Ser	Leu	Leu	Asp	Asp	Gly	Gly
Trp	Pro	Thr	Ala	325	Phe	Leu	Glu	Asn	His	330	Asp	Phe	Ala	Arg	Ser	Val	Ser
Arg	Phe	Gly	Pro	340	Gly	Glu	Gly	Thr	Tyr	345	Arg	Glu	Ala	Ala	Ala	Arg	Met
Leu	Ala	Leu	Met	355	Ala	Ala	Thr	Leu	Ser	360	Gly	Thr	Phe	Phe	Ile	Tyr	Gln
Gly	Gln	Glu	Ile	370	Gly	Met	Thr	Asn	Val	375	Pro	Ser	His	Trp	Arg	Arg	Asp
Asp	Phe	Lys	Asp	385	Asn	Ala	Asp	Leu	Glu	390	His	Ile	Glu	Asp	Ile	Asp	Pro
Ile	Lys	Asp	Ala	405	Glu	Ala	Val	Glu	Lys	410	Ser	Val	Asp	Ala	Leu	Thr	Thr
Trp	Gly	Arg	Asp	420	Asn	Gly	Arg	Thr	Pro	425	Val	Gln	Trp	Ser	Asp	Lys	Asp
His	Ala	Gly	Phe	435	Ser	Asp	Val	Lys	Pro	440	Pro	Trp	Ile	Arg	Val	Ile	Asn
Tyr	Thr	Tyr	Ile	450	Asn	Val	Glu	Gln	Leu	455	Gly	Asn	Pro	Asn	Ser	Ile	
Arg	Ser	Phe	Trp	465	Val	Lys	Met	Met	Lys	470	Met	Arg	Lys	Glu	Met	Tyr	Asp
Val	Leu	Ala	Cys	485	Gly	His	Phe	Arg	Leu	490	Val	Asp	Arg	Asp	Asn	Leu	Ser
Val	Phe	Ser	Tyr	500	Thr	Lys	Thr	Ser	Ala	505	Asp	Gly	Lys	Thr	Glu	Met	Leu
Val	Val	Leu	Asn	515	Phe	Ser	Ser	Glu	Glu	520	Ala	Arg	Ala	Pro	Arg	Gln	Ile
Gly	Lys	Asp	Leu	530	Met	His	Leu	Val	Cys	535	Thr	Thr	Ala	Ala	Met	Ser	Thr
Lys	Thr	Ser	Leu	545	Asp	Glu	Val	Thr	Leu	550	Gln	Pro	Trp	Glu	Gly	Arg	Leu
Tyr	Arg	Leu	Ala	565	Pro	Lys	Leu			570							
				580													

<210> 462
 <211> 1252
 <212> DNA
 <213> Fungi

<220>
 <221> CDS
 <222> (1)...(114)
 <223> Exon

<221> CDS
 <222> (161)...(1252)
 <223> Exon

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1001827087_1.txt

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<400> 462
atg tca aga tct tct acc att ctt ttt gtg ttg gca gca gcc aat ttg      48
Met Ser Arg Ser Ser Thr Ile Leu Phe Val Leu Ala Ala Ala Asn Leu
1      5      10      15

gcg tct cta gta gat gct cat ggc tac atg aac ata ccc tcc agt cga      96
Ala Ser Leu Val Asp Ala His Gly Tyr Met Asn Ile Pro Ser Ser Arg
20      25      30

aca agc atc ggc cat tca gtaagattcc ataaatccat ttctattacc      144
Thr Ser Ile Gly His Ser
35

acattaacaa ttgaag gtt ggc atc gac aca tgt ccc gaa tgt act att ctt      196
Val Gly Ile Asp Thr Cys Pro Glu Cys Thr Ile Leu
40      45      50

gag ccc gtc tct tca tgg cct gat ctc gac gtc gct cca gtt gga cgc      244
Glu Pro Val Ser Ser Trp Pro Asp Leu Asp Val Ala Pro Val Gly Arg
55      60      65

agc ggt cct tgc ggc tac aat gct cgg gtc agc gtc gac tac aat tac      292
Ser Gly Pro Cys Gly Tyr Asn Ala Arg Val Ser Val Asp Tyr Asn Tyr
70      75      80

cct tcc aca aac tgg ggt aac acc ccc gtc acc acc tat aca gct ggt      340
Pro Ser Thr Asn Trp Gly Asn Thr Pro Val Thr Thr Tyr Thr Ala Gly
85      90      95

caa aca gtc gat gta cag tgg tgc gta gac aac aac gga gac cat ggc      388
Gln Thr Val Asp Val Gln Trp Cys Val Asp Asn Asn Gly Asp His Gly
100      105      110

ggt atg ttc agc tgg cgc atc tgt caa gac caa tcc atc gtg gat aag      436
Gly Met Phe Ser Trp Arg Ile Cys Gln Asp Gln Ser Ile Val Asp Lys
115      120      125

ctc atc aca ccc gga tac atc ccc acg gat gcc gag aaa caa gcc gct      484
Leu Ile Thr Pro Gly Tyr Ile Pro Thr Asp Ala Glu Lys Gln Ala Ala
135      140      145

gag gat tgt ttc gac gcc gga tta ctc ccc tgt acc gac gtc acc gga      532
Glu Asp Cys Phe Asp Ala Gly Leu Leu Pro Cys Thr Asp Val Thr Gly
150      155      160

gcc acg tgc gat tac aac gca gat tgc act ccc gga caa gcc tgt tac      580
Ala Thr Cys Asp Tyr Asn Ala Asp Cys Thr Pro Gly Gln Ala Cys Tyr
165      170      175

cga aac gac tgg ttt act tgc aag gga ttc aac agc ggc aca aga tgt      628
Arg Asn Asp Trp Phe Thr Cys Lys Gly Phe Asn Ser Gly Thr Arg Cys
180      185      190

caa gga gtg gat ggc gcc gct ctc ggc tct tgc tac act tcc atc gca      676
Gln Gly Val Asp Gly Ala Ala Leu Gly Ser Cys Tyr Thr Ser Ile Ala
195      200      205

ggt gga tat aca gtg acc aag aaa gtc aag atc cca aac tac gca tca      724
Gly Gly Tyr Thr Val Thr Lys Lys Val Lys Ile Pro Asn Tyr Ala Ser
215      220      225

aat cac acc ctc ata tcc ctg aaa tgg aat tcc ttc caa aca ggc caa      772
Asn His Thr Leu Ile Ser Leu Lys Trp Asn Ser Phe Gln Thr Gly Gln
230      235      240

atc tat cta acc tgc gcc gat atc gcc att gtc ggt tcc ggc agc ggc      820
Ile Tyr Leu Thr Cys Ala Asp Ile Ala Ile Val Gly Ser Gly Ser Gly
245      250      255

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ggc Gly	ggc Gly 260	tcc Ser	tca Ser	agt Ser	acc Thr	acg Thr 265	agc Ser	aaa Lys	aca Thr	tct Ser	tcc Ser 270	acc Thr	tcc Ser	acc Thr	acc Thr	868
ctc Leu 275	agc Ser	aca Thr	tcg Ser	acc Thr	aaa Lys 280	tcg Ser	agc Ser	acc Thr	acc Thr	acc Thr 285	agc Ser	gcc Ala	acc Thr	acg Thr	gca Ala 290	916
acc Thr	aca Thr	agc Ser	tgc Cys	gcc Ala 295	tca Ser	acc Thr	cct Pro	tct Ser	ctg Leu 300	att Ile	ccc Pro	gta Val	acc Thr	ttc Phe 305	aac Asn	964
gag Glu	ctc Leu	gtc Val	acc Thr 310	acc Thr	acc Thr	tac Tyr	gga Gly	caa Gln 315	acc Thr	atc Ile	aaa Lys	gtc Val	gcc Ala 320	ggc Gly	tcg Ser	1012
gtc Val	gcc Ala	gct Ala 325	ctc Leu	gga Gly	aac Asn	tgg Trp	gat Asp 330	gtt Val	tcc Ser	gcc Ala	gcc Ala	gtc Val 335	gca Ala	ttg Leu	tca Ser	1060
gct Ala 340	gta Val	aac Asn	tac Tyr	acg Thr	agt Ser	gcg Ala 345	aat Asn	cca Pro	ttg Leu	tgg Trp	acc Thr 350	gga Gly	acc Thr	gtc Val	atg Met	1108
ttg Leu 355	ccc Pro	ccc Pro	gga Gly	cag Gln	gtg Val 360	gtg Val	cag Gln	tat Tyr	aag Lys	tat Tyr 365	gtg Val	aat Asn	gtg Val	gcg Ala	agt Ser 370	1156
agt Ser	ggg Gly	acg Thr	cca Pro	acg Thr 375	tgg Trp	gaa Glu	aaa Lys	gat Asp	ccc Pro 380	aat Asn	cgc Arg	gtt Val	ttt Phe	acg Thr 385	gtg Val	1204
ccc Pro	gcg Ala	ggt Gly	tgt Cys 390	gcg Ala	aca Thr	aag Lys	ggg Gly	acc Thr 395	gtg Val	ggg Gly	gat Asp	act Thr	tgg Trp 400	aga Arg	tag *	1252

<210> 463
<211> 38
<212> PRT
<213> Fungi

<400> 463
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1 5 10 15
Ala Ser Leu Val Asp Ala His Gly Tyr Met Asn Ile Pro Ser Ser Arg
20 25 30
Thr Ser Ile Gly His Ser
35

<210> 464
<211> 363
<212> PRT
<213> Fungi

<400> 464
Val Gly Ile Asp Thr Cys Pro Glu Cys Thr Ile Leu Glu Pro Val Ser
1 5 10 15
Ser Trp Pro Asp Leu Asp Val Ala Pro Val Gly Arg Ser Gly Pro Cys
20 25 30
Gly Tyr Asn Ala Arg Val Ser Val Asp Tyr Asn Tyr Pro Ser Thr Asn
35 40 45
Trp Gly Asn Thr Pro Val Thr Thr Tyr Thr Ala Gly Gln Thr Val Asp
50 55 60
Val Gln Trp Cys Val Asp Asn Asn Gly Asp His Gly Gly Met Phe Ser
65 70 75 80
Trp Arg Ile Cys Gln Asp Gln Ser Ile Val Asp Lys Leu Ile Thr Pro

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Gly	Tyr	Ile	Pro	85	Thr	Asp	Ala	Glu	Lys	90	Gln	Ala	Ala	Glu	Asp	95	Cys	Phe
Asp	Ala	Gly	Leu	100	Leu	Pro	Cys	Thr	Asp	105	Val	Thr	Gly	Ala	Thr	110	Cys	Asp
Tyr	Asn	Ala	Asp	115	Cys	Thr	Pro	Gly	Gln	120	Ala	Cys	Tyr	Arg	Asn	125	Asp	Trp
Phe	Thr	Cys	Lys	130	Gly	Phe	Asn	Ser	Gly	135	Thr	Arg	Cys	Gln	Gly	140	Val	Asp
Gly	Ala	Ala	Leu	145	Ser	Cys	Tyr	Thr	Ser	150	Ile	Ala	Gly	Gly	Tyr	155	Thr	Thr
Val	Thr	Lys	Lys	160	Val	Lys	Ile	Pro	Asn	165	Tyr	Ala	Ser	Asn	His	170	Thr	Leu
Ile	Ser	Leu	Lys	175	Trp	Asn	Ser	Phe	Gln	180	Thr	Gly	Gln	Ile	Tyr	185	Leu	Thr
Cys	Ala	Asp	Ile	190	Ala	Ile	Val	Gly	Ser	195	Gly	Ser	Gly	Gly	Gly	200	Ser	Ser
Ser	Thr	Thr	Ser	205	Lys	Thr	Ser	Ser	Thr	210	Ser	Thr	Thr	Leu	Ser	215	Thr	Ser
Thr	Lys	Ser	Ser	220	Thr	Thr	Thr	Ser	Ala	225	Thr	Thr	Ala	Thr	Thr	230	Ser	Cys
Ala	Ser	Thr	Pro	235	Ser	Leu	Ile	Pro	Val	240	Thr	Phe	Asn	Glu	Leu	245	Val	Thr
Thr	Thr	Tyr	Gly	250	Gln	Thr	Ile	Lys	Val	255	Ala	Gly	Ser	Val	Ala	260	Ala	Leu
Gly	Asn	Trp	Asp	265	Val	Ser	Ala	Ala	Val	270	Ala	Leu	Ser	Ala	Val	275	Asn	Tyr
Thr	Ser	Ala	Asn	280	Pro	Leu	Trp	Thr	Gly	285	Thr	Val	Met	Leu	Pro	290	Pro	Gly
Gln	Val	Val	Gln	295	Tyr	Lys	Tyr	Val	Asn	300	Val	Ala	Ser	Ser	Gly	305	Thr	Pro
Thr	Trp	Glu	Lys	310	Asp	Pro	Asn	Arg	Val	315	Phe	Thr	Val	Pro	Ala	320	Gly	Cys
Ala	Thr	Lys	Gly	325	Thr	Val	Gly	Asp	Thr	330	Trp	Arg				335		
				340						345						350		
				355						360								

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<212>	DNA
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atcgacacat	gtcccgaatg	tactattctt	gagcccgtct	cttcattggcc	tgatctcgac		180
gtcgtctccag	ttggcagcag	cggctcttgc	ggctacaatg	ctcgggtcag	cgctgcatac		240
aattaccctt	ccacaacatg	gggtaaacac	cccgtcacca	cctatacagc	tggtcaaacaca		300
gtcgaatgtac	agtgggtgct	agacaacaac	ggagaccatg	gcggatatgtt	cagctggcgc		360
atctgtcaag	accaatccat	cgtggataag	ctcatcacac	ccggatacat	ccccacggat		420
gccgagaaaac	aagccgctga	ggattgtttc	gacgccggat	tactccccctg	taccgacgtc		480
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gactggtttta	cttgcaaggg	attcacacagc	ggcacaaagt	gtcaaggagt	ggatggcgcc		600
gctctcggct	cttgctacac	ttccatcgca	ggtggatata	cagtgaacca	gaaagtcaag		660
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ggccaaatct	atctaaccctg	cgccgataatc	gccattgtcg	gttcggcgag	cggcggcggc		780
tcctcaagta	ccacgagcaa	aacatcttcc	acctccacca	ccctcagcac	atcgacaaa		840
tcgagacca	ccaccagcgc	caccacggca	accacaagct	gcgcctcaac	cccttctctg		900
attcccgtaa	ccttcaacga	gctcgtcacc	accacctacg	gacaaacctt	caaagtcgcc		960
ggctcggctg	ccgctctcgg	aaactgggat	gtttccgcgc	ccgtcgcatt	gtcagctgta		1020
aactacacga	gtgcgaatcc	attgtggacc	ggaaccgtca	tgttgcccc	cggacaggtg		1080
gtgcagtata	agtatgtgaa	tgtggcgagt	agtgggacgc	caacgtggga	aaaagatccc		1140
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aqataq							1206

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 Thr Ser Ile Gly His Ser Val Gly Ile Asp Thr Cys Pro Glu Cys Thr
 35 40 45
 Ile Leu Glu Pro Val Ser Ser Trp Pro Asp Leu Asp Val Ala Pro Val
 50 55 60
 Gly Arg Ser Gly Pro Cys Gly Tyr Asn Ala Arg Val Ser Val Asp Tyr
 65 70 75 80
 Asn Tyr Pro Ser Thr Asn Trp Gly Asn Thr Pro Val Thr Thr Tyr Thr
 85 90 95
 Ala Gly Gln Thr Val Asp Val Gln Trp Cys Val Asp Asn Asn Gly Asp
 100 105 110
 His Gly Gly Met Phe Ser Trp Arg Ile Cys Gln Asp Gln Ser Ile Val
 115 120 125
 Asp Lys Leu Ile Thr Pro Gly Tyr Ile Pro Thr Asp Ala Glu Lys Gln
 130 135 140
 Ala Ala Glu Asp Cys Phe Asp Ala Gly Leu Leu Pro Cys Thr Asp Val
 145 150 155 160
 Thr Gly Ala Thr Cys Asp Tyr Asn Ala Asp Cys Thr Pro Gly Gln Ala
 165 170 175
 Cys Tyr Arg Asn Asp Trp Phe Thr Cys Lys Gly Phe Asn Ser Gly Thr
 180 185 190
 Arg Cys Gln Gly Val Asp Gly Ala Ala Leu Gly Ser Cys Tyr Thr Ser
 195 200 205
 Ile Ala Gly Gly Tyr Thr Val Thr Lys Lys Val Lys Ile Pro Asn Tyr
 210 215 220
 Ala Ser Asn His Thr Leu Ile Ser Leu Lys Trp Asn Ser Phe Gln Thr
 225 230 235 240
 Gly Gln Ile Tyr Leu Thr Cys Ala Asp Ile Ala Ile Val Gly Ser Gly
 245 250 255
 Ser Gly Gly Gly Ser Ser Ser Thr Thr Ser Lys Thr Ser Ser Thr Ser
 260 265 270
 Thr Thr Leu Ser Thr Ser Thr Lys Ser Ser Thr Thr Thr Ser Ala Thr
 275 280 285
 Thr Ala Thr Thr Ser Cys Ala Ser Thr Pro Ser Leu Ile Pro Val Thr
 290 295 300
 Phe Asn Glu Leu Val Thr Thr Tyr Gly Gln Thr Ile Lys Val Ala
 305 310 315 320
 Gly Ser Val Ala Ala Leu Gly Asn Trp Asp Val Ser Ala Ala Val Ala
 325 330 335
 Leu Ser Ala Val Asn Tyr Thr Ser Ala Asn Pro Leu Trp Thr Gly Thr
 340 345 350
 Val Met Leu Pro Pro Gly Gln Val Val Gln Tyr Lys Tyr Val Asn Val
 355 360 365
 Ala Ser Ser Gly Thr Pro Thr Trp Glu Lys Asp Pro Asn Arg Val Phe
 370 375 380
 Thr Val Pro Ala Gly Cys Ala Thr Lys Gly Thr Val Gly Asp Thr Trp
 385 390 395 400
 Arg

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 <212> DNA
 <213> Cochliobolus heterostrophus ATCC 48331

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tgc aag acg cta gcc tgt tcc agc ata gcc aac ctt ttt ggc ggc ttc 96
Cys Lys Thr Leu Ala Cys Ser Ser Ile Ala Asn Leu Phe Gly Gly Phe
20 25 30

cct ccc tcc gga tgt cca agc agc gac gat ctt gcc cca aag tct tca 144
Pro Pro Ser Gly Cys Pro Ser Ser Asp Asp Leu Ala Pro Lys Ser Ser
35 40 45

tcc aca aga ctt gca tcg agc tgg cag aca cgt cca ggg cac cat gtt 192
Ser Thr Arg Leu Ala Ser Ser Trp Gln Thr Arg Pro Gly His His Val
50 55 60

ctt ttt tgc ttc tac acc tac ttg tct aca cca ctt cca cac cga cag 240
Leu Phe Cys Phe Tyr Thr Tyr Leu Ser Thr Pro Leu Pro His Arg Gln
65 70 75 80

ctg agt tgt acg att cat ggc agt ccg cgg cat ggc agt ttc gtg cag 288
Leu Ser Cys Thr Ile His Gly Ser Pro Arg His Gly Ser Phe Val Gln
85 90 95

gct gac gtc act g gtgctgaatg ttgacaagca cggctctccac tcacatctag 341
Ala Asp Val Thr
100

accattcaca acgcgcgggt ctcattcttg atagccag ca cag ccg aag gag tca 396
Ala Gln Pro Lys Glu Ser
105

aca tac aaa cct gct cag cca tgg acc tat aca ggg tat tca ccc gct 444
Thr Tyr Lys Pro Ala Gln Pro Trp Thr Tyr Thr Gly Tyr Ser Pro Ala
110 115 120

atg agc cgt a gtgggtataa aagtaaagaa catgagcgac aagtatggct 494
Met Ser Arg
125

ctgtcgaatt ttccggtggg cttgaaaact tcgcttgaac attgatctcc ttgggtccag 553
ct gaa cca gta cgc tgc cta cca atc atc ggg aaa tcg tac aaa aca 600
Thr Glu Pro Val Arg Cys Leu Pro Ile Ile Gly Lys Ser Tyr Lys Thr
130 135 140

gca cac ctc aag tct aaa ctg acc agc gca tcc ctg acg atg g 643

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Ala His Leu Lys Ser Lys Leu Thr	Ser Ala Ser Leu Thr Met	
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caccactgaa cag aa cag aag cca acg cct gag aat agc acg ctg ttc cag		754
	Glu Gln Lys Pro Thr Pro Glu Asn Ser Thr Leu Phe Gln	
	160	165
ggc ttc gag tgg aat gtc ccg gat gat cac aaa cac tgg aag cgt ctt		802
Gly Phe Glu Trp Asn Val Pro Asp Asp His Lys His Trp Lys Arg Leu		
170	175	180
gct gaa cag ctc cca aag ctc aaa gcg atc ggt atc act aac att tgg		850
Ala Glu Gln Leu Pro Lys Leu Lys Ala Ile Gly Ile Thr Asn Ile Trp		
185	190	195
cta cct ccg ggt tgt aaa gcc gcg aat cca aag ggt gtg ggc tat gac		898
Leu Pro Pro Gly Cys Lys Ala Ala Asn Pro Lys Gly Val Gly Tyr Asp		
	205	210
atc tat gac cta tac gac ctg gcc gaa ttt gat cag aaa ggc gcc aaa		946
Ile Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ala Lys		
	220	225
ggc acg aaa tgg gga aca aag gaa gag cta ttg gag ctc agc aag gtt		994
Gly Thr Lys Trp Gly Thr Lys Glu Glu Leu Leu Glu Leu Ser Lys Val		
	235	240
gcc aag gaa cac aac gtt gcc ctt tac tgg gat gcc gtg ttg aat cac		1042
Ala Lys Glu His Asn Val Gly Leu Tyr Trp Asp Ala Val Leu Asn His		
	250	255
aag gcg ggt gca gat agg aca gag aag tgc cgg gtt gtg gaa gtg gat		1090
Lys Ala Gly Ala Asp Arg Thr Glu Lys Cys Arg Val Val Glu Val Asp		
	265	270
gaa aac gac cgc aca aag gag ata tcc gat gcc tac gaa atc gag ggg		1138
Glu Asn Asp Arg Thr Lys Glu Ile Ser Asp Ala Tyr Glu Ile Glu Gly		
	285	290
tgg cta gga ttc gac ttc cca ggg cgc ggc gag cag tat tcc aag atg		1186
Trp Leu Gly Phe Asp Phe Pro Gly Arg Gly Glu Gln Tyr Ser Lys Met		
	300	305
aag tac cac tgg gag cac ttt tct gga acc gac tac aac cag gcc aac		1234
Lys Tyr His Trp Glu His Phe Ser Gly Thr Asp Tyr Asn Gln Ala Asn		
	315	320
gag aag aag gcc atc tac aag atc ctg ggt gaa aac aag ggc tgg tcg		1282
Glu Lys Lys Ala Ile Tyr Lys Ile Leu Gly Glu Asn Lys Gly Trp Ser		
	330	335
caa agc gtg gac acg gaa tcc ggc aac gct gat tac atg atg ttc gcc		1330
Gln Ser Val Asp Thr Glu Ser Gly Asn Ala Asp Tyr Met Met Phe Ala		
	345	350
gac att gac tac tcg cac cct gaa gtt caa gcg gat gtc aag aat tgg		1378
Asp Ile Asp Tyr Ser His Pro Glu Val Gln Ala Asp Val Lys Asn Trp		
	365	370
ggt gtc tgg ata acc aaa gaa gtg gga ctc aag gga ttc cgc ctc gat		1426
Gly Val Trp Ile Thr Lys Glu Val Gly Leu Lys Gly Phe Arg Leu Asp		
	380	385
gct gtg cag cac ttc agc tca aga ttt agc aac gaa tgg atc aag aac		1474
Ala Val Gln His Phe Ser Ser Arg Phe Ser Asn Glu Trp Ile Lys Asn		
	395	400

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ctc Leu	cgc Arg 410	gag Glu	cag Gln	tgc Cys	gga Gly	gac Asp 415	gac Asp	atg Met	ttc Phe	atc Ile	gtg Val 420	ggc Gly	gag Glu	ttc Phe	tgg Trp	1522
tcg Ser 425	ggc Gly	gat Asp	gtc Val	aag Lys	gaa Glu 430	atg Met	agt Ser	gaa Glu	tgg Trp	ctc Leu 435	gac Asp	ggc Gly	atg Met	cag Gln	cac His 440	1570
gaa Glu	att Ile	tgc Cys	ctg Leu	tac Tyr 445	gac Asp	tcg Ser	ccc Pro	ttg Leu	ctt Leu 450	aac Asn	aag Lys	ttt Phe	ggc Gly	tcg Ser 455	ttg Leu	1618
tcg Ser	acg Thr	tgc Cys	gaa Glu 460	tcg Ser	gcc Ala	gat Asp	ctg Leu	agg Arg 465	aca Thr	gtg Val	ttt Phe	gac Asp 470	gac Asp	tcg Ser	ctg Leu	1666
gtc Val	cgg Arg	ctg Leu 475	cgg Arg	ccg Pro	atg Met	gat Asp	gct Ala 480	gtg Val	acg Thr	gtt Val	gtc Val	act Thr 485	aac Asn	cac His	gac Asp	1714
acg Thr	cag Gln 490	cct Pro	gga Gly	cag Gln	gtc Val	atg Met 495	gag Glu	acc Thr	aag Lys	att Ile	gag Glu 500	ggc Gly	ttc Phe	ttt Phe	aag Lys	1762
ccg Pro 505	ctt Leu	gca Ala	tat Tyr	gca Ala	ctg Leu 510	att Ile	ctg Leu	ctc Leu	cga Arg	cag Gln 515	gaa Glu	gga Gly	tac Tyr	cca Pro	tgc Cys 520	1810
ccg Pro	ttc Phe	tat Tyr	ggc Gly	gat Asp 525	ctg Leu	tat Tyr	ggc Gly	ctg Leu	tcg Ser 530	gag Glu	ccg Pro	cac His	gaa Glu	acg Thr 535	ccg Pro	1858
gcg Ala	tcg Ser	tgc Cys	ggt Gly 540	ggc Gly	aag Lys	ctt Leu	gca Ala	gac Asp 545	ctt Leu	atc Ile	ctg Leu	gcg Ala	cgt Arg 550	aag Lys	ctg Leu	1906
ttt Phe	gca Ala	tac Tyr 555	gga Gly	gcg Ala	cag Gln	gaa Glu	gac Asp 560	tac Tyr	ttg Leu	gac Asp	gag Glu	gcc Ala 565	aat Asn	tgc Cys	att Ile	1954
ggg Gly	ttc Phe 570	gtg Val	cgt Arg	cgt Arg	ggg Gly	aca Thr 575	tgg Trp	gac Asp	aag Lys	gtg Val	gcg Ala 580	ggt Gly	ctt Leu	gca Ala	tgt Cys	2002
gtg Val 585	atg Met	agc Ser	aat Asn	gcg Ala	ggg Gly 590	cca Pro	ggg Gly	cag Gln	atc Ile	aga Arg 595	atg Met	gct Ala	gtg Val	ggc Gly	gag Glu 600	2050
atg Met	cac His	gcc Ala	ggg Gly	gag Glu 605	aag Lys	tgg Trp	agc Ser	gac Asp	gtg Val 610	ctt Leu	gga Gly	tgg Trp	gag Glu	cag Gln 615	ggc Gly	2098
gag Glu	gta Val	gag Glu	att Ile 620	gac Asp	agc Ser	gaa Glu	ggc Gly	tac Tyr 625	gga Gly	gtg Val	ttc Phe	agg Arg	tgt Cys 630	cct Pro	ggc Gly	2146
acc Thr	agc Ser	gtc Val 635	gcg Ala	gtg Val	tgg Trp	gtg Val	cga Arg 640	agt Ser	gat Asp	gct Ala	gag Glu	gga Gly 645	cgc Arg	gac Asp	cag Gln	2194
ttc Phe 650	ccc Pro	acc Thr	aac Asn	tgg Trp	gat Asp	gcg Ala 655	gat Asp	gtg Val	tat Tyr	ggc Gly	cag Gln 660	gga Gly	tc Ser	gtgtgcgtaa		2245
tggttcgcat	ggaggggtgta	cggatgggggt	agttgaatca	cggcgctggc	ggccgatgac											2305
gcaag g	ccg	ccc	acc	ctc	ttc	act	tgt	cca	agt	cag	gcg	tgt	act	ggt		2353
	Pro	Pro	Thr	Leu	Phe	Thr	Cys	Pro	Ser	Gln	Ala	Cys	Thr	Val		
				665				670					675			

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gtg	agt	agc	gat	ggg	cat	tat	atc	act	gga	aaa	aga	aga	cat	ggc	gtg	2401
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			680					685					690			
atg	cgc	tgt	cgt	cgc	cct	ctg	ccc	tgt	cgc	cgc	ggc	tcg	cag	gct	gtt	2449
Met	Arg	Cys	Arg	Arg	Pro	Leu	Pro	Cys	Arg	Arg	Gly	Ser	Gln	Ala	Val	
		695					700					705				
agc	tgg	tgg	aag	atg	cgc	gaa	agt	gcg	tgg	ggc	ggc	cga	cag	ggc	tgc	2497
Ser	Trp	Trp	Lys	Met	Arg	Glu	Ser	Ala	Trp	Gly	Gly	Arg	Gln	Gly	Cys	
	710					715					720					
cgc	gca	gca	cag	cca	atc	acc	cgc	cgc	aca	aca	ata	gcc	gtt	ttg	gac	2545
Arg	Ala	Ala	Gln	Pro	Ile	Thr	Arg	Arg	Thr	Thr	Ile	Ala	Val	Leu	Asp	
	725				730					735					740	
gtt	tcg	cgg	cct	ttc	tgg	caa	gcg	agt	atg	cga	cat	ccc	gcg	aag	cat	2593
Val	Ser	Arg	Pro	Phe	Trp	Gln	Ala	Ser	Met	Arg	His	Pro	Ala	Lys	His	
				745				750						755		
gat	cga	cgg	ccc	atg	ccg	gca	gaa	tgc	tcc	ctc	cct	ctg	tac	cat	gat	2641
Asp	Arg	Arg	Pro	Met	Pro	Ala	Glu	Cys	Ser	Leu	Pro	Leu	Tyr	His	Asp	
			760					765					770			
gca	cac	gca	cac	gca	cat	gct	cgc	acg	cgc	gcc	cgc	gac	acg	gcc	aac	2689
Ala	His	Ala	His	Ala	His	Ala	Arg	Thr	Arg	Ala	Arg	Asp	Thr	Ala	Asn	
		775					780					785				
atg	cga	ggc	caa	cgg	ccc	cgt	cct	gcc	gcc	gga	cat	gaa	gcc	gca	tgt	2737
Met	Arg	Gly	Gln	Arg	Pro	Arg	Pro	Ala	Ala	Gly	His	Glu	Ala	Ala	Cys	
	790					795					800					
gcc	atg	tga														2746
Ala	Met	*														
805																

<210> 468
 <211> 100
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

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 20 25 30
 Pro Pro Ser Gly Cys Pro Ser Ser Asp Asp Leu Ala Pro Lys Ser Ser
 35 40 45
 Ser Thr Arg Leu Ala Ser Ser Trp Gln Thr Arg Pro Gly His His Val
 50 55 60
 Leu Phe Cys Phe Tyr Thr Tyr Leu Ser Thr Pro Leu Pro His Arg Gln
 65 70 75 80
 Leu Ser Cys Thr Ile His Gly Ser Pro Arg His Gly Ser Phe Val Gln
 85 90 95
 Ala Asp Val Thr
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<210> 469
 <211> 25
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

<400> 469
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<211> 30
<212> PRT
<213> Cochliobolus heterostrophus ATCC 48331

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Ala His Leu Lys Ser Lys Leu Thr Ser Ala Ser Leu Thr Met
20 25 30

<210> 471
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<212> PRT
<213> Cochliobolus heterostrophus ATCC 48331

<400> 471
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20 25 30
Leu Pro Lys Leu Lys Ala Ile Gly Ile Thr Asn Ile Trp Leu Pro Pro
35 40 45
Gly Cys Lys Ala Ala Asn Pro Lys Gly Val Gly Tyr Asp Ile Tyr Asp
50 55 60
Leu Tyr Asp Leu Gly Glu Phe Asp Gln Lys Gly Ala Lys Gly Thr Lys
65 70 75 80
Trp Gly Thr Lys Glu Glu Leu Leu Glu Leu Ser Lys Val Ala Lys Glu
85 90 95
His Asn Val Gly Leu Tyr Trp Asp Ala Val Leu Asn His Lys Ala Gly
100 105 110
Ala Asp Arg Thr Glu Lys Cys Arg Val Val Glu Val Asp Glu Asn Asp
115 120 125
Arg Thr Lys Glu Ile Ser Asp Ala Tyr Glu Ile Glu Gly Trp Leu Gly
130 135 140
Phe Asp Phe Pro Gly Arg Gly Glu Gln Tyr Ser Lys Met Lys Tyr His
145 150 155 160
Trp Glu His Phe Ser Gly Thr Asp Tyr Asn Gln Ala Asn Glu Lys Lys
165 170 175
Ala Ile Tyr Lys Ile Leu Gly Glu Asn Lys Gly Trp Ser Gln Ser Val
180 185 190
Asp Thr Glu Ser Gly Asn Ala Asp Tyr Met Met Phe Ala Asp Ile Asp
195 200 205
Tyr Ser His Pro Glu Val Gln Ala Asp Val Lys Asn Trp Gly Val Trp
210 215 220
Ile Thr Lys Glu Val Gly Leu Lys Gly Phe Arg Leu Asp Ala Val Gln
225 230 235 240
His Phe Ser Ser Arg Phe Ser Asn Glu Trp Ile Lys Asn Leu Arg Glu
245 250 255
Gln Cys Gly Asp Asp Met Phe Ile Val Gly Glu Phe Trp Ser Gly Asp
260 265 270
Val Lys Glu Met Ser Glu Trp Leu Asp Gly Met Gln His Glu Ile Cys
275 280 285
Leu Tyr Asp Ser Pro Leu Leu Asn Lys Phe Gly Ser Leu Ser Thr Cys
290 295 300
Glu Ser Ala Asp Leu Arg Thr Val Phe Asp Asp Ser Leu Val Arg Leu
305 310 315 320
Arg Pro Met Asp Ala Val Thr Val Val Thr Asn His Asp Thr Gln Pro
325 330 335
Gly Gln Val Met Glu Thr Lys Ile Glu Gly Phe Phe Lys Pro Leu Ala
340 345 350
Tyr Ala Leu Ile Leu Leu Arg Gln Glu Gly Tyr Pro Cys Pro Phe Tyr
355 360 365
Gly Asp Leu Tyr Gly Leu Ser Glu Pro His Glu Thr Pro Ala Ser Cys
370 375 380
Gly Gly Lys Leu Ala Asp Leu Ile Leu Ala Arg Lys Leu Phe Ala Tyr

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385      390      395      400
Gly Ala Gln Glu Asp Tyr Leu Asp Glu Ala Asn Cys Ile Gly Phe Val
405      410      415
Arg Arg Gly Thr Trp Asp Lys Val Ala Gly Leu Ala Cys Val Met Ser
420      425      430
Asn Ala Gly Pro Gly Gln Ile Arg Met Ala Val Gly Glu Met His Ala
435      440      445
Gly Glu Lys Trp Ser Asp Val Leu Gly Trp Glu Gln Gly Glu Val Glu
450      455      460
Ile Asp Ser Glu Gly Tyr Gly Val Phe Arg Cys Pro Gly Thr Ser Val
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Ala Val Trp Val Arg Ser Asp Ala Glu Gly Arg Asp Gln Phe Pro Thr
485      490      495
Asn Trp Asp Ala Asp Val Tyr Gly Gln Gly Ser
500      505

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<210> 472
 <211> 144
 <212> PRT
 <213> *Cochliobolus heterostrophus* ATCC 48331

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20      25      30
Cys Arg Arg Pro Leu Pro Cys Arg Arg Gly Ser Gln Ala Val Ser Trp
35      40      45
Trp Lys Met Arg Glu Ser Ala Trp Gly Gly Arg Gln Gly Cys Arg Ala
50      55      60
Ala Gln Pro Ile Thr Arg Arg Thr Thr Ile Ala Val Leu Asp Val Ser
65      70      75      80
Arg Pro Phe Trp Gln Ala Ser Met Arg His Pro Ala Lys His Asp Arg
85      90      95
Arg Pro Met Pro Ala Glu Cys Ser Leu Pro Leu Tyr His Asp Ala His
100      105      110
Ala His Ala His Ala Arg Thr Arg Ala Arg Asp Thr Ala Asn Met Arg
115      120      125
Gly Gln Arg Pro Arg Pro Ala Ala Gly His Glu Ala Ala Cys Ala Met
130      135      140

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<210> 473
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 <212> DNA
 <213> *Cochliobolus heterostrophus* ATCC 48331

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gacgatcttg ccccaaagtc ttcattccaca agacttgcac cgagctggca gacacgtcca      180
gggcaccatg ttcttttttg cttctacacc tacttgctca caccacttcc acaccgacag      240
ctgagttgta cgattcatgg cagtccgcgg catggcagtt tcgtgcaggc tgacgtcact      300
gcacagccga aggagtcaac atacaaacct gctcagccat ggacctatac aggggtattca      360
cccgtatga gccgtactga accagtacgc tgcctacca tcatcgggaa atcgtacaaa      420
acagcacacc tcaagtctaa actgaccagc gcatccctga cgatggaaca gaagccaacg      480
cctgagaata gcacgctggt ccagggtctc gagtggaaatg tcccggatga tcacaaacac      540
tggaagcgct ttgctgaaca gctcccaaag ctcaaagcga tcggtatcac taacatttgg      600
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tacgacctgg gcgaatttga tcagaaaggc gccaaaggca cgaaatgggg aacaaaggaa      720
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 <223> Alpha amylase, catalytic domain

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 35 40 45
 Ser Thr Arg Leu Ala Ser Ser Trp Gln Thr Arg Pro Gly His His Val
 50 55 60
 Leu Phe Cys Phe Tyr Thr Tyr Leu Ser Thr Pro Leu Pro His Arg Gln
 65 70 75 80
 Leu Ser Cys Thr Ile His Gly Ser Pro Arg His Gly Ser Phe Val Gln
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 Ala Asp Val Thr Ala Gln Pro Lys Glu Ser Thr Tyr Lys Pro Ala Gln
 100 105 110
 Pro Trp Thr Tyr Thr Gly Tyr Ser Pro Ala Met Ser Arg Thr Glu Pro
 115 120 125
 Val Arg Cys Leu Pro Ile Ile Gly Lys Ser Tyr Lys Thr Ala His Leu
 130 135 140
 Lys Ser Lys Leu Thr Ser Ala Ser Leu Thr Met Glu Gln Lys Pro Thr
 145 150 155 160
 Pro Glu Asn Ser Thr Leu Phe Gln Gly Phe Glu Trp Asn Val Pro Asp
 165 170 175
 Asp His Lys His Trp Lys Arg Leu Ala Glu Gln Leu Pro Lys Leu Lys
 180 185 190
 Ala Ile Gly Ile Thr Asn Ile Trp Leu Pro Pro Gly Cys Lys Ala Ala
 195 200 205
 Asn Pro Lys Gly Val Gly Tyr Asp Ile Tyr Asp Leu Tyr Asp Leu Gly
 210 215 220
 Glu Phe Asp Gln Lys Gly Ala Lys Gly Thr Lys Trp Gly Thr Lys Glu
 225 230 235 240
 Glu Leu Leu Glu Leu Ser Lys Val Ala Lys Glu His Asn Val Gly Leu
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 Tyr Trp Asp Ala Val Leu Asn His Lys Ala Gly Ala Asp Arg Thr Glu
 260 265 270
 Lys Cys Arg Val Val Glu Val Asp Glu Asn Asp Arg Thr Lys Glu Ile
 275 280 285

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Arg	290	Gly	Glu	Gln	Tyr	Ser	295	Met	Lys	Tyr	His	300	Glu	His	Phe
Gly	305	Thr	Asp	Tyr	Asn	Gln	Ala	Asn	Glu	Lys	Lys	Ala	Ile	Tyr	Lys
Leu		Gly	Glu	Asn	Lys	Gly	Trp	Ser	Gln	Ser	Val	Asp	Thr	Glu	Ser
Asn		Ala	Asp	Tyr	Met	Met	Phe	Ala	Asp	Ile	Asp	Tyr	Ser	His	Pro
Val		Gln	Ala	Asp	Val	Lys	Asn	Trp	Gly	Val	Trp	Ile	Thr	Lys	Glu
Gly		Leu	Lys	Gly	Phe	Arg	Leu	Asp	Ala	Val	Gln	His	Phe	Ser	Ser
Phe		Ser	Asn	Glu	Trp	Ile	Lys	Asn	Leu	Arg	Glu	Gln	Cys	Gly	Asp
Met		Phe	Ile	Val	Gly	Glu	Phe	Trp	Ser	Gly	Asp	Val	Lys	Glu	Met
Glu		Trp	Leu	Asp	Gly	Met	Gln	His	Glu	Ile	Cys	Leu	Tyr	Asp	Ser
Leu		Leu	Asn	Lys	Phe	Gly	Ser	Leu	Ser	Thr	Cys	Glu	Ser	Ala	Asp
Arg		Thr	Val	Phe	Asp	Asp	Ser	Leu	Val	Arg	Leu	Arg	Pro	Met	Asp
Val		Thr	Val	Val	Thr	Asn	His	Asp	Thr	Gln	Pro	Gly	Gln	Val	Met
Thr		Lys	Ile	Glu	Gly	Phe	Phe	Lys	Pro	Leu	Ala	Tyr	Ala	Leu	Ile
Leu		Arg	Gln	Glu	Gly	Tyr	Pro	Cys	Pro	Phe	Tyr	Gly	Asp	Leu	Tyr
Leu		Ser	Glu	Pro	His	Glu	Thr	Pro	Ala	Ser	Cys	Gly	Gly	Lys	Leu
Asp		Leu	Ile	Leu	Ala	Arg	Lys	Leu	Phe	Ala	Tyr	Gly	Ala	Gln	Glu
Tyr		Leu	Asp	Glu	Ala	Asn	Cys	Ile	Gly	Phe	Val	Arg	Arg	Gly	Thr
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Gln		Ile	Arg	Met	Ala	Val	Gly	Glu	Met	His	Ala	Gly	Glu	Lys	Trp
Asp		Val	Leu	Gly	Trp	Glu	Gln	Gly	Glu	Val	Glu	Ile	Asp	Ser	Glu
Tyr		Gly	Val	Phe	Arg	Cys	Pro	Gly	Thr	Ser	Val	Ala	Val	Trp	Val
Ser		Asp	Ala	Glu	Gly	Arg	Asp	Gln	Phe	Pro	Thr	Asn	Trp	Asp	Ala
Val		Tyr	Gly	Gln	Gly	Ser	Pro	Pro	Thr	Leu	Phe	Thr	Cys	Pro	Ser
Ala		Cys	Thr	Val	Val	Ser	Ser	Asp	Gly	His	Tyr	Ile	Thr	Gly	Lys
Arg		His	Gly	Val	Met	Arg	Cys	Arg	Arg	Pro	Leu	Pro	Cys	Arg	Arg
Ser		Gln	Ala	Val	Ser	Trp	Trp	Lys	Met	Arg	Glu	Ser	Ala	Trp	Gly
Arg		Gln	Gly	Cys	Arg	Ala	Ala	Gln	Pro	Ile	Thr	Arg	Arg	Thr	Thr
Ala		Val	Leu	Asp	Val	Ser	Arg	Pro	Phe	Trp	Gln	Ala	Ser	Met	Arg
Pro		Ala	Lys	His	Asp	Arg	Arg	Pro	Met	Pro	Ala	Glu	Cys	Ser	Leu
Leu		Tyr	His	Asp	Ala	His	Ala	His	Ala	His	Ala	Arg	Thr	Arg	Ala
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 Ser Leu Asp Tyr Ile Gln Gly Met Gly Ile Lys Gly Ile Tyr Ile Ala
 20 25 30
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 Gly Ser Pro Met Ile Asn Ala Pro Trp Gly Ala Asp Gln Tyr Ser Pro
 35 40 45
 ctg gat ttc agt ctt ctc gac caa cat ttc gga gat atc gat ttg tgg 192
 Leu Asp Phe Ser Leu Leu Asp Gln His Phe Gly Asp Ile Asp Leu Trp
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 agg aaa act acc act gcc atc cat gac agg gga atg tat gtg atg ctc 240
 Arg Lys Thr Thr Thr Ala Ile His Asp Arg Gly Met Tyr Val Met Leu
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 Asp Asn Thr Met Gly Thr Met Gly Asp Leu Ile Ala Phe Lys Gly Tyr
 85 90 95
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 Glu Asn Ser Ser Thr Pro Phe Thr Leu Thr Glu His Glu Val Leu Trp
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 Arg Asn Asp Asn Arg Gln Tyr Leu Asp Phe Ser Phe Gly Ser Lys Tyr
 115 120 125
 aat gag acc tgc aac tac cct aga ttc tgg ctt gat act gga tat acc 432
 Asn Glu Thr Cys Asn Tyr Pro Arg Phe Trp Leu Asp Thr Gly Tyr Thr
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 Phe Asp Gln Tyr Gly Asp Thr Glu Ala Phe Gly Val Phe Pro Asp Trp
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 Gln Arg Gln Leu Ser Lys Phe Ala Ser Val Gln Asp Arg Leu Arg Glu
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 Trp Val Pro Ser Val Arg Thr Lys Ile Glu Ile Phe Thr Cys Met Ile
 195 200 205
 atc caa caa ttg gac att gat gga ctg cgt gtg gac aag gcg aca caa 672
 Ile Gln Gln Leu Asp Ile Asp Gly Leu Arg Val Asp Lys Ala Thr Gln

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agt Ser	ttc Phe	agt Ser	cac His 235
gca Ala	atc Ile	aga Arg	gag Glu
tgc Cys 240			
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ggt Gly	gga Gly	aac Asn	act Thr 260
ttt Phe	gga Gly	agt Ser	att Ile
tac Tyr 265	ctc Leu	ggg Gly	cgt Arg
gga Gly	cgt Arg	gga Gly	cgt Arg 270
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tat Tyr	ttt Phe	atc Ile 295	cgt Arg
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cac His	tat Tyr 310	agt Ser	ggt Val
tat Tyr	cgc Arg	aca Thr 315	ctg Leu
acc Thr	aga Arg	ttt Phe	ctg Leu 320
960			
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aac Asn 325	ttg Leu	gca Ala	agt Ser
ggt Gly	tat Tyr 330	gat Asp	aca Thr
cca Pro	act Thr	aac Asn 335	tgg Trp
1008			
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aac Asn	gaa Glu	atg Met	ctt Leu
gta Val 345	acg Thr	aac Asn	gat Asp
ttt Phe 350	ggt Val	aac Asn	cct Pro
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ttc Phe	gat Asp	cca Pro	aga Arg 360
cat His	atg Met	tac Tyr	gga Gly 365
gtc Val	agc Ser	aat Asn	cag Gln
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cag Gln	aat Asn	ggc Gly	aca Thr 380
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cac His	atg Met	cct Pro 395	ggg Gly
att Ile	cct Pro	ggt Gly	att Ile
cct Leu 400			
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gaa Glu	caa Gln	gca Ala	ttc Phe
tat Tyr 410	gtc Val	tta Leu	gac Asp
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caa Gln	gca Ala 430	tgg Trp	cag Gln
1296			
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tac Tyr	gcc Ala	ctt Leu	ggc Gly 440
agt Ser	ggt Gly	aca Thr	tat Tyr 445
caa Gln	ttc Phe	ccc Pro	
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act Thr	cga Arg	ggt Gly 455	tgc Cys
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aac Asn	agc Ser	cgc Arg	gat Asp
1392			
cac His 465	aga Arg	gat Asp	cca Pro
tct Ser	cat His 470	cca Pro	ggt Val
aga Arg	aat Asn	att Ile 475	gtc Val
aag Lys	gca Ala	atg Met	tac Tyr 480
1440			
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tat Tyr	cca Pro	ggt Val	ttg Leu
acc Thr	gat Asp	gga Gly	gca Ala
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Gly	Ile	Gln	Asp	Leu	Ser	Lys	Ala	Gly	Gly	Gln	Ala	Asn	Gln	Ser	Val	
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Trp	Met	Val	Tyr	His	Asn	Asn	Tyr	Asp	Thr	Val	Thr	Tyr	Ser	Phe	Asp	
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Cys	Ser	Ser	Asn	Glu	Thr	Ser	Leu	Val	Ala	Pro	Phe	Leu	Asp	Gly	Val	
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Ser	Ala	Val	Asn	Val	Thr	Glu	Phe	Val	Gly	Ser	Ile	Pro	Thr	Met	Trp	
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Thr	Phe	Lys	Gly	Asn	Leu	Thr	Asn	Val	Ser	Asn	Gly	Val	His	Ser	Ile	
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Asn Gly Thr Leu Tyr Val Ser His Lys Ala Ala Gly Ala Asp Arg Phe			
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aaa tat tct ttg aac tgg gga agt tca tgg tct gaa tgg tta cca tat			2448
Lys Tyr Ser Leu Asn Trp Gly Ser Ser Trp Trp Ser Glu Trp Leu Pro Tyr			
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Val Gly Gly Asn Thr Thr Leu Glu Pro Pro Gln Ala Trp Thr Gly Thr Ser			
820	825	830	
aaa caa aaa tgg gac gga gat cac gtc atc atg caa tat tgg agc aaa			2544
Lys Gln Lys Trp Asp Gly Asp His Val Ile Met Gln Tyr Trp Ser Lys			
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ctt act ggt agc agt gat gta gtt caa cat gct gat cta gaa acc aac			2592
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850	855	860	
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865	870	875	
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Gln Tyr Gly Tyr Asp Thr Gly Ile Asp Asn Gly Phe Thr Leu Asn Lys			
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			Val Met Gln
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Val Asn Val Trp Gly Met Asn Pro Asp Gly Gln Pro Asp Glu Thr Ile			
915	920	925	
gtg ttg gga gat atc gac aat gat aat atc ctt gat cgt atg cca cct			2888
Val Leu Gly Asp Ile Asp Asn Asp Asn Ile Leu Asp Arg Met Pro Pro			
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Ser Ser Leu Ser Thr Ala Leu Ile Asn Ile Thr Arg Val Pro Pro Ser			
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cca tat ctc gga tac aaa att gca atc gac gac ggt aat tat aga tac			2984
Pro Tyr Leu Gly Tyr Lys Ile Ala Ile Asp Asp Gly Asn Tyr Arg Tyr			
965	970	975	
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Tyr Gln Ile Pro Tyr Gly Asn Arg Ser Tyr Gln Met Leu Met Val Val			
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Phe Met Lys Ser Phe Tyr Ala Val Lys Leu Asn Ser Val Gly Val Lys			

1001827087_1.txt																		
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Glu	Ala	Lys	Asn	Tyr	Ile	Pro	Leu	Ala	Leu	Arg	Arg	Lys	Leu	Lys	Arg			
			1030					1035					1040					
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Glu	Lys	Lys	Glu	Asp	Pro	Ser	Glu	Lys	Glu	Met	Thr	Val	Met	Gly	Gly			
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Asp	Val	Ala	Ala	Thr	Gly	Arg	Arg	Thr	Val	Leu	Ile	Ala	Thr	Met	Glu			
					1080					1085					1090			
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Tyr	Asp	Ile	Glu	Asp	Trp	Ala	Ile	Lys	Ile	Lys	Ile	Gly	Gly	Leu	Gly			
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Val	Met	Ala	Gln	Leu	Met	Gly	Lys	Asn	Leu	Gly	His	Gln	Asp	Leu	Ile			
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Trp	Val	Val	Pro	Cys	Val	Gly	Gly	Ile	Asp	Tyr	Pro	Ile	Asp	His	Gln			
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Pro	Val	Phe	Arg	Ala	Gln	Ser	Lys	Ser	Glu	Pro	Tyr	Pro	Ala	Arg	Met			
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Asp	Asp	Leu	Asp	Ser	Ala	Val	Tyr	Tyr	Ser	Ala	Trp	Asn	Gln	Cys	Ile			
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Arg	Asn	Gln	Lys	Glu	Arg	Ser	Glu	Val	Cys	Gln	Val	Tyr	Asn	Ile	Pro			
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cct att ttc tgg gga ttg agc aag atc ggc aat ctt cct aat cct gat			4040
Pro Ile Phe Trp Gly Leu Ser Lys Ile Gly Asn Leu Pro Asn Pro Asp			
1315	1320	1325	
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Thr Ile Val Asn Gln Asp Phe Glu Ser Gln Arg Ala Asp Leu Lys Arg			
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Gln Ala Gln Glu Trp Ala Gly Leu Glu Gln Asn Pro Lys Ala Glu Leu			
1365	1370	1375	
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Leu Val Phe Val Gly Arg Trp Ser Met Gln Lys Gly Ile Asp Leu Ile			
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Ala Asp Val Met Pro Ala Val Leu Glu Glu Asn Pro Asn Val Gln Leu			
1395	1400	1405	
att tgc gtt ggt cct gtt att gat ctt tat gga aaa ttc gcc gct ctc			4328
Ile Cys Val Gly Pro Val Ile Asp Leu Tyr Gly Lys Phe Ala Ala Leu			
1415	1420	1425	
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Lys Leu Asp Val Met Met Thr Lys Tyr Lys Gly Arg Val Phe Ser Lys			
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Pro Glu Phe Thr Ala Leu Pro Pro Phe Ile Phe Ser Gly Ala Glu Phe			
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Ala Leu Ile Pro Ser Arg Asp Glu Pro Phe Gly Leu Val Ala Val Glu			
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Phe Gly Arg Lys Gly Ala Leu Gly Ile Gly Ala Arg Val Gly Gly Leu			
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Gly Gln Met Pro Gly Trp Trp Phe Thr Val Glu Ser Thr Thr Thr Thr			
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His Met Leu His Gln Phe Lys Gln Ala Ile Lys Glu Ala Leu Ala Ser			
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Lys Leu Glu Thr Arg Gln Ile Met Arg Ala Arg Ser Ala Lys Gln Arg			
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Phe Pro Val Ala Gln Trp Val Asn Asp Leu Glu Ile Leu Gln Ser Lys			
1540	1545	1550	
gca atc aag att cac gat aag gaa gag gcc aaa cat tct ggt aga cca			4760
Ala Ile Lys Ile His Asp Lys Glu Glu Ala Lys His Ser Gly Arg Pro			

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1555	aga tct aga gga aga gat gga gga aat att ttc tat tct tcc gct aag	4808
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	1575 1580 1585	
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Arg Leu Ser Ser Phe Gly Thr Ser Phe Ala Ser Thr Ser Ser Gln Leu Asp		
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Val Ala Ser Thr Ile Gly Gly Ser Thr Arg Pro Ser Ser Pro Thr Arg		
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	1670 1675 1680	
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Ile Tyr Asp Glu Tyr Val Leu Thr Pro Glu Glu Leu Glu Ala Gln Arg		
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	1715 1720 1725 1730	
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Leu Pro Arg Leu Lys Asp Gly Ser Asn Ile Tyr Ser Thr His Ser Ile		
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Pro Gly Ser Pro Gly Leu Glu Ser Pro Gly Thr Pro Gly Ala His Asp		
	1765 1770 1775	
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Ser Leu Leu Pro Pro Pro Ser Ile Phe Gly Ala Gly Gly Asp Asn Ala		
	1780 1785 1790	
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Asn Arg Ala Ser Met Leu Ser Leu Ser Ser Val Ile Gly Asp Lys Thr		
	1795 1800 1805 1810	
gat ttc aaa ctt caa aag gtc gat cca ttc ttc act gat act caa gga	5528	
Asp Phe Lys Leu Gln Lys Val Asp Pro Phe Phe Thr Asp Thr Gln Gly		
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Lys	Lys	Trp	Phe	Asp	Arg	Phe	Arg	Asp	Ala	Lys	Leu	Gly	Arg	Asn	Ser									
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Ser	Arg	Gly	Arg	Ser	Asn	Ser	Arg	Asp	Ser	Phe	Thr	Asn	Asp	Thr	Ile									
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Thr	His	Ser	Arg	Gln	Asn	Ser	Ser	Gln	Asp	Asn	Met	Ala	Leu	Thr	Gly									
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Val	Tyr	Ser	Phe	Leu	Leu	Ala	Phe	Gly	Gln	Ile	Ile	Ala	Ala	Asn	Ser									
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ctt	tac	atc	aca	gca	acg	atc	tat	ctt	ggt	acc	tcg	atc	atg	tgg	tgg	6056								
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Tyr	Ala	Val	Ala	Ala	Ser	Ser	Gly	Ser	Ile	Phe	Phe	Ala	Leu	Asn	Phe									
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Tyr Arg Gln Ala Pro Gly Lys Val Pro Ser Phe Tyr Ala Ser Leu Phe			
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Gly Met Leu Leu Leu Gln Thr Leu Thr Arg Val His Ile Ala Phe Thr			
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2375

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Gly Ser Pro Met Ile Asn Ala Pro Trp Gly Ala Asp Gln Tyr Ser Pro
35 40 45
Leu Asp Phe Ser Leu Leu Asp Gln His Phe Gly Asp Ile Asp Leu Trp
50 55 60
Arg Lys Thr Thr Thr Ala Ile His Asp Arg Gly Met Tyr Val Met Leu
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Asp Asn Thr Met Gly Thr Met Gly Asp Leu Ile Ala Phe Lys Gly Tyr
85 90 95
Glu Asn Ser Ser Thr Pro Phe Thr Leu Thr Glu His Glu Val Leu Trp
100 105 110
Arg Asn Asp Asn Arg Gln Tyr Leu Asp Phe Ser Phe Gly Ser Lys Tyr
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Phe Asp Gln Tyr Gly Asp Thr Glu Ala Phe Gly Val Phe Pro Asp Trp
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Gln Arg Gln Leu Ser Lys Phe Ala Ser Val Gln Asp Arg Leu Arg Glu
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225 230 235 240
Ala Arg Asp Val Asn Lys Thr Asn Phe Met Val Thr Gly Glu Ile Thr
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260 265 270
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Asp Val Phe Arg Trp Pro Ala Ile Gln Asn Gly Thr Gln Lys Gln Leu
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Leu Gly Phe Phe Ile Thr Thr Ile His Met Pro Gly Ile Pro Leu Leu
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405 410 415
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Phe Thr Leu Ile Ala Ala Gln Val Leu Gly Ser Ile Ala Thr Met Val
2305      2310      2315      2320
Ala Arg Ala Cys Ala Pro Asn Lys Ile Gly Pro Gly Thr Val Phe Pro
2325      2330      2335
Asp Phe Ser Asp Gly Ala Ala Gly Leu Gly Asn Gly Trp Phe Trp Ile
2340      2345      2350
Gly Leu Ile Leu Gln Leu Val Ile Cys Gly Gly Phe Phe Thr Phe Phe
2355      2360      2365
Arg Lys Glu Gln Leu Ser Lys Pro
2370      2375

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<210> 480
<211> 1511
<212> DNA
<213> Fungi

<220>
<221> CDS
<222> (1)...(263)
<223> Exon

<221> CDS
<222> (312)...(417)
<223> Exon

<221> CDS
<222> (498)...(1510)
<223> Exon

<400> 480
atg aag ttc tct ctt ctt gcg acc atc gtc gct agc att agt ccg ctt      48
Met Lys Phe Ser Leu Leu Ala Thr Ile Val Ala Ser Ile Ser Pro Leu
  1                               10                      15

gca cgc gca gca gat gca aac gct tgg aag tcg cga aat att tac ttc      96
Ala Arg Ala Ala Asp Ala Asn Ala Trp Lys Ser Arg Asn Ile Tyr Phe
                20                      25                      30

gca ctt act gat cgt gtt gcg cgg agt gct gat gat aat ggc ggc agt      144
Ala Leu Thr Asp Arg Val Ala Arg Ser Ala Asp Asp Asn Gly Gly Ser
                35                      40                      45

gca tgc gga aac ctc ggg aat tac tgt ggt gga act ttt aag ggc tta      192
Ala Cys Gly Asn Leu Gly Asn Tyr Cys Gly Gly Thr Phe Lys Gly Leu
                50                      55                      60

gag tca aag ctt gat tat atc aag ggc atg gga ttt gat gct att tgg      240
Glu Ser Lys Leu Asp Tyr Ile Lys Gly Met Gly Phe Asp Ala Ile Trp
  65                               70                      75                      80

att act cct gtt gtt gac agt ag gtggtgcgaa attgagtga aaaatgtgac      293
Ile Thr Pro Val Val Asp Ser Ser
                85

tgacaggggtg atagatac t gat ggg gga tac cat gga tac tgg gct aag gat      345
                Asp Gly Gly Tyr His Gly Tyr Trp Ala Lys Asp
                90                      95

ctg tat gcg gtc aac tcc aag tat ggt act gca gat gat ttg aag agt      393
Leu Tyr Ala Val Asn Ser Lys Tyr Gly Thr Ala Asp Asp Leu Lys Ser
 100                               105                      110                      115

ctt gtc aaa tct gct cat gac aag gtaagcaaag ttctctctct catgctgttc      447
Leu Val Lys Ser Ala His Asp Lys
                120

ttctgtgatc gccttggtct ttcttggtga tcccgtgctg actgttccag aac atg      503
                Asn Met
                125

tac gtc atg tgc gac gtc gtc gca aac cac atg ggc aaa ggc atc tca      551
Tyr Val Met Cys Asp Val Val Ala Asn His Met Gly Lys Gly Ile Ser
                130                      135                      140

gac cac aaa ccc tcc ccc ctc aac gaa caa agc tca tac cac acc ccc      599
Asp His Lys Pro Ser Pro Leu Asn Glu Gln Ser Ser Tyr His Thr Pro
                145                      150                      155

tgc gat atc gac tac agc aac cag acc agc atc gag cag tgc gaa atc      647

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1001827087_1.txt															
Cys	Asp	Ile	Asp	Tyr	Ser	Asn	Gln	Thr	Ser	Ile	Glu	Gln	Cys	Glu	Ile
		160					165					170			
gcc	ggt	ctt	ccc	gac	ctc	aac	acc	ggc	agc	gac	acc	gtc	aag	aaa	gtc
Ala	Gly	Leu	Pro	Asp	Leu	Asn	Thr	Gly	Ser	Asp	Thr	Val	Lys	Lys	Val
	175					180					185				
ctc	tac	gac	tgg	atc	aag	tgg	ctt	gtc	tcc	gag	tac	agc	ttc	gac	ggc
Leu	Tyr	Asp	Trp	Ile	Lys	Trp	Leu	Val	Ser	Glu	Tyr	Ser	Phe	Asp	Gly
	190				195					200					205
atc	cgc	atc	gat	aca	gtc	aaa	cat	gtc	gaa	aag	ccg	ttc	tgg	cct	ggc
Ile	Arg	Ile	Asp	Thr	Val	Lys	His	Val	Glu	Lys	Pro	Phe	Trp	Pro	Gly
				210					215					220	
ttt	caa	gac	gct	gct	ggc	gtt	tac	gcc	atc	ggc	gag	gtt	tgg	gat	gga
Phe	Gln	Asp	Ala	Ala	Gly	Val	Tyr	Ala	Ile	Gly	Glu	Val	Trp	Asp	Gly
			225					230					235		
ggg	cct	gat	tac	cta	gct	ggc	tat	gca	cag	gtc	atg	cct	ggc	ctc	ttg
Gly	Pro	Asp	Tyr	Leu	Ala	Gly	Tyr	Ala	Gln	Val	Met	Pro	Gly	Leu	Leu
		240					245					250			
aac	tat	gct	atg	tat	tac	ccc	atg	aac	cgg	ttc	tac	cag	caa	aaa	ggc
Asn	Tyr	Ala	Met	Tyr	Tyr	Pro	Met	Asn	Arg	Phe	Tyr	Gln	Gln	Lys	Gly
	255					260					265				
gat	cct	tca	gat	gtt	gtc	gct	atg	cac	gat	gag	att	agc	aac	aaa	ttc
Asp	Pro	Ser	Asp	Val	Val	Ala	Met	His	Asp	Glu	Ile	Ser	Asn	Lys	Phe
	270				275					280					285
ccc	gat	ccc	act	atc	ctc	gga	aca	ttc	atc	gac	aac	cac	gat	aac	cct
Pro	Asp	Pro	Thr	Ile	Leu	Gly	Thr	Phe	Ile	Asp	Asn	His	Asp	Asn	Pro
				290					295					300	
cgc	tgg	ttg	agc	caa	aag	aac	gac	aaa	gct	ctc	ctc	aag	aac	gcc	ctc
Arg	Trp	Leu	Ser	Gln	Lys	Asn	Asp	Lys	Ala	Leu	Leu	Lys	Asn	Ala	Leu
			305					310					315		
gca	tac	gtc	atc	cta	gcc	cga	ggc	att	ccc	atc	gtc	tac	tac	gga	aca
Ala	Tyr	Val	Ile	Leu	Ala	Arg	Gly	Ile	Pro	Ile	Val	Tyr	Tyr	Gly	Thr
		320					325					330			
gag	caa	ggc	tac	gct	ggc	ggc	aat	gat	ccc	gcc	aac	cgt	gag	gat	ctc
Glu	Gln	Gly	Tyr	Ala	Gly	Gly	Asn	Asp	Pro	Ala	Asn	Arg	Glu	Asp	Leu
	335				340						345				
tgg	cga	agc	agc	ttc	agc	acc	gac	gcg	gat	ctg	tat	caa	cac	att	tca
Trp	Arg	Ser	Ser	Phe	Ser	Thr	Asp	Ala	Asp	Leu	Tyr	Gln	His	Ile	Ser
	350				355					360					365
cgc	ctc	tcc	aag	gct	cgc	tcg	gcg	gtc	ggc	ggc	ctt	ggc	gga	aat	gat
Arg	Leu	Ser	Lys	Ala	Arg	Ser	Ala	Val	Gly	Gly	Leu	Gly	Gly	Asn	Asp
				370					375					380	
cac	aag	cat	ctc	tac	tcg	cag	gac	agc	gcg	tat	gct	tgg	agt	cgt	gct
His	Lys	His	Leu	Tyr	Ser	Gln	Asp	Ser	Ala	Tyr	Ala	Trp	Ser	Arg	Ala
			385					390					395		
gat	ggc	gat	ctc	att	gtt	ctt	act	ttg	aac	cgc	gga	cag	ggg	tac	tcg
Asp	Gly	Asp	Leu	Ile	Val	Leu	Thr	Leu	Asn	Arg	Gly	Gln	Gly	Tyr	Ser
		400					405					410			
ggg	caa	tac	tgc	ttt	aac	act	gga	aag	aac	aac	aag	act	tgg	gac	aga
Gly	Gln	Tyr	Cys	Phe	Asn	Thr	Gly	Lys	Asn	Asn	Lys	Thr	Trp	Asp	Arg
	415					420					425				
gtc	ttt	gga	agt	gga	act	gtt	acc	tct	gat	ggc	aat	ggc	cag	gtt	tgt

1001827087_1.txt

Val Phe Gly Ser Gly Thr Val Thr Ser Asp Gly Asn Gly Gln Val Cys
430 435 440 445

gtt agc tac act aac ggc gag cct gaa gtc ttg gtt gcg tct agc ta 1510
Val Ser Tyr Thr Asn Gly Glu Pro Glu Val Leu Val Ala Ser Ser 460

g 1511

<210> 481
<211> 88
<212> PRT
<213> Fungi

<400> 481
Met Lys Phe Ser Leu Leu Ala Thr Ile Val Ala Ser Ile Ser Pro Leu
1 5 10 15
Ala Arg Ala Ala Asp Ala Asn Ala Trp Lys Ser Arg Asn Ile Tyr Phe
20 25 30
Ala Leu Thr Asp Arg Val Ala Arg Ser Ala Asp Asp Asn Gly Gly Ser
35 40 45
Ala Cys Gly Asn Leu Gly Asn Tyr Cys Gly Gly Thr Phe Lys Gly Leu
50 55 60
Glu Ser Lys Leu Asp Tyr Ile Lys Gly Met Gly Phe Asp Ala Ile Trp
65 70 75 80
Ile Thr Pro Val Val Asp Ser Ser
85

<210> 482
<211> 35
<212> PRT
<213> Fungi

<400> 482
Asp Gly Gly Tyr His Gly Tyr Trp Ala Lys Asp Leu Tyr Ala Val Asn
1 5 10 15
Ser Lys Tyr Gly Thr Ala Asp Asp Leu Lys Ser Leu Val Lys Ser Ala
20 25 30
His Asp Lys
35

<210> 483
<211> 337
<212> PRT
<213> Fungi

<400> 483
Asn Met Tyr Val Met Cys Asp Val Val Ala Asn His Met Gly Lys Gly
1 5 10 15
Ile Ser Asp His Lys Pro Ser Pro Leu Asn Glu Gln Ser Ser Tyr His
20 25 30
Thr Pro Cys Asp Ile Asp Tyr Ser Asn Gln Thr Ser Ile Glu Gln Cys
35 40 45
Glu Ile Ala Gly Leu Pro Asp Leu Asn Thr Gly Ser Asp Thr Val Lys
50 55 60
Lys Val Leu Tyr Asp Trp Ile Lys Trp Leu Val Ser Glu Tyr Ser Phe
65 70 75 80
Asp Gly Ile Arg Ile Asp Thr Val Lys His Val Glu Lys Pro Phe Trp
85 90 95
Pro Gly Phe Gln Asp Ala Ala Gly Val Tyr Ala Ile Gly Glu Val Trp
100 105 110
Asp Gly Gly Pro Asp Tyr Leu Ala Gly Tyr Ala Gln Val Met Pro Gly
115 120 125
Leu Leu Asn Tyr Ala Met Tyr Tyr Pro Met Asn Arg Phe Tyr Gln Gln
130 135 140
Lys Gly Asp Pro Ser Asp Val Val Ala Met His Asp Glu Ile Ser Asn
145 150 155 160
Lys Phe Pro Asp Pro Thr Ile Leu Gly Thr Phe Ile Asp Asn His Asp

1001827087_1.txt

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      165      170      175
Asn Pro Arg Trp Leu Ser Gln Lys Asn Asp Lys Ala Leu Leu Lys Asn
      180      190
Ala Leu Ala Tyr Val Ile Leu Ala Arg Gly Ile Pro Ile Val Tyr Tyr
      195      200      205
Gly Thr Glu Gln Gly Tyr Ala Gly Gly Asn Asp Pro Ala Asn Arg Glu
      210      215      220
Asp Leu Trp Arg Ser Ser Phe Ser Thr Asp Ala Asp Leu Tyr Gln His
      225      230      235
Ile Ser Arg Leu Ser Lys Ala Arg Ser Ala Val Gly Gly Leu Gly Gly
      245      250      255
Asn Asp His Lys His Leu Tyr Ser Gln Asp Ser Ala Tyr Ala Trp Ser
      260      265      270
Arg Ala Asp Gly Asp Leu Ile Val Leu Thr Leu Asn Arg Gly Gln Gly
      275      280      285
Tyr Ser Gly Gln Tyr Cys Phe Asn Thr Gly Lys Asn Asn Lys Thr Trp
      290      295      300
Asp Arg Val Phe Gly Ser Gly Thr Val Thr Ser Asp Gly Asn Gly Gln
      305      310      315
Val Cys Val Ser Tyr Thr Asn Gly Glu Pro Glu Val Leu Val Ala Ser
      325      330      335
Ser

```

<210> 484
 <211> 1383
 <212> DNA
 <213> Fungi

```

<400> 484
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agtgcctgatg ataatggcgg cagtgcctgc ggaacctcgc ggaattactg tgggtggaact      180
tttaagggct tagagtcaaa gcttgattat atcaagggca tgggatttga tgctatttgg      240
attactcctg ttgttgacag tagtgatggg ggataccatg gatactgggc taaggatctg      300
tatgcgggtca actccaagta tgggtactgca gatgatttga agagtcttgt caaatctgct      360
catgacaaga acatgtacgt catgtgcgac gtcgtcgcaa accacatggg caaaggcatc      420
tcagaccaca aaccctcccc cctcaacgaa caaagctcat accacacccc ctgcatatc      480
gactacagca accagaccag catcgagcag tgcgaaatcg ccggtcttcc cgacctcaac      540
accggcagcg acaccgtcaa gaaagtcctc tacgactgga tcaagtggct tgtctccgag      600
tacagcttcg acggtatccg catcgataca gtcaaacatg tcgaaaagcc gttctggcct      660
ggttttcaag acgctgctgg tgtttacgcc atcgggtgagg tttgggatgg agggcctgat      720
tacctagctg gttatgcaca ggtcatgcct ggtctcttga actatgctat gtattacccc      780
atgaaccggt tctaccagca aaaaggcgat ccttcagatg ttgtcgctat gcacgatgag      840
attagcaaca aattccccga tcccactatc ctcggaacat tcatcgacaa ccacgataac      900
cctcgctggg tgagccaaaa gaacgacaaa gctctctcctc agaacgccct cgcatacgtc      960
atcctagccc gaggtattcc catcgtctac tacggaacag agcaagggtta cgctggcggc      1020
aatgatcccc ccaaccgtga ggatctcttg cgaagcagct tcagcaccga cgcggatctg      1080
tatcaacaca tttcacgcct ctccaaggct cgcctggcgg tcggtgggtc tgggtggaat      1140
gatcacaagc atctctactc gcaggacagc gcgtatgctt ggagtcgtgc tgatggcgat      1200
ctcattgttc ttactttgaa ccgcgagacag gggactctcg ggcaatactg ctttaacact      1260
ggaaagaaca acaagacttg ggacagagtc tttggaagtg gaactgttac ctctgatggc      1320
aatggtcagg tttgtgttag ctacactaac ggcgagcctg aagtcttggt tgcgtctagc      1380
tag

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<210> 485
 <211> 460
 <212> PRT
 <213> Fungi

<220>
 <221> SIGNAL
 <222> (1)...(19)

<221> DOMAIN
 <222> (32)...(371)
 <223> Alpha amylase, catalytic domain

1001827087_1.txt

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<400> 485
Met Lys Phe Ser Leu Leu Ala Thr Ile Val Ala Ser Ile Ser Pro Leu
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20 25 30
Ala Leu Thr Asp Arg Val Ala Arg Ser Ala Asp Asp Asn Gly Gly Ser
35 40 45
Ala Cys Gly Asn Leu Gly Asn Tyr Cys Gly Gly Thr Phe Lys Gly Leu
50 55 60
Glu Ser Lys Leu Asp Tyr Ile Lys Gly Met Gly Phe Asp Ala Ile Trp
65 70 75 80
Ile Thr Pro Val Val Asp Ser Ser Asp Gly Gly Tyr His Gly Tyr Trp
85 90 95
Ala Lys Asp Leu Tyr Ala Val Asn Ser Lys Tyr Gly Thr Ala Asp Asp
100 105 110
Leu Lys Ser Leu Val Lys Ser Ala His Asp Lys Asn Met Tyr Val Met
115 120 125
Cys Asp Val Val Ala Asn His Met Gly Lys Gly Ile Ser Asp His Lys
130 135 140
Pro Ser Pro Leu Asn Glu Gln Ser Ser Tyr His Thr Pro Cys Asp Ile
145 150 155 160
Asp Tyr Ser Asn Gln Thr Ser Ile Glu Gln Cys Glu Ile Ala Gly Leu
165 170 175
Pro Asp Leu Asn Thr Gly Ser Asp Thr Val Lys Lys Val Leu Tyr Asp
180 185 190
Trp Ile Lys Trp Leu Val Ser Glu Tyr Ser Phe Asp Gly Ile Arg Ile
195 200 205
Asp Thr Val Lys His Val Glu Lys Pro Phe Trp Pro Gly Phe Gln Asp
210 215 220
Ala Ala Gly Val Tyr Ala Ile Gly Glu Val Trp Asp Gly Gly Pro Asp
225 230 235 240
Tyr Leu Ala Gly Tyr Ala Gln Val Met Pro Gly Leu Leu Asn Tyr Ala
245 250 255
Met Tyr Tyr Pro Met Asn Arg Phe Tyr Gln Gln Lys Gly Asp Pro Ser
260 265 270
Asp Val Val Ala Met His Asp Glu Ile Ser Asn Lys Phe Pro Asp Pro
275 280 285
Thr Ile Leu Gly Thr Phe Ile Asp Asn His Asp Asn Pro Arg Trp Leu
290 295 300
Ser Gln Lys Asn Asp Lys Ala Leu Leu Lys Asn Ala Leu Ala Tyr Val
305 310 315 320
Ile Leu Ala Arg Gly Ile Pro Ile Val Tyr Tyr Gly Thr Glu Gln Gly
325 330 335
Tyr Ala Gly Gly Asn Asp Pro Ala Asn Arg Glu Asp Leu Trp Arg Ser
340 345 350
Ser Phe Ser Thr Asp Ala Asp Leu Tyr Gln His Ile Ser Arg Leu Ser
355 360 365
Lys Ala Arg Ser Ala Val Gly Gly Leu Gly Gly Asn Asp His Lys His
370 375 380
Leu Tyr Ser Gln Asp Ser Ala Tyr Ala Trp Ser Arg Ala Asp Gly Asp
385 390 395 400
Leu Ile Val Leu Thr Leu Asn Arg Gly Gln Gly Tyr Ser Gly Gln Tyr
405 410 415
Cys Phe Asn Thr Gly Lys Asn Asn Lys Thr Trp Asp Arg Val Phe Gly
420 425 430 435
Ser Gly Thr Val Thr Ser Asp Gly Asn Gly Gln Val Cys Val Ser Tyr
440 445
Thr Asn Gly Glu Pro Glu Val Leu Val Ala Ser Ser
450 455 460

```

```

<210> 486
<211> 2131
<212> DNA
<213> Fungi

```

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<220>
<221> CDS
<222> (1)...(61)

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<223> Exon
<221> CDS
<222> (108)...(356)
<223> Exon

<221> CDS
<222> (410)...(625)
<223> Exon

<221> CDS
<222> (673)...(1270)
<223> Exon

<221> CDS
<222> (1324)...(2131)
<223> Exon

<400> 486
atg aga cga aaa agt aca gac aag tac aag aaa gta tca ata aga gct      48
Met Arg Arg Lys Ser Thr Asp Lys Tyr Lys Lys Val Ser Ile Arg Ala
 1              5              10              15

cat ctt gct gca t gttatattgt ttgtacgaca gttgtttctt cttttcttga      101
His Leu Ala Ala
          20

cttgag gt gaa caa ttg gca ata tcc aag atg ctg ttt tct cga act gcc      151
Cys Glu Gln Leu Ala Ile Ser Lys Met Leu Phe Ser Arg Thr Ala
          25              30              35

acg att ttg tct ctg cta tgt gta caa gca act gca att tca ccg aga      199
Thr Ile Leu Ser Leu Leu Cys Val Gln Ala Thr Ala Ile Ser Pro Arg
          40              45              50

ggc tcg gcc gca agt tat agt agt ctt gat tcc ttc att gct tcc gag      247
Gly Ser Ala Ala Ser Tyr Ser Ser Leu Asp Ser Phe Ile Ala Ser Glu
          55              60              65

aga aag att gct ctg cag ggt gtt ctc aac aat att gga cca gac gga      295
Arg Lys Ile Ala Leu Gln Gly Val Leu Asn Asn Ile Gly Pro Asp Gly
          70              75              80

tcg aaa gca tca gga gca ggc aat tat gtc att gca agt cct tcc aag      343
Ser Lys Ala Ser Gly Ala Gly Asn Tyr Val Ile Ala Ser Pro Ser Lys
          85              90              95

acc aat cct gat t gtaagaaata ccaatgttat gtttccaggc cgaatcttgc      396
Thr Asn Pro Asp
100

taattgattc cag ac ttc tat acc tgg act cga gat tct gcg tta acg tta      447
Tyr Phe Tyr Thr Trp Thr Arg Asp Ser Ala Leu Thr Leu
          105              110              115

aaa atg ctt gtt gat gaa ttt att ttt ggc aat cgt gca ctc cag cct      495
Lys Met Leu Val Asp Glu Phe Ile Phe Gly Asn Arg Ala Leu Gln Pro
          120              125              130

cag att gag aaa tac gtg aaa gct caa gcc atc ctt caa act gtt acg      543
Gln Ile Glu Lys Tyr Val Lys Ala Gln Ala Ile Leu Gln Thr Val Thr
          135              140              145

aat cct tct ggc act ttc ctt cct aac ggt tta gga ctt ggt gaa ccc      591
Asn Pro Ser Gly Thr Phe Leu Pro Asn Gly Leu Gly Leu Gly Glu Pro
          150              155              160

aaa tac gag gtt gac ggt act cga ttt aat ggt g gtatgtaatt      635

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Lys 165	Tyr	Glu	Val	Asp	Gly 170	Thr	Arg	Phe	Asn	Gly 175							
ttcagaggta acttctggca ctgctgacgt tccccag ca tgg ggt aga cct cag																	689
Ala Trp Gly Arg Pro Gln																	
cgc	gat	ggt	cca	gct	ctt	cgt	gct	atc	gca	tta	atg	gcc	tac	agc	gat		
Arg	Asp	Gly	Pro 185	Ala	Leu	Arg	Ala	Ile 190	Ala	Leu	Met	Ala	Tyr 195	Ser	Asp		
737																	
tgg	tta	att	aat	cat	aat	cag	ctt	caa	aga	gcc	aag	cta	gtt	ata	tgg		
Trp	Leu	Ile 200	Asn	His	Asn	Gln	Leu 205	Gln	Arg	Ala	Lys	Leu 210	Val	Ile	Trp		
785																	
cct	ata	atc	tca	aat	gac	tta	tca	tac	att	ggc	gaa	tat	tgg	aac	caa		
Pro	Ile 215	Ile	Ser	Asn	Asp	Leu 220	Ser	Tyr	Ile	Gly	Glu 225	Tyr	Trp	Asn	Gln		
833																	
act	gga	tat	gat	tta	tgg	gaa	gaa	gtt	cta	gga	tct	agc	ttt	ttt	acc		
Thr 230	Gly	Tyr	Asp	Leu	Trp 235	Glu	Glu	Val	Leu	Gly 240	Ser	Ser	Phe	Phe	Thr 245		
881																	
gta	caa	aat	caa	cac	cgt	gct	ttt	gtt	gaa	gcc	gga	aat	ctg	gcc	aat		
Val	Gln	Asn	Gln	His 250	Arg	Ala	Phe	Val	Glu 255	Ala	Gly	Asn	Leu	Ala 260	Asn		
929																	
aga	ctc	ggt	gtt	gaa	tct	cca	gga	tat	ctt	aat	gca	tta	gaa	gct	cta		
Arg	Leu	Gly	Val 265	Glu	Ser	Pro	Gly	Tyr 270	Leu	Asn	Ala	Leu	Glu 275	Ala	Leu		
977																	
tgc	ttc	ctt	aat	caa	agt	ttt	tgg	aac	gga	gaa	tat	atc	gtt	tcg	aac		
Cys	Phe	Leu 280	Asn	Gln	Ser	Phe	Trp 285	Asn	Gly	Glu	Tyr	Ile 290	Val	Ser	Asn		
1025																	
atc	aat	gtc	gat	aac	gga	aga	act	ggg	tta	gac	gga	gcc	tca	att	ctt		
Ile	Asn 295	Val	Asp	Asn	Gly	Arg 300	Thr	Gly	Leu	Asp	Gly 305	Ala	Ser	Ile	Leu		
1073																	
gga	tct	att	tct	atc	ttc	gat	att	gat	ggc	tct	tgc	gat	agc	cca	aat		
Gly 310	Ser	Ile	Ser	Ile	Phe 315	Asp	Ile	Asp	Gly	Ser 320	Cys	Asp	Ser	Pro	Asn 325		
1121																	
tta	caa	ccg	tgt	agt	agc	aga	gcg	ttg	tct	aat	ttt	agg	gca	gtc	gtg		
Leu	Gln	Pro	Cys	Ser 330	Ser	Arg	Ala	Leu	Ser 335	Asn	Phe	Arg	Ala	Val 340	Val		
1169																	
aac	agc	ttc	cga	ttt	tat	agt	atc	aat	gct	ggg	att	aaa	aac	aat	tca		
Asn	Ser	Phe	Arg 345	Phe	Tyr	Ser	Ile	Asn 350	Ala	Gly	Ile	Lys	Asn 355	Asn	Ser		
1217																	
ggt	atc	gcc	gtt	gga	aga	tat	gcc	gaa	gac	att	tat	caa	gga	gga	aat		
Gly	Ile	Ala 360	Val	Gly	Arg	Tyr	Ala 365	Glu	Asp	Ile	Tyr	Gln 370	Gly	Gly	Asn		
1265																	
cca	tg	gtactccatt	ccacccctct		gcggctcatt		acctttgagg		ctaactctcg							1320	
Pro 375																	
cag	g	tac	tta	ctt	acc	act	gcg	gcc	gca	gaa	ttt	ctt	tat	gat	gct	gtt	
		Tyr	Leu	Leu	Thr	Thr 380	Ala	Ala	Ala	Glu	Phe 385	Leu	Tyr	Asp	Ala	Val 390	
1369																	
gca	caa	tgg	aga	aat	cag	aaa	aaa	ctt	acc	gtt	gat	tca	aca	tcc	att		
Ala	Gln	Trp	Arg	Asn 395	Gln	Lys	Lys	Leu	Thr 400	Val	Asp	Ser	Thr	Ser 405	Ile		
1417																	
aat	ttc	ttc	cgc	gat	ctt	tac	cca	tct	atc	aaa	ctt	aag	agt	tac	aca		
1465																	

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Asn	Phe	Phe	Arg 410	Asp	Leu	Tyr	Pro	Ser 415	Ile	Lys	Leu	Lys	Ser 420	Tyr	Thr		
aac	agg	gac	aac	gaa	ttt	aac	aag	att	ctt	aat	gtc	gta	aca	aaa	tac	1513	
Asn	Arg	Asp 425	Asn	Glu	Phe	Asn	Lys 430	Ile	Leu	Asn	Val	Val 435	Thr	Lys	Tyr		
gcc	gac	tct	ttt	gtc	gca	att	gct	caa	aaa	tat	aca	ccc	agt	agt	gga	1561	
Ala	Asp 440	Ser	Phe	Val	Ala	Ile 445	Ala	Gln	Lys	Tyr	Thr 450	Pro	Ser	Ser	Gly		
tct	tta	gcc	gaa	caa	ttc	gat	cgc	aat	aca	gga	gcc	ccc	att	tct	gcc	1609	
Ser 455	Leu	Ala	Glu	Gln	Phe 460	Asp	Arg	Asn	Thr	Gly 465	Ala	Pro	Ile	Ser	Ala 470		
aat	gac	ctt	aca	tgg	agt	tat	gcg	gca	ttt	gtc	acc	atg	gct	caa	aga	1657	
Asn	Asp	Leu	Thr	Trp 475	Ser	Tyr	Ala	Ala	Phe 480	Val	Thr	Met	Ala	Gln	Arg		
cgt	tca	ggg	caa	tat	cct	gct	agt	tgg	gga	agt	gcg	gca	gca	tcg	gtt	1705	
Arg	Ser	Gly	Gln 490	Tyr	Pro	Ala	Ser	Trp 495	Gly	Ser	Ala	Ala	Ala 500	Ser	Val		
cca	ccc	cca	aat	tgc	gca	tct	tca	tct	tct	caa	gga	gtg	tat	gtt	cca	1753	
Pro	Pro	Pro 505	Asn	Cys	Ala	Ser	Ser 510	Ser	Ser	Gln	Gly	Val 515	Tyr	Val	Pro		
gcc	att	gca	gca	ggg	gcc	cca	aat	gta	acg	gca	aca	tgt	caa	tat	cca	1801	
Ala	Ile 520	Ala	Ala	Gly	Ala	Pro 525	Asn	Val	Thr	Ala	Thr 530	Cys	Gln	Tyr	Pro		
gtg	cga	ttt	tat	gtt	aat	gcc	aca	aca	tat	tat	ggg	gaa	aac	ctc	tac	1849	
Val 535	Arg	Phe	Tyr	Val	Asn 540	Ala	Thr	Thr	Tyr	Tyr 545	Gly	Glu	Asn	Leu	Tyr 550		
att	att	gga	aat	aca	acg	gat	ttg	gga	gca	tgg	aac	ttg	aac	agt	gca	1897	
Ile	Ile	Gly	Asn	Thr 555	Thr	Asp	Leu	Gly	Ala 560	Trp	Asn	Leu	Asn	Ser 565	Ala		
tta	cct	atg	aat	gca	ggg	atg	tac	acg	acg	gaa	aac	cct	gtt	tgg	tat	1945	
Leu	Pro	Met	Asn 570	Ala	Gly	Met	Tyr	Thr 575	Thr	Glu	Asn	Pro	Val 580	Trp	Tyr		
gtg	gat	gca	cag	tta	acg	gct	ggg	gag	cca	gtc	agt	tat	gtt	tat	gtg	1993	
Val	Asp	Ala 585	Gln	Leu	Thr	Ala	Gly 590	Glu	Pro	Val	Ser	Tyr 595	Val	Tyr	Val		
aga	gaa	caa	gat	tgc	ggg	caa	gct	cca	att	tac	gag	acg	aat	aat	cga	2041	
Arg	Glu 600	Gln	Asp	Cys	Gly	Gln 605	Ala	Pro	Ile	Tyr	Glu 610	Thr	Asn	Asn	Arg		
aca	agt	gtg	gtt	ccg	gaa	tgt	ggg	aca	ggg	ggg	tcg	acg	atc	agg	cgt	2089	
Thr 615	Ser	Val	Val	Pro	Glu 620	Cys	Gly	Thr	Gly	Gly 625	Ser	Thr	Ile	Arg	Arg 630		
gat	gta	tgg	gtc	ggg	agt	gga	gga	agt	agt	gga	ggg	tgt	taa			2131	
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 <212> PRT
 <213> Fungi

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His Leu Ala Ala
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<211> 83
<212> PRT
<213> Fungi

<400> 488
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Ser Ala Ala Ser Tyr Ser Ser Leu Asp Ser Phe Ile Ala Ser Glu Arg
35 40 45
Lys Ile Ala Leu Gln Gly Val Leu Asn Asn Ile Gly Pro Asp Gly Ser
50 55 60
Lys Ala Ser Gly Ala Gly Asn Tyr Val Ile Ala Ser Pro Ser Lys Thr
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Asn Pro Asp

<210> 489
<211> 72
<212> PRT
<213> Fungi

<400> 489
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Val Asp Glu Phe Ile Phe Gly Asn Arg Ala Leu Gln Pro Gln Ile Glu
20 25 30
Lys Tyr Val Lys Ala Gln Ala Ile Leu Gln Thr Val Thr Asn Pro Ser
35 40 45
Gly Thr Phe Leu Pro Asn Gly Leu Gly Leu Gly Glu Pro Lys Tyr Glu
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Val Asp Gly Thr Arg Phe Asn Gly
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<211> 200
<212> PRT
<213> Fungi

<400> 490
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Leu Met Ala Tyr Ser Asp Trp Leu Ile Asn His Asn Gln Leu Gln Arg
20 25 30
Ala Lys Leu Val Ile Trp Pro Ile Ile Ser Asn Asp Leu Ser Tyr Ile
35 40 45
Gly Glu Tyr Trp Asn Gln Thr Gly Tyr Asp Leu Trp Glu Glu Val Leu
50 55 60
Gly Ser Ser Phe Phe Thr Val Gln Asn Gln His Arg Ala Phe Val Glu
65 70 75 80
Ala Gly Asn Leu Ala Asn Arg Leu Gly Val Glu Ser Pro Gly Tyr Leu
85 90 95
Asn Ala Leu Glu Ala Leu Cys Phe Leu Asn Gln Ser Phe Trp Asn Gly
100 105 110
Glu Tyr Ile Val Ser Asn Ile Asn Val Asp Asn Gly Arg Thr Gly Leu
115 120 125
Asp Gly Ala Ser Ile Leu Gly Ser Ile Ser Ile Phe Asp Ile Asp Gly
130 135 140
Ser Cys Asp Ser Pro Asn Leu Gln Pro Cys Ser Ser Arg Ala Leu Ser
145 150 155 160
Asn Phe Arg Ala Val Asn Ser Phe Arg Phe Tyr Ser Ile Asn Ala
165 170 175
Gly Ile Lys Asn Asn Ser Gly Ile Ala Val Gly Arg Tyr Ala Glu Asp

180
Ile Tyr Gln Gly Gly Asn Pro Trp
195 200

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<212> PRT
<213> Fungi

<400> 491
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20 25 30
Phe Phe Arg Asp Leu Tyr Pro Ser Ile Lys Leu Lys Ser Tyr Thr Asn
35 40 45
Arg Asp Asn Glu Phe Asn Lys Ile Leu Asn Val Val Thr Lys Tyr Ala
50 55 60
Asp Ser Phe Val Ala Ile Ala Gln Lys Tyr Thr Pro Ser Ser Gly Ser
65 70 75 80
Leu Ala Glu Gln Phe Asp Arg Asn Thr Gly Ala Pro Ile Ser Ala Asn
85 90 95
Asp Leu Thr Trp Ser Tyr Ala Ala Phe Val Thr Met Ala Gln Arg Arg
100 105 110
Ser Gly Gln Tyr Pro Ala Ser Trp Gly Ser Ala Ala Ser Val Pro
115 120 125
Pro Pro Asn Cys Ala Ser Ser Ser Ser Gln Gly Val Tyr Val Pro Ala
130 135 140
Ile Ala Ala Gly Ala Pro Asn Val Thr Ala Thr Cys Gln Tyr Pro Val
145 150 155 160
Arg Phe Tyr Val Asn Ala Thr Thr Tyr Tyr Gly Glu Asn Leu Tyr Ile
165 170 175
Ile Gly Asn Thr Thr Asp Leu Gly Ala Trp Asn Leu Asn Ser Ala Leu
180 185 190
Pro Met Asn Ala Gly Met Tyr Thr Thr Glu Asn Pro Val Trp Tyr Val
195 200 205
Asp Ala Gln Leu Thr Ala Gly Glu Pro Val Ser Tyr Val Tyr Val Arg
210 215 220
Glu Gln Asp Cys Gly Gln Ala Pro Ile Tyr Glu Thr Asn Asn Arg Thr
225 230 235 240
Ser Val Val Pro Glu Cys Gly Thr Gly Gly Ser Thr Ile Arg Arg Asp
245 250 255
Val Trp Val Gly Ser Gly Gly Ser Ser Gly Gly Cys
260 265

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<211> 1932
<212> DNA
<213> Fungi

<400> 492
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ctatgtgtac aagcaactgc aatttcaccg agagggttcgg ccgcaagtta tagtagtctt 180
gattccttca ttgcttccga gagaaagatt gctctgcagg gtgttctcaa caatattgga 240
ccagacggat cgaaagcatc aggagcaggc aattatgtca ttgcaagtcc ttccaagacc 300
aatcctgatt acttctatac ctggactcga gattctgcgt taacgttaaa aatgcttggt 360
gatgaattta tttttggcaa tcgtgcactc cagcctcaga ttgagaaata cgtgaaagct 420
caagccatcc ttcaaactgt tacgaatcct tctggcactt tccttcctaa cggtttagga 480
cttggtgaac ccaaatacga ggttgacggt actcgattta atggtgcatg gggtagacct 540
cagcgcgatg gtccagctct tcgtgctatc gcattaatgg cctacagcga ttggttaatt 600
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tcatacattg gcgaatatgt gaaccaaact ggatatgatt tatgggaaga agttctagga 720
tctagctttt ttaccgtaca aaatcaacac cgtgcttttg ttgaagccgg aaatctggcc 780
aatagactcg gtgttgaaatc tccaggatat cttaatgcat tagaagctct atgcttcctt 840
aatcaaaagt tttggaacgg agaatatatc gtttcgaaca tcaatgtcga taacggaaga 900
actgggttag acggagcctc aattcttgga tctatttcta tcttcgatat tgatggctct 960
tgcgatagcc caaatttaca accgtgtagt agcagagcgt tgtctaattt tagggcagtc 1020

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gttggaagat	atgccgaaga	catttatcaa	ggaggaaatc	catggtactt	acttaccact	1140
gcggccgcag	aattttcttta	tgatgctggt	gcacaatgga	gaaatcagaa	aaaacttacc	1200
gttgattcaa	catccattaa	tttcttccgc	gatctttacc	catctatcaa	acttaagagt	1260
tacacaaaca	gggacaacga	atttaacaag	attcttaatg	tcgtaacaaa	atacgccgac	1320
tcttttgtcg	caattgctca	aaaatataca	cccagtagtg	gatcttttagc	cgaacaattc	1380
gatcgcaata	caggagcccc	catttctgcc	aatgacctta	catggagtta	tgcggcattt	1440
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tcggttccac	ccccaaattg	cgcacatctca	tcttctcaag	gagtgtatgt	tccagccatt	1560
gcagcagggg	ccccaaatgt	aacggcaaca	tgtcaatatc	cagtgcgatt	ttatgttaat	1620
gccacaacat	attatggtga	aaacctctac	attattggaa	atacaacgga	tttgggagca	1680
tggaacttga	acagtgcatt	acctatgaat	gcagggatgt	acacgacgga	aaaccctggt	1740
tggtatgtgg	atgcgcgatt	aacggctggg	gagccagtca	gttatgttta	tgtgagagaa	1800
caagattgcg	gtcaagctcc	aatttacgag	acgaataatc	gaacaagtgt	ggttccggaa	1860
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 <211> 643
 <212> PRT
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<220>
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 <222> (532)...(628)
 <223> Starch binding domain

<221> DOMAIN
 <222> (69)...(485)
 <223> Glycosyl hydrolases family 15

<221> SIGNAL
 <222> (1)...(54)

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 35 40 45
 Ser Pro Arg Gly Ser Ala Ala Ser Tyr Ser Ser Leu Asp Ser Phe Ile
 50 55 60
 Ala Ser Glu Arg Lys Ile Ala Leu Gln Gly Val Leu Asn Asn Ile Gly
 65 70 75 80
 Pro Asp Gly Ser Lys Ala Ser Gly Ala Gly Asn Tyr Val Ile Ala Ser
 85 90 95
 Pro Ser Lys Thr Asn Pro Asp Tyr Phe Tyr Thr Trp Thr Arg Asp Ser
 100 105 110
 Ala Leu Thr Leu Lys Met Leu Val Asp Glu Phe Ile Phe Gly Asn Arg
 115 120 125
 Ala Leu Gln Pro Gln Ile Glu Lys Tyr Val Lys Ala Gln Ala Ile Leu
 130 135 140
 Gln Thr Val Thr Asn Pro Ser Gly Thr Phe Leu Pro Asn Gly Leu Gly
 145 150 155 160
 Leu Gly Glu Pro Lys Tyr Glu Val Asp Gly Thr Arg Phe Asn Gly Ala
 165 170 175
 Trp Gly Arg Pro Gln Arg Asp Gly Pro Ala Leu Arg Ala Ile Ala Leu
 180 185 190
 Met Ala Tyr Ser Asp Trp Leu Ile Asn His Asn Gln Leu Gln Arg Ala
 195 200 205
 Lys Leu Val Ile Trp Pro Ile Ile Ser Asn Asp Leu Ser Tyr Ile Gly
 210 215 220
 Glu Tyr Trp Asn Gln Thr Gly Tyr Asp Leu Trp Glu Glu Val Leu Gly
 225 230 235 240
 Ser Ser Phe Phe Thr Val Gln Asn Gln His Arg Ala Phe Val Glu Ala
 245 250 255
 Gly Asn Leu Ala Asn Arg Leu Gly Val Glu Ser Pro Gly Tyr Leu Asn

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Ala	Leu	Glu	260	Ala	Leu	Cys	Phe	Leu	265	Asn	Gln	Ser	Phe	Trp	270	Asn	Gly	Glu
Tyr	Ile	Val	275	Ser	Asn	Ile	Asn	Val	280	Asp	Asn	Gly	Arg	Thr	285	Gly	Leu	Asp
Gly	Ala	Ser	290	Ile	Leu	Gly	Ser	Ile	295	Ser	Ile	Phe	Asp	Ile	300	Asp	Gly	Ser
Cys	Asp	Ser	305	Pro	Asn	Leu	Gln	Pro	310	Cys	Ser	Ser	Arg	Ala	315	Leu	Ser	Asn
Phe	Arg	Ala	325	Val	Val	Asn	Ser	Phe	330	Arg	Phe	Tyr	Ser	Ile	335	Asn	Ala	Gly
Ile	Lys	Asn	340	Asn	Ser	Gly	Ile	Ala	345	Val	Gly	Arg	Tyr	Ala	350	Glu	Asp	Ile
Tyr	Gln	Gly	355	Gly	Asn	Pro	Trp	Tyr	360	Leu	Leu	Thr	Thr	Ala	365	Ala	Ala	Glu
Phe	Leu	Tyr	370	Asp	Ala	Val	Ala	Gln	375	Trp	Arg	Asn	Gln	Lys	380	Lys	Leu	Thr
Val	Asp	Ser	385	Thr	Ser	Ile	Asn	Phe	390	Phe	Arg	Asp	Leu	Tyr	395	Pro	Ser	Ile
Lys	Leu	Lys	405	Tyr	Thr	Asn	Arg	Asp	410	Asn	Glu	Phe	Asn	Lys	415	Ile	Leu	
Asn	Val	Val	420	Thr	Lys	Tyr	Ala	Asp	425	Ser	Phe	Val	Ala	Ile	430	Ala	Gln	Lys
Tyr	Thr	Pro	435	Ser	Ser	Gly	Ser	Leu	440	Ala	Glu	Gln	Phe	Asp	445	Arg	Asn	Thr
Gly	Ala	Pro	450	Ile	Ser	Ala	Asn	Asp	455	Leu	Thr	Trp	Ser	Tyr	460	Ala	Ala	Phe
Val	Thr	Met	465	Ala	Gln	Arg	Arg	Ser	470	Gly	Gln	Tyr	Pro	Ala	475	Ser	Trp	Gly
Ser	Ala	Ala	485	Ser	Val	Pro	Pro	Pro	490	Asn	Cys	Ala	Ser	Ser	495	Ser	Ser	Ser
Gln	Gly	Val	500	Tyr	Val	Pro	Ala	Ile	505	Ala	Gly	Ala	Pro	Asn	510	Val	Thr	
Ala	Thr	Cys	515	Gln	Tyr	Pro	Val	Arg	520	Phe	Tyr	Val	Asn	Ala	525	Thr	Thr	Tyr
Tyr	Gly	Glu	530	Asn	Leu	Tyr	Ile	Ile	535	Gly	Asn	Thr	Thr	Asp	540	Leu	Gly	Ala
Trp	Asn	Leu	545	Asn	Ser	Ala	Leu	Pro	550	Met	Asn	Ala	Gly	Met	555	Tyr	Thr	Thr
Glu	Asn	Pro	565	Val	Trp	Tyr	Val	Asp	570	Ala	Gln	Leu	Thr	Ala	575	Gly	Glu	Pro
Val	Ser	Tyr	580	Val	Tyr	Val	Arg	Glu	585	Gln	Asp	Cys	Gly	Gln	590	Ala	Pro	Ile
Tyr	Glu	Thr	595	Asn	Asn	Arg	Thr	Ser	600	Val	Val	Pro	Glu	Cys	605	Gly	Thr	Gly
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Gly	Gly	Cys	625			630			635									640

<210> 494
 <211> 1586
 <212> DNA
 <213> Fungi

<220>
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 <222> (1)...(225)
 <223> Exon

<221> CDS
 <222> (283)...(450)
 <223> Exon

<221> CDS
 <222> (501)...(1586)
 <223> Exon

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<400> 494
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tta aac ggc gtt caa tca gca acg ccc gac gaa tgg agg act cgt tca      96
Leu Asn Gly Val Gln Ser Ala Thr Pro Asp Glu Trp Arg Thr Arg Ser
20      25      30

ata tat caa gta ttt aca gat cga ttt gca aga aca gat gtt tca aat      144
Ile Tyr Gln Val Phe Thr Asp Arg Phe Ala Arg Thr Asp Val Ser Asn
35      40      45

act acg gat tgc cct tct caa act cgg gga tac tgt ggt ggg aca tgg      192
Thr Thr Asp Cys Pro Ser Gln Thr Arg Gly Tyr Cys Gly Gly Thr Trp
50      55      60

caa ggc ata att aac aag ctc gat tat atc cag gtatgagagt agtgcaatca      245
Gln Gly Ile Ile Asn Lys Leu Asp Tyr Ile Gln
65      70      75

catattggta aaattacaag tgctaataagg ctattag gat atg ggg ttc aca gca      300
Asp Met Gly Phe Thr Ala
80

att tgg ata tca cca gtt gtt gaa caa gtt ggg gat cca ggt cgc ggc      348
Ile Trp Ile Ser Pro Val Val Glu Gln Val Gly Asp Pro Gly Arg Gly
85      90      95

ttt cat ggg tat tcg gct cag aat ttg tat ggt cta aac agt cac ttc      396
Phe His Gly Tyr Ser Ala Gln Asn Leu Tyr Gly Leu Asn Ser His Phe
100      105      110

gga gac gca gcg gat ttg aaa gcg cta gct acc gcg ctc cat gat cga      444
Gly Asp Ala Ala Asp Leu Lys Ala Leu Ala Thr Ala Leu His Asp Arg
115      120      125

gga atg gtgagactga acaagaaatt tgcacagata aagaaagcta attctggtag      500
Gly Met
130

tat ttg atg gtt gat gtc gtg gcg aac cat atg gga tct gat gat act      548
Tyr Leu Met Val Asp Val Val Ala Asn His Met Gly Ser Asp Asp Thr
135      140      145

gcg cat aac gtt gat ttc agc atc atg aat cca ttc aat gac agc aaa      596
Ala His Asn Val Asp Phe Ser Ile Met Asn Pro Phe Asn Asp Ser Lys
150      155      160

tac ttt cac agt gtc tgc ttt atc aat gat tac aac aat cag aca aat      644
Tyr Phe His Ser Val Cys Phe Ile Asn Asp Tyr Asn Asn Gln Thr Asn
165      170      175

gtt gaa cta tgc gag ctt gga acg gag cgg tat cca tta cct gat cta      692
Val Glu Leu Cys Glu Leu Gly Thr Glu Arg Tyr Pro Leu Pro Asp Leu
180      185      190      195

aac act act cga caa gat gtt cga gat ctc cat act act tgg atc aag      740
Asn Thr Thr Arg Gln Asp Val Arg Asp Leu His Thr Thr Trp Ile Lys
200      205      210

tca cta gta gcc aac tac tca att gac gga ttg aga gtg gac act gtt      788
Ser Leu Val Ala Asn Tyr Ser Ile Asp Gly Leu Arg Val Asp Thr Val
215      220      225

aga cat gta gag aag gat ttt tgg cct tta ttc aat gaa acc gcc ggt      836
Arg His Val Glu Lys Asp Phe Trp Pro Leu Phe Asn Glu Thr Ala Gly
230      235      240

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gtt Val	tat Tyr 245	tgt Cys	gtt Val	ggc Gly	gaa Glu	gtg Val 250	gca Ala	gat Asp	gga Gly	gac Asp	gta Val 255	gat Asp	tat Tyr	cta Leu	tgt Cys	884
ccc Pro 260	tac Tyr	caa Gln	gat Asp	tac Tyr	ata Ile 265	gat Asp	ggg Gly	ctt Leu	ctc Leu	tcc Ser 270	tat Tyr	gcc Ala	tca Ser	tac Tyr	ttc Phe 275	932
caa Gln	ctc Leu	aca Thr	aaa Lys	ttc Phe 280	ttt Phe	tcc Ser	aac Asn	acc Thr	tct Ser 285	gca Ala	aca Thr	tca Ser	gag Glu	aac Asn 290	tta Leu	980
ata Ile	gga Gly	caa Gln	tta Leu 295	caa Gln	tcc Ser	cag Gln	aat Asn	tat Tyr 300	caa Gln	tgt Cys	aaa Lys	gac Asp	acg Thr 305	aca Thr	ctt Leu	1028
ctg Leu	gga Gly	tcc Ser 310	ttc Phe	acc Thr	gag Glu	aac Asn	cat His 315	gat Asp	caa Gln	ccg Pro	cgc Arg	ttc Phe 320	gga Gly	aat Asn	tac Tyr	1076
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gca Ala 340	gat Asp	ggg Gly	ata Ile	cca Pro	att Ile 345	atc Ile	tat Tyr	caa Gln	ggg Gly	caa Gln 350	gaa Glu	cag Gln	cat His	ttc Phe	cat His 355	1172
ggc Gly	gca Ala	act Thr	gac Asp	cct Pro 360	tat Tyr	gat Asp	cgt Arg	gaa Glu	ccc Pro 365	ata Ile	tgg Trp	cct Pro	aca Thr	ggg Gly 370	tac Tyr	1220
gat Asp	act Thr	act Thr	tcc Ser 375	cca Pro	ctt Leu	tat Tyr	gtc Val	ctg Leu 380	gtc Val	aaa Lys	caa Gln	ttg Leu	aac Asn 385	gct Ala	ata Ile	1268
cgt Arg	tct Ser	ttg Leu 390	gcg Ala	ata Ile	gca Ala	cgc Arg	tcg Ser 395	gat Asp	aca Thr	tat Tyr	gct Ala	act Thr 400	tat Tyr	caa Gln	acc Thr	1316
caa Gln 405	atc Ile	gct Ala	tac Tyr	tca Ser	gat Asp	ccg Pro 410	cac His	aat Asn	att Ile	gcg Ala	ttt Phe 415	cgg Arg	agg Arg	gga Gly	gat Asp	1364
gag Glu 420	aaa Lys	tgc Cys	atg Met	agt Ser	cta Leu 425	atg Met	gtg Val	ctg Leu	aac Asn	aat Asn 430	atc Ile	ggg Gly	gag Glu	agt Ser	gca Ala 435	1412
gag Glu	gat Asp	tat Tyr	aca Thr	gtc Val 440	gag Glu	att Ile	gag Glu	aat Asn	gtg Val 445	ggg Gly	ttt Phe	gaa Glu	gct Ala	gga Gly 450	tcg Ser	1460
ata Ile	gtg Val	acg Thr	gat Asp 455	gta Val	ttg Leu	agt Ser	tgt Cys	aga Arg 460	aat Asn	gta Val	atg Met	gta Val	gat Asp 465	gag Glu	tat Tyr	1508
ggg Gly	ggg Gly	atg Met 470	gaa Glu	gtg Val	cca Pro	ttt Phe	gtg Val 475	agc Ser	ggg Gly	tta Leu	cct Pro	agt Ser 480	gta Val	agt Ser	cta Leu	1556
atc Ile 485	ctc Leu	tcc Ser	atg Met	aga Arg	ctg Leu	ttt Phe 490	aaa Lys	atc Ile	taa *							1586

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<211> 75

<212> PRT
<213> Fungi

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20 25 30
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35 40 45
Thr Thr Asp Cys Pro Ser Gln Thr Arg Gly Tyr Cys Gly Gly Thr Trp
50 55 60
Gln Gly Ile Ile Asn Lys Leu Asp Tyr Ile Gln
65 70 75

<210> 496
<211> 56
<212> PRT
<213> Fungi

<400> 496
Asp Met Gly Phe Thr Ala Ile Trp Ile Ser Pro Val Val Glu Gln Val
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Gly Asp Pro Gly Arg Gly Phe His Gly Tyr Ser Ala Gln Asn Leu Tyr
20 25 30
Gly Leu Asn Ser His Phe Gly Asp Ala Ala Asp Leu Lys Ala Leu Ala
35 40 45
Thr Ala Leu His Asp Arg Gly Met
50 55

<210> 497
<211> 361
<212> PRT
<213> Fungi

<400> 497
Tyr Leu Met Val Asp Val Val Ala Asn His Met Gly Ser Asp Asp Thr
1 5 10 15
Ala His Asn Val Asp Phe Ser Ile Met Asn Pro Phe Asn Asp Ser Lys
20 25 30
Tyr Phe His Ser Val Cys Phe Ile Asn Asp Tyr Asn Asn Gln Thr Asn
35 40 45
Val Glu Leu Cys Glu Leu Gly Thr Glu Arg Tyr Pro Leu Pro Asp Leu
50 55 60
Asn Thr Thr Arg Gln Asp Val Arg Asp Leu His Thr Thr Trp Ile Lys
65 70 75 80
Ser Leu Val Ala Asn Tyr Ser Ile Asp Gly Leu Arg Val Asp Thr Val
85 90 95
Arg His Val Glu Lys Asp Phe Trp Pro Leu Phe Asn Glu Thr Ala Gly
100 105 110
Val Tyr Cys Val Gly Glu Val Ala Asp Gly Asp Val Asp Tyr Leu Cys
115 120 125
Pro Tyr Gln Asp Tyr Ile Asp Gly Leu Leu Ser Tyr Ala Ser Tyr Phe
130 135 140
Gln Leu Thr Lys Phe Phe Ser Asn Thr Ser Ala Thr Ser Glu Asn Leu
145 150 155 160
Ile Gly Gln Leu Gln Ser Gln Asn Tyr Gln Cys Lys Asp Thr Thr Leu
165 170 175
Leu Gly Ser Phe Thr Glu Asn His Asp Gln Pro Arg Phe Gly Asn Tyr
180 185 190
Thr Ser Asp Leu Thr Leu Ala Lys Asn Ile Ile Thr Tyr Thr Met Leu
195 200 205
Ala Asp Gly Ile Pro Ile Ile Tyr Gln Gly Gln Glu Gln His Phe His
210 215 220
Gly Ala Thr Asp Pro Tyr Asp Arg Glu Pro Ile Trp Pro Thr Gly Tyr
225 230 235 240
Asp Thr Thr Ser Pro Leu Tyr Val Leu Val Lys Gln Leu Asn Ala Ile
245 250 255

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Arg Ser Leu Ala Ile Ala Arg Ser Asp Thr Tyr Ala Thr Tyr Gln Thr
260 270
Gln Ile Ala Tyr Ser Asp Pro His Asn Ile Ala Phe Arg Arg Gly Asp
275 280
Glu Lys Cys Met Ser Leu Met Val Leu Asn Asn Ile Gly Glu Ser Ala
290 295 300
Glu Asp Tyr Thr Val Glu Ile Glu Asn Val Gly Phe Glu Ala Gly Ser
305 310 315
Ile Val Thr Asp Val Leu Ser Cys Arg Asn Val Met Val Asp Glu Tyr
325 330 335
Gly Gly Met Glu Val Pro Phe Val Ser Gly Leu Pro Ser Val Ser Leu
340 345 350
Ile Leu Ser Met Arg Leu Phe Lys Ile
355 360

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<210> 498
 <211> 1479
 <212> DNA
 <213> Fungi

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<400> 498
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tttgcagaaa cagatgtttc aaatactacg gattgccctt ctcaaactcg gggatactgt 180
ggtgggacat ggcaaggcat aattaacaag ctcgattata tccaggatat ggggttcaca 240
gcaatttgga tatcaccagt tgttgaacaa gttggggatc caggtcgcgg cttcatggg 300
tattcggctc agaatttgta tggctaaac agtcacttcg gagacgcagc ggatttgaaa 360
gcgctagcta ccgcgctcca tgatcgagga atgtatttga tggttgatgt cgtggcgaac 420
catatgggat ctgatgatac tgcgcataac gttgatattc gcatcatgaa tccattcaat 480
gacagcaaat actttcacag tgtctgcttt atcaatgatt acaacaatca gacaaatggt 540
gaactatgcg agcttggaac ggagcgggat ccattacctg atctaaacac tactcgacaa 600
gatgttcgag atctccatac tacttgatc aagtcactag tagccaacta ctcaattgac 660
ggattgagag tggacactgt tagacatgta gagaaggatt tttggccttt attcaatgaa 720
accgccggtg tttatttgtt tggcgaagtg gcagatggag acgtagatta tctatgtccc 780
taccaagatt acatagatgg gcttctctcc tatgcctcat acttccaact cacaaaattc 840
ttttccaaca cctctgcaac atcagagaac ttaataggac aattacaatc ccagaattat 900
caatgtaaa acacgacact tctgggatcc ttcaccgaga accatgatca accgcgcttc 960
ggaaattaca ctagtgcact cacgttagct aaaaacatca tcacatatac gatgttagca 1020
gatggtatac caattatcta tcaagggcaa gaacagcatt tccatggcgc aactgaccct 1080
tatgatcgtg aaccatattg gcctacaggt tacgatacta cttccccact ttatgtcctg 1140
gtcaaacaat tgaacgctat acgttctttg gcgatagcac gctcggatac atatgtctact 1200
tatcaaacc aaatcgctta ctcagatccg cacaatattg cgtttcggag gggagatgag 1260
aaatgcatga gtctaattgt gctgaacaat atcggtgaga gtgcagagga ttatacagtc 1320
gagattgaga atgtggggtt tgaagctgga tcgatagtga cggatgtatt gagttgtaga 1380
aatgtaatgg tagatgagta tgggtgggatg gaagtgccat ttgtgagcgg gttacctagt 1440
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<210> 499
 <211> 492
 <212> PRT
 <213> Fungi

<220>
 <221> SIGNAL
 <222> (1)...(22)

<221> DOMAIN
 <222> (35)...(388)
 <223> Alpha amylase, catalytic domain

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<400> 499
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Leu Asn Gly Val Gln Ser Ala Thr Pro Asp Glu Trp Arg Thr Arg Ser
20 25 30
Ile Tyr Gln Val Phe Thr Asp Arg Phe Ala Arg Thr Asp Val Ser Asn
35 40 45
Thr Thr Asp Cys Pro Ser Gln Thr Arg Gly Tyr Cys Gly Gly Thr Trp

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50	Gln	Gly	Ile	Ile	Asn	Lys	55	Leu	Asp	Tyr	Ile	Gln	60	Asp	Met	Gly	Phe	Thr
65	Ala	Ile	Trp	Ile	Ser	Pro	70	Val	Val	Glu	Gln	75	Val	Gly	Asp	Pro	Gly	Arg
	Gly	Phe	His	Gly	Tyr	Ser	85	Ala	Gln	Asn	90	Leu	Tyr	Gly	Leu	Asn	Ser	His
	Phe	Gly	Asp	Ala	Ala	Asp	100	Leu	Lys	105	Ala	Leu	Ala	Thr	Ala	Leu	His	Asp
	Arg	Gly	Met	Tyr	Leu	Met	115	Val	Asp	120	Val	Val	Ala	Asn	His	Met	Gly	Ser
	Asp	Asp	Thr	Ala	His	Asn	130	Val	Asp	135	Phe	Ser	Ile	Met	Asn	Pro	Phe	Asn
145	Asp	Ser	Lys	Tyr	Phe	His	150	Ser	Val	Cys	Phe	155	Ile	Asn	Asp	Tyr	Asn	Asn
	Gln	Thr	Asn	Val	Glu	Leu	165	Cys	Glu	Leu	Gly	170	Thr	Glu	Arg	Tyr	Pro	Leu
	Pro	Asp	Leu	Asn	Thr	Thr	180	Arg	Gln	Asp	Val	185	Arg	Asp	Leu	His	Thr	Thr
	Trp	Ile	Lys	Ser	Leu	Val	195	Ala	Asn	Tyr	Ser	200	Ile	Asp	Gly	Leu	Arg	Val
	Asp	Thr	Val	Arg	His	Val	210	Glu	Lys	Asp	Phe	215	Trp	Pro	Leu	Phe	Asn	Glu
225	Thr	Ala	Gly	Val	Tyr	Cys	225	Val	Gly	Glu	Val	230	Ala	Asp	Gly	Asp	Val	Asp
	Tyr	Leu	Cys	Pro	Tyr	Gln	245	Asp	Tyr	Ile	Asp	250	Gly	Leu	Leu	Ser	Tyr	Ala
	Ser	Tyr	Phe	Gln	Leu	Thr	260	Lys	Phe	Phe	Ser	265	Asn	Thr	Ser	Ala	Thr	Ser
	Glu	Asn	Leu	Ile	Gly	Gln	275	Leu	Gln	Ser	Gln	280	Asn	Thr	Ser	Ala	Thr	Ser
	Thr	Thr	Leu	Leu	Gly	Ser	290	Phe	Thr	Glu	Asn	300	His	Asp	Gln	Pro	Arg	Phe
305	Gly	Asn	Tyr	Thr	Ser	Asp	310	Leu	Thr	Leu	Ala	315	Lys	Asn	Ile	Ile	Thr	Tyr
	Thr	Met	Leu	Ala	Asp	Gly	325	Ile	Pro	Ile	Ile	330	Tyr	Gln	Gly	Gln	Glu	Gln
	His	Phe	His	Gly	Ala	Thr	340	Asp	Pro	Tyr	Asp	345	Arg	Glu	Pro	Ile	Trp	Pro
	Thr	Gly	Tyr	Asp	Thr	Thr	355	Ser	Pro	Leu	Tyr	360	Val	Leu	Val	Lys	Gln	Leu
	Asn	Ala	Ile	Arg	Ser	Leu	370	Ala	Ile	Ala	Arg	375	Ser	Asp	Thr	Tyr	Ala	Thr
385	Tyr	Gln	Thr	Gln	Ile	Ala	385	Tyr	Ser	Asp	Pro	390	His	Asn	Ile	Ala	Phe	Arg
	Arg	Gly	Asp	Glu	Lys	Cys	405	Met	Ser	Leu	Met	410	Val	Leu	Asn	Asn	Ile	Gly
	Glu	Ser	Ala	Glu	Asp	Tyr	420	Thr	Val	Glu	Ile	425	Glu	Asn	Val	Gly	Phe	Glu
	Ala	Gly	Ser	Ile	Val	Thr	435	Val	Leu	Ser	Cys	440	Arg	Asn	Val	Met	Val	Val
	Asp	Glu	Tyr	Gly	Gly	Met	450	Glu	Val	Pro	Phe	455	Val	Ser	Gly	Leu	Pro	Ser
465	Val	Ser	Leu	Ile	Leu	Ser	465	Met	Arg	Leu	Phe	470	Lys	Ile				
							485					490						

<210> 500
 <211> 2236
 <212> DNA
 <213> Fungi

<220>
 <221> CDS
 <222> (1)...(63)
 <223> Exon

<221> CDS
 <222> (114)...(207)

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<223> Exon
<221> CDS
<222> (264)...(287)
<223> Exon

<221> CDS
<222> (341)...(378)
<223> Exon

<221> CDS
<222> (435)...(560)
<223> Exon

<221> CDS
<222> (623)...(1521)
<223> Exon

<221> CDS
<222> (1571)...(1973)
<223> Exon

<221> CDS
<222> (2051)...(2236)
<223> Exon

<400> 500
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Met Ser Gln Val Ser Gly Lys Val Pro Asp Arg Lys Trp Trp Lys Glu
 1               5               10               15

gct att gtc tac cag gtttgaacta cttttctctt gtcactgcat gcgacttata      103
Ala Ile Val Tyr Gln
                20

cggcggcctag atc tat ccc gcg tcg ttt ctt gat acc gat tct aat ggc      152
            Ile Tyr Pro Ala Ser Phe Leu Asp Thr Asp Ser Asn Gly
                25               30

gtc ggg aac atc aat ggg atc aca gct aag ctg gac tac ctc aag gaa      200
Val Gly Asn Ile Asn Gly Ile Thr Ala Lys Leu Asp Tyr Leu Lys Glu
 35               40               45               50

ttg ggc g gtaagaactg acctccctcc tcctatgctc atgatctaac gcgtgtgtgt      257
Leu Gly

tcacag tg gac att ctc tgg att tca ccc g gtaagctcta tcccagccgt      307
      Val Asp Ile Leu Trp Ile Ser Pro
          55               60

agcgacgcgc ttccagtcta acacgcgctc tag tt tac gaa agc ccc cag aag      360
                        Val Tyr Glu Ser Pro Gln Lys
                              65

gac atg gga tac gac att gtacgacacc catctcttta taaaaattga      408
Asp Met Gly Tyr Asp Ile
          70

tgtagcaccc actgaccttg tcgcag tcg aat tac cgc gat atc cat cgc ccc      461
                        Ser Asn Tyr Arg Asp Ile His Arg Pro
                              75               80

tac ggc acc atg caa gac gtg cac cac ttg atc aac gag ctc aaa gca      509
Tyr Gly Thr Met Gln Asp Val His His Leu Ile Asn Glu Leu Lys Ala
          85               90               95

cgc gac atg aaa ctg atc atg gat ctg gtg gtc aat cac acc tcg acc      557

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Arg Asp Met Lys Leu Ile Met Asp Leu Val Val Asn His Thr Ser Thr
100 105 110

gag gtaagatagc ctttcgaact ctgtatcctg tttaacgtga ctgatttcac 610
Glu
115

ctgacgctat ag cat ccc tgg ttc atc gac tca gcc tcg tcc gca gta agc 661
His Pro Trp Phe Ile Asp Ser Ala Ser Ser Ala Val Ser
120 125

gcg cga cga gac tgg tac atc tgg cgc aag cca aag tac gac gag cag 709
Ala Arg Arg Asp Trp Tyr Ile Trp Arg Lys Pro Lys Tyr Asp Glu Gln
130 135 140

ggc aat cgg cag cca ccg aat aac tgg tgc agc ctc ttt gac gag act 757
Gly Asn Arg Gln Pro Pro Asn Asn Trp Cys Ser Leu Phe Asp Glu Thr
145 150 155 160

gaa tca gcc tgg acc tac gac gct aag acg gat gag tac tat ctc tcg 805
Glu Ser Ala Trp Thr Tyr Asp Ala Lys Thr Asp Glu Tyr Tyr Leu Ser
165 170 175

ctg ttt agt ccg ttc cag gca gat ctt aac tgg gag acc cca ttc gtc 853
Leu Phe Ser Pro Phe Gln Ala Asp Leu Asn Trp Glu Thr Pro Phe Val
180 185 190

aga gag gaa gtc tgc gac atc ctg cgc ttc tgg ctc gac aag ggc gtc 901
Arg Glu Glu Val Cys Asp Ile Leu Arg Phe Trp Leu Asp Lys Gly Val
195 200 205

tct ggt ttc cgg atg gac gtc atc aac ctc atc tcc aag gac cag agg 949
Ser Gly Phe Arg Met Asp Val Ile Asn Leu Ile Ser Lys Asp Gln Arg
210 215 220

ttc cct gac gca gag atc cgc cac ccg ggc agg aag tac cag ccg ggt 997
Phe Pro Asp Ala Glu Ile Arg His Pro Gly Arg Lys Tyr Gln Pro Gly
225 230 235 240

gaa tgc tac ttc gcc aac ggg gtc cga ctc atg gac tat ctc cag gag 1045
Glu Cys Tyr Phe Ala Asn Gly Val Arg Leu Met Asp Tyr Leu Gln Glu
245 250 255

atg aag acc gcc gta ttc tca aag tat gac gcc gtc acg gtc gcc gag 1093
Met Lys Thr Ala Val Phe Ser Lys Tyr Asp Ala Val Thr Val Gly Glu
260 265 270

atg ccg tac ctt gaa gat gaa gag cag aga ctc agg atg gtc gcg gct 1141
Met Pro Tyr Leu Glu Asp Glu Glu Gln Arg Leu Arg Met Val Ala Ala
275 280 285

gaa gaa ggc gtg ctg aac atg atc ttc act ttt gag atg att ggc ctc 1189
Glu Glu Gly Val Leu Asn Met Ile Phe Thr Phe Glu Met Ile Gly Leu
290 295 300

gac atc gtc ccc gaa aaa ggc cgc ttc agc aat aaa gct tgg aag gtc 1237
Asp Ile Val Pro Glu Lys Gly Arg Phe Ser Asn Lys Ala Trp Lys Val
305 310 315 320

tgt gag ctg aag gac ata gtt gcc aag gcc tgc aag ctc gtt aac cag 1285
Cys Glu Leu Lys Asp Ile Val Ala Lys Ala Cys Lys Leu Val Asn Gln
325 330 335

gac ggg tgg cat acg ctc ttc tgc gag aat cat gac cag ccg cgt tcc 1333
Asp Gly Trp His Thr Leu Phe Cys Glu Asn His Asp Gln Pro Arg Ser
340 345 350

gtc acg cgt ttc tgc gac gac tcg gat gag cat cgt gtc gca ggc acc 1381

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Val	Thr	Arg 355	Phe	Cys	Asp	Asp	Ser 360	Asp	Glu	His	Arg	Val 365	Ala	Gly	Thr	
aag Lys	ctc Leu 370	ctc Leu	tcc Ser	atc Ile	atg Met	cag Gln 375	aca Thr	agc Ser	ctt Leu	cct Pro	ggt Gly 380	acc Thr	ctg Leu	tat Tyr	gtc Val	1429
tac Tyr 385	cag Gln	ggc Gly	gaa Glu	gag Glu	ctg Leu 390	ggt Gly	atg Met	cga Arg	aac Asn	gtg Val 395	ccc Pro	aag Lys	tcc Ser	tgg Trp	gga Gly 400	1477
ttt Phe	gaa Glu	gaa Glu	tac Tyr	ata Ile 405	gac Asp	att Ile	gag Glu	tct Ser	gtc Val 410	tcc Ser	tac Tyr	atc Ile	aag Lys	aa Asn 415		1521
gtaagtctca		attctcttat		cgactttaac		atctactgac		atgggtatag		t	gtc Val	tgc Cys				1577
gct Ala	caa Gln	ttc Phe 420	cca Pro	gaa Glu	ggg Gly	tct Ser	ccg Pro 425	gaa Glu	aga Arg	aaa Lys	gag Glu	gct Ala 430	aag Lys	aat Asn	ctt Leu	1625
ctg Leu	cgc Arg 435	ctc Leu	aaa Lys	gct Ala	cgT Arg	gat Asp 440	aat Asn	gcc Ala	agg Arg	acc Thr	cca Pro 445	atg Met	cag Gln	tgg Trp	gac Asp	1673
tct Ser 450	aca Thr	tct Ser	aac Asn	ggc Gly	ggc Gly 455	ttc Phe	tgt Cys	ccc Pro	gag Glu	gga Gly 460	gtc Val	aag Lys	ccc Pro	tgg Trp	atg Met 465	1721
cgC Arg	gtc Val	aac Asn	gac Asp	gac Asp 470	tac Tyr	acg Thr	atg Met	gtc Val	aac Asn 475	gca Ala	gct Ala	cta Leu	cag Gln	act Thr 480	tcc Ser	1769
gca Ala	ggc Gly	cag Gln	gca Ala 485	acc Thr	gac Asp	cgC Arg	acc Thr	atg Met 490	ctt Leu	gtg Val	tca Ser	ccc Pro	tat Tyr 495	agg Arg	ttc Phe	1817
tgg Trp	cag Gln	cga Arg 500	agc Ser	atc Ile	cag Gln	ata Ile	cgC Arg 505	aag Lys	aag Lys	ttc Phe	aaa Lys	gac Asp 510	ctg Leu	ctc Leu	gtg Val	1865
tac Tyr	gga Gly 515	gac Asp	ctt Leu	gag Glu	att Ile	att Ile 520	gat Asp	ggc Gly	acg Thr	cac His	gcc Ala 525	aac Asn	gtc Val	ttt Phe	gcg Ala	1913
ttc Phe 530	aag Lys	agg Arg	atc Ile	agc Ser	aac Asn 535	ggc Gly	aga Arg	cat His	acc Thr	atc Ile 540	acg Thr	atc Ile	ctt Leu	aac Asn	ttt Phe 545	1961
agt Ser	aag Lys	gag Glu	gag Glu	gtgagtttca		agttttccgga		agggtgagggg		gtcaggggct						2013
gggctctcgg		gagttatgat		gttttgtcctt		tggcgag		gcc Ala 550	gtg Val	gac Asp	ggg Gly	gga Gly	gat Asp 555			2068
aca Thr	gct Ala	ttt Phe	acc Thr	ttg Leu 560	gga Gly	ggg Gly	ctt Leu	gct Ala	tgg Trp 565	tat Tyr	tgc Cys	ttg Leu	atg Met	gat Asp 570	ctc Leu	2116
tca Ser	ttt Phe	tgg Trp	aat Asn 575	gga Gly	agt Ser	aaa Lys	aga Arg	ttg Leu 580	gta Val	tca Ser	atg Met	gct Ala	acg Thr 585	ata Ile	ttg Leu	2164
tct	tta	tat	agc	cag	tac	aga	atg	cca	cgg	act	gtt	agt	aac	atg	caa	2212

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Ser Leu Tyr Ser Gln Tyr Arg Met Pro Arg Thr Val Ser Asn Met Gln
590 595 600

tgt ctt gtg ata cca tcc agt taa
Cys Leu Val Ile Pro Ser Ser *
605 610

2236

<210> 501
<211> 21
<212> PRT
<213> Fungi

<400> 501
Met Ser Gln Val Ser Gly Lys Val Pro Asp Arg Lys Trp Trp Lys Glu
1 5 10 15
Ala Ile Val Tyr Gln
20

<210> 502
<211> 31
<212> PRT
<213> Fungi

<400> 502
Ile Tyr Pro Ala Ser Phe Leu Asp Thr Asp Ser Asn Gly Val Gly Asn
1 5 10 15
Ile Asn Gly Ile Thr Ala Lys Leu Asp Tyr Leu Lys Glu Leu Gly
20 25 30

<210> 503
<211> 8
<212> PRT
<213> Fungi

<400> 503
Val Asp Ile Leu Trp Ile Ser Pro
1 5

<210> 504
<211> 13
<212> PRT
<213> Fungi

<400> 504
Val Tyr Glu Ser Pro Gln Lys Asp Met Gly Tyr Asp Ile
1 5 10

<210> 505
<211> 42
<212> PRT
<213> Fungi

<400> 505
Ser Asn Tyr Arg Asp Ile His Arg Pro Tyr Gly Thr Met Gln Asp Val
1 5 10 15
His His Leu Ile Asn Glu Leu Lys Ala Arg Asp Met Lys Leu Ile Met
20 25 30
Asp Leu Val Val Asn His Thr Ser Thr Glu
35 40

<210> 506
<211> 300
<212> PRT
<213> Fungi

<400> 506
His Pro Trp Phe Ile Asp Ser Ala Ser Ser Ala Val Ser Ala Arg Arg

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1      5      10      15
Asp Trp Tyr Ile Trp Arg Lys Pro Lys Tyr Asp Glu Gln Gly Asn Arg
20    35    40    45    50    55    60    65    70    75    80
Gln Pro Pro Asn Asn Trp Cys Ser Leu Phe Asp Glu Thr Glu Ser Ala
35    40    45    50    55    60    65    70    75    80
Trp Thr Tyr Asp Ala Lys Thr Asp Glu Tyr Tyr Leu Ser Leu Phe Ser
50    55    60    65    70    75    80
Pro Phe Gln Ala Asp Leu Asn Trp Glu Thr Pro Phe Val Arg Glu Glu
65    70    75    80
Val Cys Asp Ile Leu Arg Phe Trp Leu Asp Lys Gly Val Ser Gly Phe
85    90    95
Arg Met Asp Val Ile Asn Leu Ile Ser Lys Asp Gln Arg Phe Pro Asp
100   105   110
Ala Glu Ile Arg His Pro Gly Arg Lys Tyr Gln Pro Gly Glu Cys Tyr
115   120   125
Phe Ala Asn Gly Val Arg Leu Met Asp Tyr Leu Gln Glu Met Lys Thr
130   135   140
Ala Val Phe Ser Lys Tyr Asp Ala Val Thr Val Gly Glu Met Pro Tyr
145   150   155   160
Leu Glu Asp Glu Glu Gln Arg Leu Arg Met Val Ala Ala Glu Glu Gly
165   170   175
Val Leu Asn Met Ile Phe Thr Phe Glu Met Ile Gly Leu Asp Ile Val
180   185   190
Pro Glu Lys Gly Arg Phe Ser Asn Lys Ala Trp Lys Val Cys Glu Leu
195   200   205
Lys Asp Ile Val Ala Lys Ala Cys Lys Leu Val Asn Gln Asp Gly Trp
210   215   220
His Thr Leu Phe Cys Glu Asn His Asp Gln Pro Arg Ser Val Thr Arg
225   230   235   240
Phe Cys Asp Asp Ser Asp Glu His Arg Val Ala Gly Thr Lys Leu Leu
245   250   255
Ser Ile Met Gln Thr Ser Leu Pro Gly Thr Leu Tyr Val Tyr Gln Gly
260   265   270
Glu Glu Leu Gly Met Arg Asn Val Pro Lys Ser Trp Gly Phe Glu Glu
275   280   285
Tyr Ile Asp Ile Glu Ser Val Ser Tyr Ile Lys Asn
290   295   300

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<210> 507
 <211> 134
 <212> PRT
 <213> Fungi

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<400> 507
Val Cys Ala Gln Phe Pro Glu Gly Ser Pro Glu Arg Lys Glu Ala Lys
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Asn Leu Leu Arg Leu Lys Ala Arg Asp Asn Ala Arg Thr Pro Met Gln
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Trp Asp Ser Thr Ser Asn Gly Gly Phe Cys Pro Glu Gly Val Lys Pro
35    40    45    50    55    60    65    70    75    80
Trp Met Arg Val Asn Asp Asp Tyr Thr Met Val Asn Ala Ala Leu Gln
50    55    60    65    70    75    80
Thr Ser Ala Gly Gln Ala Thr Asp Arg Thr Met Leu Val Ser Pro Tyr
65    70    75    80
Arg Phe Trp Gln Arg Ser Ile Gln Ile Arg Lys Lys Phe Lys Asp Leu
85    90    95
Leu Val Tyr Gly Asp Leu Glu Ile Ile Asp Gly Thr His Ala Asn Val
100   105   110   115
Phe Ala Phe Lys Arg Ile Ser Asn Gly Arg His Thr Ile Thr Ile Leu
115   120   125
Asn phe Ser Lys Glu Glu
130

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<210> 508
 <211> 61
 <212> PRT
 <213> Fungi

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<400> 508
 Ala Val Asp Gly Gly Asp Thr Ala Phe Thr Leu Gly Gly Leu Ala Trp
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 Tyr Cys Leu Met Asp Leu Ser Phe Trp Asn Gly Ser Lys Arg Leu Val
 20 25 30
 Ser Met Ala Thr Ile Leu Ser Leu Tyr Ser Gln Tyr Arg Met Pro Arg
 35 40 45
 Thr Val Ser Asn Met Gln Cys Leu Val Ile Pro Ser Ser
 50 55 60

<210> 509
 <211> 1833
 <212> DNA
 <213> Fungi

<400> 509
 atgagtcaag tttctggaag agtgcccgac aggaagtggg ggaaggaagc tattgtctac 60
 cagatctatc ccgcgtcggt tcttgatacc gattctaata gcgtcgggaa catcaatggg 120
 atcacagcta agctggacta cctcaaggaa ttgggcgtgg acattctctg gatttcaccc 180
 gtttacgaaa gccccagaa ggacatggga tacgacattt cgaattaccg cgatatccat 240
 cgcccctacg gcaccatgca agacgtgcac cacttgatca acgagctcaa agcacgcgac 300
 atgaaactga tcatggatct ggtggtcaat cacacctcga ccgagcatcc ctggttcac 360
 gactcagcct cgtccgcagt aagcgcgcga cgagactggg acatctggcg caagccaaag 420
 tacgacgagc agggcaatcg gcagccaccg aataactggg gcagcctctt tgacgagact 480
 gaatcagcct ggacctacga cgctaagacg gatgagtact atctctcgct gtttagtccg 540
 ttccaggcag atcttaactg ggagacccca ttcgtcagag aggaagtctg cgacatcctg 600
 cgcttctggc tcgacaaggg cgtctctggg ttccggatgg acgtcatcaa cctcatctcc 660
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 <212> PRT
 <213> Fungi

<220>
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 <222> (21)...(443)
 <223> Alpha amylase, catalytic domain

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 20 25 30
 Asn Gly Val Gly Asn Ile Asn Gly Ile Thr Ala Lys Leu Asp Tyr Leu
 35 40 45
 Lys Glu Leu Gly Val Asp Ile Leu Trp Ile Ser Pro Val Tyr Glu Ser
 50 55 60
 Pro Gln Lys Asp Met Gly Tyr Asp Ile Ser Asn Tyr Arg Asp Ile His

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65	Arg	Pro	Tyr	Gly	Thr	70	Met	Gln	Asp	Val	His	75	His	Leu	Ile	Asn	Glu	80	Leu
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	Ser	Thr	Glu	His	Pro	Trp	Phe	Ile	Asp	Ser	Ala	Ser	Ser	Ala	Val	Ser			
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	Ala	Arg	Arg	Asp	Trp	Tyr	Ile	Trp	Arg	Lys	Pro	Lys	Tyr	Asp	Glu	Gln			
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	Gly	Asn	Arg	Gln	Pro	Pro	Asn	Asn	Trp	Cys	Ser	Leu	Phe	Asp	Glu	Thr			
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	Glu	Ser	Ala	Trp	Thr	Tyr	Asp	Ala	Lys	Thr	Asp	Glu	Tyr	Tyr	Leu	Ser			
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	Leu	Phe	Ser	Pro	Phe	Gln	Ala	Asp	Leu	Asn	Trp	Glu	Thr	Pro	Phe	Val			
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	Arg	Glu	Glu	Val	Cys	Asp	Ile	Leu	Arg	Phe	Trp	Leu	Asp	Lys	Gly	Val			
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	Ser	Gly	Phe	Arg	Met	Asp	Val	Ile	Asn	Leu	Ile	Ser	Lys	Asp	Gln	Arg			
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	Asp	Ile	Val	Pro	Glu	Lys	Gly	Arg	Phe	Ser	Asn	Lys	Ala	Trp	Lys	Val			
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	Cys	Glu	Leu	Lys	Asp	Ile	Val	Ala	Lys	Ala	Cys	Lys	Leu	Val	Asn	Gln			
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	Cys	Ala	Gln	Phe	Pro	Glu	Gly	Ser	Pro	Glu	Arg	Lys	Glu	Ala	Lys	Asn			
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	Leu	Leu	Arg	Leu	Lys	Ala	Arg	Asp	Asn	Ala	Arg	Thr	Pro	Met	Gln	Trp			
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	Asp	Ser	Thr	Ser	Asn	Gly	Gly	Phe	Cys	Pro	Glu	Gly	Val	Lys	Pro	Trp			
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	Ala	Phe	Lys	Arg	Ile	Ser	Asn	Gly	Arg	His	Thr	Ile	Thr	Ile	Leu	Asn			
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610

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<211> 1792
<212> DNA
<213> Fungi

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<222> (1)...(300)
<223> Exon

<221> CDS
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<223> Exon

<221> CDS
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<223> Exon

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cga gca gca agt agt gat gac tgg gcg ggg agg tct atc tat cag gtt 96
Arg Ala Ala Ser Ser Asp Asp Trp Ala Gly Arg Ser Ile Tyr Gln Val
20 25 30

atc aca gat aga tat cat cga tcc tca gat tcg gat gct cct tgt aat 144
Ile Thr Asp Arg Tyr His Arg Ser Ser Asp Ser Ala Pro Cys Asn
35 40 45

atc acg aat tat tgt gga ggt acc tgg aat ggc atc acg gag aat ctg 192
Ile Thr Asn Tyr Cys Gly Gly Thr Trp Asn Gly Ile Thr Glu Asn Leu
50 55 60

gat tat atc cag aat atg ggg ttc acg gcc att caa att agt ccc gtc 240
Asp Tyr Ile Gln Asn Met Gly Phe Thr Ala Ile Gln Ile Ser Pro Val
65 70 75 80

aat ttg aac att aac agt acg acg att tac gga caa gct ttc cat gga 288
Asn Leu Asn Ile Asn Ser Thr Thr Ile Tyr Gly Gln Ala Phe His Gly
85 90 95

tat tgg cag act gtaggttttc tcttggcaga gcttcctct atttgcatatc 340
Tyr Trp Gln Thr
100

aattggcttg ttgttggggg agcatgaaga gttctaacat caatactgca g tct ctc 397
Ser Leu

tac gat ctc aat ccg aac ttc ggt tca gca gat gat ttg ttg aaa cta 445
Tyr Asp Leu Asn Pro Asn Phe Gly Ser Ala Asp Asp Leu Leu Lys Leu
105 110 115

agt gca gaa gtt cac aag aga aaa atg tat ctc cta gtt gat gtc gtt 493
Ser Ala Glu Val His Lys Arg Lys Met Tyr Leu Leu Val Asp Val Val
120 125 130

gct agt gag atg gcc gtt gac att gga gat cac aac atg acc gct gga 541
Ala Ser Glu Met Ala Val Asp Ile Gly Asp His Asn Met Thr Ala Gly
135 140 145 150

acc aag atc gat tat tca gct ttc tcc cct gct cca ttc aac gat gaa 589
Thr Lys Ile Asp Tyr Ser Ala Phe Ser Pro Ala Pro Phe Asn Asp Glu
155 160 165

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gaa Glu	tat Tyr	caa Gln 185	gat Asp	tgt Cys	tgg Trp	ttg Leu	gga Gly 190	tac Tyr	acc Thr	gga Gly	gtc Val	gca Ala 195	act Thr	cca Pro	gat Asp	685
atc Ile	aaa Lys 200	aca Thr	gac Asp	aat Asn	aaa Lys	gaa Glu 205	atc Ile	gca Ala	gcc Ala	gca Ala	tta Leu 210	ggc Gly	aca Thr	tgg Trp	att Ile	733
aag Lys 215	gag Glu	ctt Leu	gtc Val	tcc Ser	aat Asn 220	tac Tyr	tct Ser	att Ile	gat Asp	gga Gly 225	att Ile	cga Arg	gtc Val	gat Asp	ggg Gly 230	781
gct Ala	aaa Lys	cag Gln	att Ile	aca Thr 235	tat Tyr	gac Asp	ttc Phe	ttc Phe	caa Gln 240	gat Asp	ttc Phe	gtc Val	gat Asp	gcc Ala 245	gcc Ala	829
ggg Gly	gta Val	tat Tyr	cct Pro 250	ttg Leu	gga Gly	gaa Glu	gta Val	gag Glu 255	gat Asp	ggc Gly	gat Asp	gct Ala	gcc Ala 260	ttt Phe	act Thr	877
tgc Cys	aat Asn	tac Tyr 265	caa Gln	cag Gln	tac Tyr	acc Thr	gag Glu 270	gga Gly	ttg Leu	gag Glu	aac Asn	tat Tyr 275	ccg Pro	gtt Val	tat Tyr	925
tac Tyr	aca Thr 280	acc Thr	att Ile	caa Gln	gcc Ala	ttt Phe 285	acc Thr	gct Ala	gga Gly	aaa Lys	atg Met 290	gct Ala	ggg Gly	ctg Leu	gtc Val	973
gat Asp 295	atg Met	gtc Val	aag Lys	agt Ser	atg Met 300	gga Gly	gct Ala	ttg Leu	tgc Cys	aaa Lys 305	tct Ser	cca Pro	caa Gln	tat Tyr	ctc Leu 310	1021
gcg Ala	aat Asn	ttc Phe	att Ile	gag Glu 315	aat Asn	caa Gln	gat Asp	caa Gln	cca Pro 320	cgt Arg	ttt Phe	gca Ala	tct Ser	tat Tyr 325	aat Asn	1069
gat Asp	gat Asp	gag Glu	act Thr 330	gtaagtttct tctacttcat gttctaatac aaaatcacta												1121
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ggg Gly	atc Ile 345	ccc Pro	aag Lys	tac Tyr	tat Tyr	tac Tyr 350	gga Gly	caa Gln	gaa Glu	caa Gln	cat His 355	ctg Leu	cag Gln	ggc Gly	aac Asn	1219
tac Tyr 360	tct Ser	cca Pro	tac Tyr	aac Asn	cgt Arg 365	caa Gln	gct Ala	ctc Leu	tgg Trp	gaa Glu 370	agt Ser	aag Lys	cca Pro	agc Ser	aca Thr 375	1267
act Thr	act Thr	cct Pro	ctc Leu	tac Tyr 380	act Thr	atg Met	aca Thr	gct Ala	act Thr 385	ctc Leu	aac Asn	acc Thr	ctc Leu	cgt Arg 390	aac Asn	1315
cac His	gcc Ala	atc Ile	tcc Ser 395	ata Ile	aac Asn	tca Ser	aac Asn	tac Tyr 400	atc Ile	gcc Ala	aac Asn	atg Met	tcc Ser	att Ile	ctg Leu	1363
ctc Leu	cac His	aat Asn 410	gac Asp	gat Asp	tcc Ser	aca Thr	tac Tyr 415	gca Ala	aca Thr	cgt Arg	aaa Lys	gga Gly 420	cca Pro	aac Asn	ggg Gly	1411

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gtt	caa	atc	gtc	tcg	gta	ctt	tct	aac	caa	ggg	tcc	aaa	gga	gga	agc	1459
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Tyr	Thr	Leu	Gly	Ile	Asn	Asn	Ala	Ala	Asp	Ala	Gly	Thr	Asn	Met	Thr	
440					445					450					455	
gaa	gta	atc	aca	tgc	aca	tcg	tac	att	gca	agt	gag	aac	gga	acg	ctc	1555
Glu	Val	Ile	Thr	Cys	Thr	Ser	Tyr	Ile	Ala	Ser	Glu	Asn	Gly	Thr	Leu	
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ttc	att	gag	atg	gac	aaa	gga	gag	cca	aga	gta	ctg	ttc	cct	agc	tac	1603
Phe	Ile	Glu	Met	Asp	Lys	Gly	Glu	Pro	Arg	Val	Leu	Phe	Pro	Ser	Tyr	
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caa	atg	aat	gga	acg	ggc	att	tgt	gga	ttc	gat	gaa	gat	gct	agt	gcg	1651
Gln	Met	Asn	Gly	Thr	Gly	Ile	Cys	Gly	Phe	Asp	Glu	Asp	Ala	Ser	Ala	
		490					495					500				
gga	aca	gca	aca	agc	act	gga	acg	gcg	act	gga	agt	gca	gct	agt	gcg	1699
Gly	Thr	Ala	Thr	Ser	Thr	Gly	Thr	Ala	Thr	Gly	Ser	Ala	Ala	Ser	Ala	
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aca	tca	tcg	aag	aag	gga	gat	gcg	agt	gga	ttg	aga	gtt	gcc	ggg	tgg	1747
Thr	Ser	Ser	Lys	Lys	Gly	Asp	Ala	Ser	Gly	Leu	Arg	Val	Ala	Gly	Trp	
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gtg	ttc	gcg	ggg	gcg	atg	ggg	gcg	tct	gct	ttg	gct	ttc	atg	tag		1792
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 <213> Fungi

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 35 40 45
 Ile Thr Asn Tyr Cys Gly Gly Thr Trp Asn Gly Ile Thr Glu Asn Leu
 50 55 60
 Asp Tyr Ile Gln Asn Met Gly Phe Thr Ala Ile Gln Ile Ser Pro Val
 65 70 75 80
 Asn Leu Asn Ile Asn Ser Thr Thr Ile Tyr Gly Gln Ala Phe His Gly
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 Tyr Trp Gln Thr
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<210> 513
 <211> 230
 <212> PRT
 <213> Fungi

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 Lys Leu Ser Ala Glu Val His Lys Arg Lys Met Tyr Leu Leu Val Asp
 20 25 30
 Val Val Ala Ser Glu Met Ala Val Asp Ile Gly Asp His Asn Met Thr
 35 40 45
 Ala Gly Thr Lys Ile Asp Tyr Ser Ala Phe Ser Pro Ala Pro Phe Asn

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Val Thr Glu Tyr Gln Asp Cys Trp Leu Gly Tyr Thr Gly Val Ala Thr
Pro Asp Ile Lys 85 Thr Asp Asn Lys Glu 90 Ile Ala Ala Ala Leu Gly Thr
Trp Ile Lys 100 Glu Leu Val Ser Asn 105 Tyr Ser Ile Asp Gly 110 Ile Arg Val
Asp Gly Ala Lys Gln Ile Thr 120 Tyr Asp Phe Phe Gln Asp Phe Val Asp
Ala 130 Ala Gly Val Tyr Pro 135 Leu Gly Glu Val Glu 140 Asp Gly Asp Ala Ala
145 Phe Thr Cys Asn Tyr 150 Gln Gln Tyr Thr Glu 155 Gly Leu Glu Asn Tyr 160
Val Tyr Tyr Thr Thr Ile Gln Ala Phe 170 Thr Ala Gly Lys Met Ala Gly
Leu Val Asp 180 Met Val Lys Ser Met 185 Gly Ala Leu Cys Lys Ser Pro Gln
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Leu Tyr Thr Met Thr Ala 50 Thr 55 Leu Asn Thr Leu Arg Asn His Ala Ile
Ser 65 Ile Asn Ser Asn Tyr 70 Ile Ala Asn Met Ser 75 Ile Leu Leu His Asn 80
Asp Asp Ser Thr Tyr 85 Ala Thr Arg Lys Gly 90 Pro Asn Gly Val Gln Ile
Val Ser Val 100 Ser Asn Gln Gly Ser 105 Lys Gly Gly Ser Tyr 110 Thr Leu
Gly Ile Asn 115 Asn Ala Ala Asp Ala 120 Gly Thr Asn Met Thr 125 Glu Val Ile
Thr Cys 130 Thr Ser Tyr Ile Ala 135 Ser Glu Asn Gly Thr 140 Leu Phe Ile Glu
Met 145 Asp Lys Gly Glu Pro 150 Arg Val Leu Phe Pro 155 Ser Tyr Gln Met Asn 160
Gly Thr Gly Ile Cys 165 Gly Phe Asp Glu Asp 170 Ala Ser Ala Gly Thr Ala
Thr Ser Thr 180 Gly Thr Ala Thr Gly Ser 185 Ala Ala Ser Ala Thr 190 Ser Ser
Lys Lys Gly 195 Asp Ala Ser Gly Leu 200 Arg Val Ala Gly Trp 205 Val Phe Ala
Gly Ala Met Gly Ala Ser Ala 215 Leu Ala Phe Met
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 <211> 1650
 <212> DNA
 <213> Fungi

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gaagttcaca agagaaaaat gtatctccta gttgatgtcg ttgctagtga gatggcgggt 420
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atgggtgcgt ctgctttggc tttcatgtag 1650

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 <212> PRT
 <213> Fungi

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 <222> (1)...(21)

<221> DOMAIN
 <222> (31)...(390)
 <223> Alpha amylase, catalytic domain

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Ile Thr Asp Arg Tyr His Arg Ser Ser Asp Ser Asp Ala Pro Cys Asn
          35          40          45
Ile Thr Asn Tyr Cys Gly Gly Thr Trp Asn Gly Ile Thr Glu Asn Leu
          50          55          60
Asp Tyr Ile Gln Asn Met Gly Phe Thr Ala Ile Gln Ile Ser Pro Val
          65          70          75          80
Asn Leu Asn Ile Asn Ser Thr Thr Ile Tyr Gly Gln Ala Phe His Gly
          85          90          95
Tyr Trp Gln Thr Ser Leu Tyr Asp Leu Asn Pro Asn Phe Gly Ser Ala
          100          105          110
Asp Asp Leu Leu Lys Leu Ser Ala Glu Val His Lys Arg Lys Met Tyr
          115          120          125
Leu Leu Val Asp Val Val Ala Ser Glu Met Ala Val Asp Ile Gly Asp
          130          135          140
His Asn Met Thr Ala Gly Thr Lys Ile Asp Tyr Ser Ala Phe Ser Pro
          145          150          155          160
Ala Pro Phe Asn Asp Glu Ser Ser Tyr Asn Pro Tyr Cys Pro Ile Ile
          165          170          175
Asp Trp Gln Asn Val Thr Glu Tyr Gln Asp Cys Trp Leu Gly Tyr Thr
          180          185          190
Gly Val Ala Thr Pro Asp Ile Lys Thr Asp Asn Lys Glu Ile Ala Ala
          195          200          205
Ala Leu Gly Thr Trp Ile Lys Glu Leu Val Ser Asn Tyr Ser Ile Asp
          210          215          220

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1001827087_1.txt

Gly Ile Arg Val Asp Gly Ala Lys Gln Ile Thr Tyr Asp Phe Phe Gln
 225 230 235 240
 Asp Phe Val Asp Ala Gly Val Tyr Pro Leu Gly Glu Val Glu Asp
 245 250 255
 Gly Asp Ala Ala Phe Thr Cys Asn Tyr Gln Gln Tyr Thr Glu Gly Leu
 260 265 270
 Glu Asn Tyr Pro Val Tyr Tyr Thr Thr Ile Gln Ala Phe Thr Ala Gly
 275 280 285
 Lys Met Ala Gly Leu Val Asp Met Val Lys Ser Met Gly Ala Leu Cys
 290 295 300
 Lys Ser Pro Gln Tyr Leu Ala Asn Phe Ile Glu Asn Gln Asp Gln Pro
 305 310 315 320
 Arg Phe Ala Ser Tyr Asn Asp Asp Glu Thr Leu Ala Ala Asn Ala Met
 325 330 335
 Ala Phe Thr Ile Leu Ala Asp Gly Ile Pro Lys Tyr Tyr Tyr Gly Gln
 340 345 350
 Glu Gln His Leu Gln Gly Asn Tyr Ser Pro Tyr Asn Arg Gln Ala Leu
 355 360 365
 Trp Glu Ser Lys Pro Ser Thr Thr Thr Pro Leu Tyr Thr Met Thr Ala
 370 375 380
 Thr Leu Asn Thr Leu Arg Asn His Ala Ile Ser Ile Asn Ser Asn Tyr
 385 390 395 400
 Ile Ala Asn Met Ser Ile Leu Leu His Asn Asp Asp Ser Thr Tyr Ala
 405 410 415
 Thr Arg Lys Gly Pro Asn Gly Val Gln Ile Val Ser Val Leu Ser Asn
 420 425 430
 Gln Gly Ser Lys Gly Gly Ser Tyr Thr Leu Gly Ile Asn Asn Ala Ala
 435 440 445
 Asp Ala Gly Thr Asn Met Thr Glu Val Ile Thr Cys Thr Ser Tyr Ile
 450 455 460
 Ala Ser Glu Asn Gly Thr Leu Phe Ile Glu Met Asp Lys Gly Glu Pro
 465 470 475 480
 Arg Val Leu Phe Pro Ser Tyr Gln Met Asn Gly Thr Gly Ile Cys Gly
 485 490 495
 Phe Asp Glu Asp Ala Ser Ala Gly Thr Ala Thr Ser Thr Gly Thr Ala
 500 505 510
 Thr Gly Ser Ala Ala Ser Ala Thr Ser Ser Lys Lys Gly Asp Ala Ser
 515 520 525
 Gly Leu Arg Val Ala Gly Trp Val Phe Ala Gly Ala Met Gly Ala Ser
 530 535 540
 Ala Leu Ala Phe Met
 545

<210> 517
 <211> 1758
 <212> DNA
 <213> Fungi

<220>
 <221> CDS
 <222> (1)...(1758)
 <223> Exon

<400> 517
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 Met Asn Met Asn Ile Phe Leu Leu Ile Ile Ser Leu Ala Phe Phe Ser
 1 5 10 15
 acc gtc aat tgt tac acc atg tcc aat gcc gag gcc tgg aaa agg cgt 96
 Thr Val Asn Cys Tyr Thr Met Ser Asn Ala Glu Ala Trp Lys Arg Arg
 20 25 30
 tcc ata tac caa gtg ctc acc gac aga ttt gcc ata gaa gat gag aat 144
 Ser Ile Tyr Gln Val Leu Thr Asp Arg Phe Ala Ile Glu Asp Glu Asn
 35 40 45
 aat cct agt aga ggc cca gat cat ggg atg aga gag tat tgt ggt ggg 192
 Asn Pro Ser Arg Gly Pro Asp His Gly Met Arg Glu Tyr Cys Gly Gly

1001827087_1.txt
60

50	aca Thr 65	tgg Trp	agg Arg	ggg Gly	ata Ile	ata Ile 70	tct Ser	aaa Lys	ctt Leu	gat Asp	tat Tyr 75	att Ile	caa Gln	gga Gly	atg Met	ggg Gly 80	240
	ttt Phe	gat Asp	gca Ala	gtt Val	tgg Trp 85	att Ile	tcg Ser	ccc Pro	gtt Val	tgt Cys 90	ttc Phe	ttc Phe	agc Ser	ctc Leu	acc Thr 95	agc Ser	288
	gtt Val	ttg Leu	att Ile	gca Ala 100	ttt Phe	act Thr	cac His	aat Asn	tcc Ser 105	ata Ile	cag Gln	atc Ile	acc Thr	aag Lys 110	aac Asn	atc Ile	336
	gaa Glu	gaa Glu	ttg Leu 115	act Thr	gaa Glu	tat Tyr	gga Gly	act Thr 120	gcc Ala	tat Tyr	cac His	ggg Gly	tat Tyr 125	tgg Trp	cca Pro	gag Glu	384
	gat Asp	att Ile 130	tat Tyr	tcg Ser	gta Val	aat Asn	cca Pro 135	cat His	ttt Phe	gga Gly	acc Thr	gct Ala 140	gat Asp	gat Asp	ttg Leu	ctc Leu	432
	gct Ala 145	cta Leu	tcg Ser	gat Asp	gcc Ala 150	ttg Leu 150	cac His	gct Ala	cgg Arg	ggg Gly	atg Met 155	ttt Phe	ttg Leu	atg Met	gtt Val	gat Asp 160	480
	ata Ile	gtt Val	gtc Val	aat Asn	cat His 165	atg Met	ggg Gly	tcg Ser	cct Pro	cct Pro 170	aca Thr	gtg Val	gac Asp	tat Tyr	tct Ser 175	aga Arg	528
	tac Tyr	gta Val	cct Pro	ttc Phe 180	aat Asn	gat Asp	tcg Ser	tca Ser	tac Tyr 185	tat Tyr	cat His	cca Pro	cag Gln	tca Ser 190	ttc Phe	gtc Val	576
	acg Thr	aac Asn	tac Tyr 195	gaa Glu	aat Asn	cag Gln	cag Gln	gag Glu 200	acc Thr	gag Glu	gaa Glu	gga Gly	tgg Trp 205	ttg Leu	ggg Gly	gat Asp	624
	gaa Glu 210	cat His	gtg Val	cca Pro	tta Leu	ccc Pro	gat Asp 215	cta Leu	aat Asn	acg Thr	gag Glu	gat Asp 220	cct Pro	act Thr	gtg Val	gta Val	672
	ctc Leu 225	acc Thr	ctc Leu	caa Gln	acc Thr	tgg Trp 230	atc Ile	aca Thr	aaa Lys	ctt Leu	gta Val 235	cag Gln	agg Arg	ttc Phe	aaa Lys	atc Ile 240	720
	gat Asp	ggg Gly	ttg Leu	aga Arg	att Ile 245	gac Asp	acc Thr	ttc Phe	aaa Lys	cat His 250	gtc Val	agg Arg	aag Lys	aat Asn	ttt Phe 255	tgg Trp	768
	ccg Pro	gat Asp	ttc Phe	gtt Val 260	cgg Arg	gct Ala	gcg Ala	ggg Gly	gtc Val 265	tgg Trp	tcg Ser	gtg Val	ggg Gly	gag Glu 270	gtg Val	ttg Leu	816
	tca Ser	ggg Gly	gac Asp 275	ccc Pro	gac Asp	tac Tyr	ttg Leu	agc Ser 280	tca Ser	tat Tyr	caa Gln	cct Pro	tat Tyr 285	gtt Val	gga Gly	ggc Gly	864
	ctt Leu	cta Leu 290	gac Asp	tat Tyr	gga Gly	aca Thr	tat Tyr 295	tat Tyr	cct Pro	ttg Leu	aag Lys	aga Arg 300	gcc Ala	ttt Phe	cat His	agg Arg	912
	gac Asp 305	ggg Gly	gga Gly	agt Ser	atg Met	tat Tyr 310	gag Glu	ttg Leu	acc Thr	aac Asn	ctc Leu 315	ctc Leu	act Thr	cct Pro	caa Gln	tat Tyr 320	960
	cga Arg	tca Ser	aaa Lys	ttc Phe	cgc Arg	gac Asp	atg Met	cag Gln	caa Gln	atg Met	tcg Ser	act Thr	ttt Phe	atg Met	gag Glu	aat Asn	1008

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                                1001827087_1.txt
                                330
                                335
325
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His Asp Met Pro Arg Phe Thr Asn Asp Thr Asn Ala Asp Leu Ala Val
340
gta aag aat gct atg gca tgg aat cta ctc acc gat ggt gtg cca att 1104
Val Lys Asn Ala Met Ala Trp Asn Leu Leu Thr Asp Gly Val Pro Ile
355
att tac tat gga caa gaa caa tct tat tct gga gat ggt gat cct ggc 1152
Ile Tyr Tyr Gly Gln Glu Gln Ser Tyr Ser Gly Asp Gly Asp Pro Gly
370
tca cga gag ctg atg tgg aca tcc aat tat aac atc aca cct cta tac 1200
Ser Arg Glu Leu Met Trp Thr Ser Asn Tyr Asn Ile Thr Pro Leu Tyr
385
caa tat atc gca cgt ttg aat cag atc aga aag ctg ggt tgg gat gta 1248
Gln Tyr Ile Ala Arg Leu Asn Gln Ile Arg Lys Leu Gly Trp Asp Val
405
gga ttt ggt aca cat cta acc aca cgt ctt tac gtt gag gat cat gtc 1296
Gly Phe Gly Thr His Leu Thr Thr Arg Leu Tyr Val Glu Asp His Val
420
gct gtt acg cag aag ggc ccc gtg ctc atg gtg ctg acc aac cta ggt 1344
Ala Val Thr Gln Lys Gly Pro Val Leu Met Val Leu Thr Asn Leu Gly
435
agc cag gcc aga tcc aaa act atc tct aca cct acc atg ttc aaa aac 1392
Ser Gln Ala Arg Ser Lys Thr Ile Ser Thr Pro Thr Met Phe Lys Asn
450
ggt acc ata ata gtc gac att ctc aca gga gaa tcc ttc aga gtc aaa 1440
Gly Thr Ile Ile Val Asp Ile Leu Thr Gly Glu Ser Phe Arg Val Lys
465
cga tcg aca aac atc acc ata gta tct gga gaa cct cga att ttt ctt 1488
Arg Ser Thr Asn Ile Thr Ile Val Ser Gly Glu Pro Arg Ile Phe Leu
485
cct ctt ggt cta gca gag aat atc tgt gaa cag atc tta cca cca gcg 1536
Pro Leu Gly Leu Ala Glu Asn Ile Cys Glu Gln Ile Leu Pro Pro Ala
500
cag agt gcc ata tcc aaa cta tta tct aga atg ttt ggt gcc tcc tct 1584
Gln Ser Ala Ile Ser Lys Leu Leu Ser Arg Met Phe Gly Ala Ser Ser
515
tcc ttg aaa act gcc gcc gaa atc acc gat tgg cca tat ggt gaa ccg 1632
Ser Leu Lys Thr Ala Ala Glu Ile Thr Asp Trp Pro Tyr Gly Glu Pro
530
gtc aac agc tca gat att ata ctc gaa gaa aaa gat cca gtc gcg gag 1680
Val Asn Ser Ser Asp Ile Ile Leu Glu Glu Lys Asp Pro Val Ala Glu
545
cag gca gga gcc aag aca ttg atg ccg agc gtt caa gag agt tca tcc 1728
Gln Ala Gly Ala Lys Thr Leu Met Pro Ser Val Gln Glu Ser Ser Ser
565
ccg tgg tct ata ttc tat acg gcc aga taa 1758
Pro Trp Ser Ile Phe Tyr Thr Ala Arg
580

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<210> 518

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<211> 585
<212> PRT
<213> Fungi

<220>
<221> DOMAIN
<222> (36)...(410)
<223> Alpha amylase, catalytic domain

<221> SIGNAL
<222> (1)...(26)

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Thr Val Asn Cys Tyr Thr Met Ser Asn Ala Glu Ala Trp Lys Arg Arg
20 25 30
Ser Ile Tyr Gln Val Leu Thr Asp Arg Phe Ala Ile Glu Asp Glu Asn
35 40 45
Asn Pro Ser Arg Gly Pro Asp His Gly Met Arg Glu Tyr Cys Gly Gly
50 55 60
Thr Trp Arg Gly Ile Ile Ser Lys Leu Asp Tyr Ile Gln Gly Met Gly
65 70 75 80
Phe Asp Ala Val Trp Ile Ser Pro Val Cys Phe Phe Ser Leu Thr Ser
85 90 95
Val Leu Ile Ala Phe Thr His Asn Ser Ile Gln Ile Thr Lys Asn Ile
100 105 110
Glu Glu Leu Thr Glu Tyr Gly Thr Ala Tyr His Gly Tyr Trp Pro Glu
115 120 125
Asp Ile Tyr Ser Val Asn Pro His Phe Gly Thr Ala Asp Asp Leu Leu
130 135 140
Ala Leu Ser Asp Ala Leu His Ala Arg Gly Met Phe Leu Met Val Asp
145 150 155 160
Ile Val Val Asn His Met Gly Ser Pro Pro Thr Val Asp Tyr Ser Arg
165 170 175 180
Tyr Val Pro Phe Asn Asp Ser Ser Tyr Tyr His Pro Gln Ser Phe Val
185 190
Thr Asn Tyr Glu Asn Gln Gln Glu Thr Glu Glu Gly Trp Leu Gly Asp
195 200 205
Glu His Val Pro Leu Pro Asp Leu Asn Thr Glu Asp Pro Thr Val Val
210 215 220
Leu Thr Leu Gln Thr Trp Ile Thr Lys Leu Val Gln Arg Phe Lys Ile
225 230 235 240
Asp Gly Leu Arg Ile Asp Thr Phe Lys His Val Arg Lys Asn Phe Trp
245 250 255
Pro Asp Phe Val Arg Ala Ala Gly Val Trp Ser Val Gly Glu Val Leu
260 265 270
Ser Gly Asp Pro Asp Tyr Leu Ser Ser Tyr Gln Pro Tyr Val Gly Gly
275 280 285
Leu Leu Asp Tyr Gly Thr Tyr Tyr Pro Leu Lys Arg Ala Phe His Arg
290 295 300
Asp Gly Gly Ser Met Tyr Glu Leu Thr Asn Leu Leu Thr Pro Gln Tyr
305 310 315 320
Arg Ser Lys Phe Arg Asp Met Gln Gln Met Ser Thr Phe Met Glu Asn
325 330 335
His Asp Met Pro Arg Phe Thr Asn Asp Thr Asn Ala Asp Leu Ala Val
340 345 350
Val Lys Asn Ala Met Ala Trp Asn Leu Leu Thr Asp Gly Val Pro Ile
355 360 365
Ile Tyr Tyr Gly Gln Glu Gln Ser Tyr Ser Gly Asp Gly Asp Pro Gly
370 375 380
Ser Arg Glu Leu Met Trp Thr Ser Asn Tyr Asn Ile Thr Pro Leu Tyr
385 390 395 400
Gln Tyr Ile Ala Arg Leu Asn Gln Ile Arg Lys Leu Gly Trp Asp Val
405 410 415
Gly Phe Gly Thr His Leu Thr Thr Arg Leu Tyr Val Glu Asp His Val
420 425 430
Ala Val Thr Gln Lys Gly Pro Val Leu Met Val Leu Thr Asn Leu Gly

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      435      440      445
Ser Gln Ala Arg Ser Lys Thr Ile Ser Thr Pro Thr Met Phe Lys Asn
450      455      460
Gly Thr Ile Ile Val Asp Ile Leu Thr Gly Glu Ser Phe Arg Val Lys
465      470      475
Arg Ser Thr Asn Ile Thr Ile Val Ser Gly Glu Pro Arg Ile Phe Leu
      485      490      495
Pro Leu Gly Leu Ala Glu Asn Ile Cys Glu Gln Ile Leu Pro Pro Ala
500      505      510
Gln Ser Ala Ile Ser Lys Leu Leu Ser Arg Met Phe Gly Ala Ser Ser
515      520      525
Ser Leu Lys Thr Ala Ala Glu Ile Thr Asp Trp Pro Tyr Gly Glu Pro
530      535      540
Val Asn Ser Ser Asp Ile Ile Leu Glu Glu Lys Asp Pro Val Ala Glu
545      550      555
Gln Ala Gly Ala Lys Thr Leu Met Pro Ser Val Gln Glu Ser Ser Ser
565      570      575
Pro Trp Ser Ile Phe Tyr Thr Ala Arg
580      585

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<210> 519
 <211> 1843
 <212> DNA
 <213> Fungi

<220>
 <221> CDS
 <222> (1)...(887)
 <223> Exon

<221> CDS
 <222> (943)...(1843)
 <223> Exon

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atg cag cct gtc atc cca aac gtc gag cga gcc tcg tgg cgg gaa gcc      48
Met Gln Pro Val Ile Pro Asn Val Glu Arg Ala Ser Trp Arg Glu Ala
1      5      10

tct gta tat cag atc tac cca cct tct ttt aaa gat acc aat ggg gat      96
Ser Val Tyr Gln Ile Tyr Pro Pro Ser Phe Lys Asp Thr Asn Gly Asp
20      25      30

ggt ata gga gat att caa gga att atc tca gag ctt gac tat atc aag      144
Gly Ile Gly Asp Ile Gln Gly Ile Ile Ser Glu Leu Asp Tyr Ile Lys
35      40      45

tct ctt ggc gtt gat atg gta tgg ctg tct ccc att ttg gca tct ccg      192
Ser Leu Gly Val Asp Met Val Trp Leu Ser Pro Ile Leu Ala Ser Pro
50      55      60

cag gtt gat atg ggt tac gat atc tct gac tat aag aat atc tat ccg      240
Gln Val Asp Met Gly Tyr Asp Ile Ser Asp Tyr Lys Asn Ile Tyr Pro
65      70      75

cct tat ggt acc atg gcc gat cat gat gct tta atc aaa ggc ctc cat      288
Pro Tyr Gly Thr Met Ala Asp His Asp Ala Leu Ile Lys Gly Leu His
85      90      95

gac cgc ggt ttg aag tat atc ctt gac ctt gtt gtg aac cat aca tcg      336
Asp Arg Gly Leu Lys Tyr Ile Leu Asp Leu Val Val Asn His Thr Ser
100      105      110

gat caa cat cct tgg ttc aaa gag tcc aag tct tca aag aca agc aga      384
Asp Gln His Pro Trp Phe Lys Glu Ser Lys Ser Ser Lys Thr Ser Arg
115      120      125

tat cgc gac tgg tat ttc tgg cgg ccg gct cgc tgg gat ccc gaa acc      432

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Tyr	Arg	Asp	Trp	Tyr	Phe	Trp	Arg	Pro	Ala	Arg	Trp	Asp	Pro	Glu	Thr		
130						135					140						
ggt	aaa	cgg	atg	cca	cct	aac	aat	tgg	gaa	tcc	ttt	ttt	agt	aca	agc	480	
Gly	Lys	Arg	Met	Pro	Pro	Asn	Asn	Trp	Glu	Ser	Phe	Phe	Ser	Thr	Ser	160	
145					150					155							
gct	tgg	act	tgg	gac	gct	acc	aca	caa	gaa	tat	tat	ctt	cac	ctt	tgg	528	
Ala	Trp	Thr	Trp	Asp	Ala	Thr	Thr	Gln	Glu	Tyr	Tyr	Leu	His	Leu	Trp	175	
				165					170					175			
tca	tct	gga	cag	cct	gac	ttg	aac	tgg	gaa	aat	ccc	gat	gtc	gtc	aat	576	
Ser	Ser	Gly	Gln	Pro	Asp	Leu	Asn	Trp	Glu	Asn	Pro	Asp	Val	Val	Asn	190	
			180					185					190				
gct	gtg	cac	gat	atc	gta	aga	ttc	tgg	ctt	gat	cgt	ggc	ggt	gat	ggg	624	
Ala	Val	His	Asp	Ile	Val	Arg	Phe	Trp	Leu	Asp	Arg	Gly	Val	Asp	Gly	205	
		195					200					205					
ttt	cga	atg	gat	gtc	att	aac	tat	att	tcg	aaa	aca	cct	ggc	ctg	ccc	672	
Phe	Arg	Met	Asp	Val	Ile	Asn	Tyr	Ile	Ser	Lys	Thr	Pro	Gly	Leu	Pro	220	
	210					215					220						
gac	gca	aag	atc	aag	aaa	ccg	gga	ttt	ttt	cag	aag	ccc	acc	gag	cat	720	
Asp	Ala	Lys	Ile	Lys	Lys	Pro	Gly	Phe	Phe	Gln	Lys	Pro	Thr	Glu	His	240	
					230					235					240		
tgt	gct	tgt	gga	cca	cgg	ctg	cac	gaa	tac	cta	cgt	ggg	att	gga	tcg	768	
Cys	Ala	Cys	Gly	Pro	Arg	Leu	His	Glu	Tyr	Leu	Arg	Gly	Ile	Gly	Ser	255	
				245					250					255			
ata	ctc	cag	gaa	tac	gat	gca	ttt	agt	gtc	ggc	gag	atg	cca	gat	gca	816	
Ile	Leu	Gln	Glu	Tyr	Asp	Ala	Phe	Ser	Val	Gly	Glu	Met	Pro	Asp	Ala	270	
			260					265					270				
gat	ccc	gaa	gaa	gtt	ctc	aaa	gct	gtg	ggc	cag	gat	cgt	gga	gaa	tta	864	
Asp	Pro	Glu	Glu	Val	Leu	Lys	Ala	Val	Gly	Gln	Asp	Arg	Gly	Glu	Leu	285	
		275					280					285					
gct	atg	gcg	ttt	cac	ttt	gac	at	gta	agtcata	acagaatata	gtctttttaca					917	
Ala	Met	Ala	Phe	His	Phe	Asp	Ile									290	
	290					295											
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				Asp	Gly	Leu	Asp	Leu	Gly	Ala	Asn	His				305	
								300									
aga	ttt	gac	ccc	gcc	gtg	ttc	caa	cca	aca	gca	tta	aag	aac	gta	gtc	1018	
Arg	Phe	Asp	Pro	Ala	Val	Phe	Gln	Pro	Thr	Ala	Leu	Lys	Asn	Val	Val	320	
				310					315					320			
aac	aaa	tgg	caa	aca	ttc	atg	gca	tcc	aac	gca	ggt	tgg	aac	gcc	ctg	1066	
Asn	Lys	Trp	Gln	Thr	Phe	Met	Ala	Ser	Asn	Ala	Gly	Trp	Asn	Ala	Leu	335	
			325					330					335				
cat	atg	gag	aac	cac	gac	gag	agc	cga	aca	gtt	agt	cgt	tac	gct	tgc	1114	
His	Met	Glu	Asn	His	Asp	Glu	Ser	Arg	Thr	Val	Ser	Arg	Tyr	Ala	Cys	345	
		340					345					350					
gac	tct	ccc	gcc	atg	cga	aca	ata	tca	gcc	aag	atg	cta	gca	aac	cac	1162	
Asp	Ser	Pro	Ala	Met	Arg	Thr	Ile	Ser	Ala	Lys	Met	Leu	Ala	Asn	His	365	
	355					360					365						
ctt	gcc	ttc	cag	tct	gga	aca	ctt	ttc	atc	tat	cag	gga	cag	gaa	ctt	1210	
Leu	Ala	Phe	Gln	Ser	Gly	Thr	Leu	Phe	Ile	Tyr	Gln	Gly	Gln	Glu	Leu	385	
	370				375					380					385		
gca	atg	gtt	aat	ctt	ccc	cga	gaa	tgg	gga	atg	gag	aaa	tac	aaa	gac	1258	

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Ala	Met	Val	Asn	Leu	Pro	Arg	Glu	Trp	Gly	Met	Glu	Lys	Tyr	Lys	Asp	
				390					395					400		
atc	gaa	tgt	ctg	aat	cac	tgg	gag	ttg	gta	cag	cgt	tgt	tac	cac	gac	1306
Ile	Glu	Cys	Leu	Asn	His	Trp	Glu	Leu	Val	Gln	Arg	Cys	Tyr	His	Asp	
			405					410					415			
gat	gta	aaa	aag	cag	aag	atg	tat	agg	gac	aga	tac	cga	cag	gtc	gga	1354
Asp	Val	Lys	Lys	Gln	Lys	Met	Tyr	Arg	Asp	Arg	Tyr	Arg	Gln	Val	Gly	
		420					425					430				
cgt	gac	aac	gct	cgc	acg	ccc	atg	cag	tgg	aac	tcg	gag	agt	cca	tat	1402
Arg	Asp	Asn	Ala	Arg	Thr	Pro	Met	Gln	Trp	Asn	Ser	Glu	Ser	Pro	Tyr	
	435					440					445					
gct	gga	ttc	gtg	ccg	att	ggg	tct	aag	caa	ggt	gag	ccc	tgg	atg	tct	1450
Ala	Gly	Phe	Val	Pro	Ile	Gly	Ser	Lys	Gln	Val	Glu	Pro	Trp	Met	Ser	
	450				455				460						465	
atc	cac	cct	gac	ttt	ggg	act	tgg	aac	ggt	tcc	acg	atg	tta	gcc	gat	1498
Ile	His	Pro	Asp	Phe	Gly	Thr	Trp	Asn	Val	Ser	Thr	Met	Leu	Ala	Asp	
				470					475					480		
tcg	cag	tcc	agc	ctt	cac	cat	tgg	cgc	cgg	gtg	ctc	aag	ttt	cga	aag	1546
Ser	Gln	Ser	Ser	Leu	His	His	Trp	Arg	Arg	Val	Leu	Lys	Phe	Arg	Lys	
			485					490					495			
cag	cat	tcc	aac	ata	ttt	gtt	tac	gga	gga	ttc	gaa	atg	cta	aat	ata	1594
Gln	His	Ser	Asn	Ile	Phe	Val	Tyr	Gly	Gly	Phe	Glu	Met	Leu	Asn	Ile	
		500					505					510				
gaa	atc	gaa	gaa	gac	gtc	att	gcg	tat	ata	cgc	act	gac	aac	act	act	1642
Glu	Ile	Glu	Glu	Asp	Val	Ile	Ala	Tyr	Ile	Arg	Thr	Asp	Asn	Thr	Thr	
	515					520					525					
ggg	ggg	agc	cta	act	gga	ccg	acg	gag	gcg	ctg	ggt	gtg	acc	agt	ttt	1690
Gly	Gly	Ser	Leu	Thr	Gly	Pro	Thr	Glu	Ala	Leu	Val	Val	Thr	Ser	Phe	
	530				535				540						545	
agt	gcc	act	gat	atc	tgg	tgg	acc	att	cct	ccc	aag	gca	atg	act	att	1738
Ser	Ala	Thr	Asp	Ile	Trp	Trp	Thr	Ile	Pro	Pro	Lys	Ala	Met	Thr	Ile	
				550					555					560		
cta	ctc	ggg	gcg	tcc	aca	gat	tca	aga	aag	ccc	aat	atc	aat	gtg	acg	1786
Leu	Leu	Gly	Ala	Ser	Thr	Asp	Ser	Arg	Lys	Pro	Asn	Ile	Asn	Val	Thr	
			565					570					575			
ggg	atg	gta	acc	gct	ttg	ggc	aat	tat	gag	ggg	gtg	aac	gaa	ttg	atg	1834
Gly	Met	Val	Thr	Ala	Leu	Gly	Asn	Tyr	Glu	Gly	Val	Asn	Glu	Leu	Met	
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gtc	aat	tag														1843
Val	Asn	*														
		595														

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 <212> PRT
 <213> Fungi

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 Ser Val Tyr Gln Ile Tyr Pro Pro Ser Phe Lys Asp Thr Asn Gly Asp
 20 25 30
 Gly Ile Gly Asp Ile Gln Gly Ile Ser Glu Leu Asp Tyr Ile Lys
 35 40 45

1001827087_1.txt

Ser Leu Gly Val Asp Met Val Trp Leu Ser Pro Ile Leu Ala Ser Pro
 50 55 60
 Gln Val Asp Met Gly Tyr Asp Ile Ser Asp Tyr Lys Asn Ile Tyr Pro
 65 70 75 80
 Pro Tyr Gly Thr Met Ala Asp His Asp Ala Leu Ile Lys Gly Leu His
 85 90 95
 Asp Arg Gly Leu Lys Tyr Ile Leu Asp Leu Val Val Asn His Thr Ser
 100 105 110
 Asp Gln His Pro Trp Phe Lys Glu Ser Lys Ser Lys Thr Ser Arg
 115 120 125
 Tyr Arg Asp Trp Tyr Phe Trp Arg Pro Ala Arg Trp Asp Pro Glu Thr
 130 135 140
 Gly Lys Arg Met Pro Pro Asn Asn Trp Glu Ser Phe Phe Ser Thr Ser
 145 150 155 160
 Ala Trp Thr Trp Asp Ala Thr Thr Gln Glu Tyr Tyr Leu His Leu Trp
 165 170 175
 Ser Ser Gly Gln Pro Asp Leu Asn Trp Glu Asn Pro Asp Val Val Asn
 180 185 190
 Ala Val His Asp Ile Val Arg Phe Trp Leu Asp Arg Gly Val Asp Gly
 195 200 205
 Phe Arg Met Asp Val Ile Asn Tyr Ile Ser Lys Thr Pro Gly Leu Pro
 210 215 220
 Asp Ala Lys Ile Lys Lys Pro Gly Phe Phe Gln Lys Pro Thr Glu His
 225 230 235 240
 Cys Ala Cys Gly Pro Arg Leu His Glu Tyr Leu Arg Gly Ile Gly Ser
 245 250 255
 Ile Leu Gln Glu Tyr Asp Ala Phe Ser Val Gly Glu Met Pro Asp Ala
 260 265 270
 Asp Pro Glu Glu Val Leu Lys Ala Val Gly Gln Asp Arg Gly Glu Leu
 275 280 285
 Ala Met Ala Phe His Phe Asp Ile
 290 295

<210> 521
 <211> 299
 <212> PRT
 <213> Fungi

<400> 521

Asp Gly Leu Asp Leu Gly Ala Asn His Arg Phe Asp Pro Ala Val Phe
 1 5 10 15
 Gln Pro Thr Ala Leu Lys Asn Val Val Asn Lys Trp Gln Thr Phe Met
 20 25 30
 Ala Ser Asn Ala Gly Trp Asn Ala Leu His Met Glu Asn His Asp Glu
 35 40 45
 Ser Arg Thr Val Ser Arg Tyr Ala Cys Asp Ser Pro Ala Met Arg Thr
 50 55 60
 Ile Ser Ala Lys Met Leu Ala Asn His Leu Ala Phe Gln Ser Gly Thr
 65 70 75 80
 Leu Phe Ile Tyr Gln Gly Gln Glu Leu Ala Met Val Asn Leu Pro Arg
 85 90 95
 Glu Trp Gly Met Glu Lys Tyr Lys Asp Ile Glu Cys Leu Asn His Trp
 100 105 110
 Glu Leu Val Gln Arg Cys Tyr His Asp Asp Val Lys Lys Gln Lys Met
 115 120 125
 Tyr Arg Asp Arg Tyr Arg Gln Val Gly Arg Asp Asn Ala Arg Thr Pro
 130 135 140
 Met Gln Trp Asn Ser Glu Ser Pro Tyr Ala Gly Phe Val Pro Ile Gly
 145 150 155 160
 Ser Lys Gln Val Glu Pro Trp Met Ser Ile His Pro Asp Phe Gly Thr
 165 170 175
 Trp Asn Val Ser Thr Met Leu Ala Asp Ser Gln Ser Ser Leu His His
 180 185 190
 Trp Arg Arg Val Leu Lys Phe Arg Lys Gln His Ser Asn Ile Phe Val
 195 200 205
 Tyr Gly Gly Phe Glu Met Leu Asn Ile Glu Ile Glu Glu Asp Val Ile
 210 215 220
 Ala Tyr Ile Arg Thr Asp Asn Thr Thr Gly Gly Ser Leu Thr Gly Pro

1001827087_1.txt

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225      230      235      240
Thr Glu Ala Leu Val Val Thr Ser Phe Ser Ala Thr Asp Ile Trp Trp
      245      250      255
Thr Ile Pro Pro Lys Ala Met Thr Ile Leu Leu Gly Ala Ser Thr Asp
      260      265      270
Ser Arg Lys Pro Asn Ile Asn Val Thr Gly Met Val Thr Ala Leu Gly
      275      280      285
Asn Tyr Glu Gly Val Asn Glu Leu Met Val Asn
      290      295

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<210> 522
 <211> 1788
 <212> DNA
 <213> Fungi

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<400> 522
atgcagcctg tcatcccaaa cgctcgagcga gcctcgtggc gggaagcctc tgtatatcag      60
atctaccac cttcttttaa agataccaat ggggatggta taggagatat tcaaggaatt      120
atctcagagc ttgactatat caagtctctt ggcgttgata tgggatggct gtctcccat      180
ttggcatctc cgcaggttga tatgggttac gatatctctg actataagaa tatctatccg      240
ccttatggta ccatggccga tcatgatgct ttaatcaaa gacctcatga ccgcggtttg      300
aagtatatcc ttgaccttgt tgtgaacct acatcggatc aacatccttg gttcaaagag      360
tccaagtctt caaagacaag cagatatcgc gactgggtatt tctggcggcc ggctcgtcgg      420
gatcccgaaa ccggtaaacg gatgccacct aacaattggg aatccttttt tagtacaagc      480
gcttggactt gggacgctac cacacaagaa tattatcttc acctttgggtc atctggacag      540
cttgacttga actgggaaaa tcccgatgtc gtcaatgctg tgcacgatat cgtaagattc      600
tggcttgatc gtggcgttga tgggtttcga atggatgtca ttaactatat ttcgaaaaca      660
cctggcctgc ccgacgcaa gatcaagaaa ccgggatttt ttcagaagcc caccgagcat      720
tgtgcttggt gaccacggct gcacgaatac ctacgtggga ttggatcgat actccaggaa      780
tacgatgcat ttagtgctcg cgagatgcca gatgcagatc ccgaagaagt tctcaaagct      840
gtgggccagg atcgtggaga attagctatg gcgtttcact ttgacataga tgggcttgac      900
ctaggtgcaa atcatagatt tgaccccgcc gtgttccaac caacagcatt aaagaacgta      960
gtcaacaaat ggcaaacatt catggcatcc aacgcagggtt ggaacgccct gcatatggag      1020
aaccacgacg agagccgaac agttagtcgt tacgcttgcg actctcccg catgcgaaca      1080
atatcagcca agatgctagc aaaccacctt gccttccagt ctggaacact tttcatctat      1140
cagggacagg aacttgcaat ggttaatctt ccccgagaat ggggaatgga gaaatacaaa      1200
gacatcgaat gtctgaatca ctgggagttg gtacagcgtt gttaccacga cgaatgtaaa      1260
aagcagaaga tgtataggga cagataccga caggtcggac gtgacaacgc tcgcacgcc      1320
atgcagtgga actcggagag tccatatgct ggattcgtgc cgattgggtc taagcaagtt      1380
gagccctgga tgtctatcca ccctgacttt gggacttgga acgtttccac gatgttagcc      1440
gattcgcagt ccagccttca ccattggcgc cgggtgctca agtttcgaaa gcagcattcc      1500
aacatatttg tttacggagg attcgaaatg ctaaatatag aaatcgaaga agacgtcatt      1560
gcgtatatac gcactgacaa cactactggt gggagcctaa ctggaccgac ggaggcgctg      1620
gtgttgacca gttttagtgc cactgatata tggtgacca ttcctcccaa ggcaatgact      1680
attctactcg gtgcgtccac agattcaaga aagcccaata tcaatgtgac ggggatggta      1740
accgctttgg gcaattatga ggggtgtgaac gaattgatgg tcaattag      1788

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<210> 523
 <211> 595
 <212> PRT
 <213> Fungi

<220>
 <221> DOMAIN
 <222> (20)...(430)
 <223> Alpha amylase, catalytic domain

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<400> 523
Met Gln Pro Val Ile Pro Asn Val Glu Arg Ala Ser Trp Arg Glu Ala
  1      5      10      15
Ser Val Tyr Gln Ile Tyr Pro Pro Ser Phe Lys Asp Thr Asn Gly Asp
      20      25      30
Gly Ile Gly Asp Ile Gln Gly Ile Ile Ser Glu Leu Asp Tyr Ile Lys
      35      40      45
Ser Leu Gly Val Asp Met Val Trp Leu Ser Pro Ile Leu Ala Ser Pro
      50      55      60
Gln Val Asp Met Gly Tyr Asp Ile Ser Asp Tyr Lys Asn Ile Tyr Pro
      65      70      75      80

```

1001827087_1.txt

Pro	Tyr	Gly	Thr	Met	Ala	Asp	His	Asp	Ala	Leu	Ile	Lys	Gly	Leu	His
Asp	Arg	Gly	Leu	Lys	Tyr	Ile	Leu	Asp	Leu	Val	Val	Asn	His	Thr	Ser
Asp	Gln	His	Pro	Trp	Phe	Lys	Glu	Ser	Lys	Ser	Ser	Lys	Thr	Ser	Arg
Tyr	Arg	Asp	Trp	Tyr	Phe	Trp	Arg	Pro	Ala	Arg	Trp	Asp	Pro	Glu	Thr
Gly	Lys	Arg	Met	Pro	Pro	Asn	Asn	Trp	Glu	Ser	Phe	Phe	Ser	Thr	Ser
Ala	Trp	Thr	Trp	Asp	Ala	Thr	Thr	Gln	Glu	Tyr	Tyr	Leu	His	Leu	Trp
Ser	Ser	Gly	Gln	Pro	Asp	Leu	Asn	Trp	Glu	Asn	Pro	Asp	Val	Val	Asn
Ala	Val	His	Asp	Ile	Val	Arg	Phe	Trp	Leu	Asp	Arg	Gly	Val	Asp	Gly
Phe	Arg	Met	Asp	Val	Ile	Asn	Tyr	Ile	Ser	Lys	Thr	Pro	Gly	Leu	Pro
Asp	Ala	Lys	Ile	Lys	Lys	Pro	Gly	Phe	Phe	Gln	Lys	Pro	Thr	Glu	His
Cys	Ala	Cys	Gly	Pro	Arg	Leu	His	Glu	Tyr	Leu	Arg	Gly	Ile	Gly	Ser
Ile	Leu	Gln	Glu	Tyr	Asp	Ala	Phe	Ser	Val	Gly	Glu	Met	Pro	Asp	Ala
Asp	Pro	Glu	Glu	Val	Leu	Lys	Ala	Val	Gly	Gln	Asp	Arg	Gly	Glu	Leu
Ala	Met	Ala	Phe	His	Phe	Asp	Ile	Asp	Gly	Leu	Asp	Leu	Gly	Ala	Asn
His	Arg	Phe	Asp	Pro	Ala	Val	Phe	Gln	Pro	Thr	Ala	Leu	Lys	Asn	Val
Val	Asn	Lys	Trp	Gln	Thr	Phe	Met	Ala	Ser	Asn	Ala	Gly	Trp	Asn	Ala
Leu	His	Met	Glu	Asn	His	Asp	Glu	Ser	Arg	Thr	Val	Ser	Arg	Tyr	Ala
Cys	Asp	Ser	Pro	Ala	Met	Arg	Thr	Ile	Ser	Ala	Lys	Met	Leu	Ala	Asn
His	Leu	Ala	Phe	Gln	Ser	Gly	Thr	Leu	Phe	Ile	Tyr	Gln	Gly	Gln	Glu
Leu	Ala	Met	Val	Asn	Leu	Pro	Arg	Glu	Trp	Gly	Met	Glu	Lys	Tyr	Lys
Asp	Ile	Glu	Cys	Leu	Asn	His	Trp	Glu	Leu	Val	Gln	Arg	Cys	Tyr	His
Asp	Asp	Val	Lys	Lys	Gln	Lys	Met	Tyr	Arg	Asp	Arg	Tyr	Arg	Gln	Val
Gly	Arg	Asp	Asn	Ala	Arg	Thr	Pro	Met	Gln	Trp	Asn	Ser	Glu	Ser	Pro
Tyr	Ala	Gly	Phe	Val	Pro	Ile	Gly	Ser	Lys	Gln	Val	Glu	Pro	Trp	Met
Ser	Ile	His	Pro	Asp	Phe	Gly	Thr	Trp	Asn	Val	Ser	Thr	Met	Leu	Ala
Asp	Ser	Gln	Ser	Ser	Leu	His	His	Trp	Arg	Arg	Val	Leu	Lys	Phe	Arg
Lys	Gln	His	Ser	Asn	Ile	Phe	Val	Tyr	Gly	Gly	Phe	Glu	Met	Leu	Asn
Ile	Glu	Ile	Glu	Glu	Asp	Val	Ile	Ala	Tyr	Ile	Arg	Thr	Asp	Asn	Thr
Thr	Gly	Gly	Ser	Leu	Thr	Gly	Pro	Thr	Glu	Ala	Leu	Val	Val	Thr	Ser
Phe	Ser	Ala	Thr	Asp	Ile	Trp	Trp	Thr	Ile	Pro	Pro	Lys	Ala	Met	Thr
Ile	Leu	Leu	Gly	Ala	Ser	Thr	Asp	Ser	Arg	Lys	Pro	Asn	Ile	Asn	Val
Thr	Gly	Met	Val	Thr	Ala	Leu	Gly	Asn	Tyr	Glu	Gly	Val	Asn	Glu	Leu
Met	Val	Asn													

<210> 524

<211> 1820
 <212> DNA
 <213> Fungi

<220>
 <221> CDS
 <222> (1)...(277)
 <223> Exon

<221> CDS
 <222> (343)...(1820)
 <223> Exon

<221> misc_feature
 <222> (791)...(800)
 <223> n = a, g, c or t

<400> 524
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 Met Gly Ser Ile Ala Arg Arg Thr Ile Gly Ser Thr Pro Gln Trp Trp
 1 5 10 15

aaa gaa gca gtc gta tat caa atc tat ccc gct tcg tat ctc gac acc 96
 Lys Glu Ala Val Val Tyr Gln Ile Tyr Pro Ala Ser Tyr Leu Asp Thr
 20 25 30

acc gga tct gga gat ggt gat ctc aat ggt atc acc tcg aaa ttg cca 144
 Thr Gly Ser Gly Asp Gly Asp Leu Asn Gly Ile Thr Ser Lys Leu Pro
 35 40 45

tat atc aga tct ctc ggt gta gat gta gta tgg att tca cct atc tat 192
 Tyr Ile Arg Ser Leu Gly Val Asp Val Val Trp Ile Ser Pro Ile Tyr
 50 55 60

gca tcc cct atg aat gat atg ggc tat gat att tcg gat tat cga gct 240
 Ala Ser Pro Met Asn Asp Met Gly Tyr Asp Ile Ser Asp Tyr Arg Ala
 65 70 75 80

atc aac cct atg ttt ggt act atg gaa gat tgg gaa c gtgagtcacc 287
 Ile Asn Pro Met Phe Gly Thr Met Glu Asp Trp Glu
 85 90

tcttcctcat tcttttctca aagctctcaa aaaaaaaaaa attaacagcg cctag gt 344
 Arg

ctt tgc gcc aga gca cac gaa cta ggt ttg aaa ctc gtg atg gat ctc 392
 Leu Cys Ala Arg Ala His Glu Leu Gly Leu Lys Leu Val Met Asp Leu
 95 100 105

gtg gta aat cac act agt tct gaa cac ccc tgg ttc aaa gaa tcc gtc 440
 Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Lys Glu Ser Val
 110 115 120 125

tcc ggc ggt ccc aac ggt cct aaa aga gac ttc tac tac tgg caa ccg 488
 Ser Gly Gly Pro Asn Gly Pro Lys Arg Asp Phe Tyr Tyr Trp Gln Pro
 130 135 140

ccc aaa aat ggt aaa gag ccc aat aac tgg ggt gcc atg ttt gga ggt 536
 Pro Lys Asn Gly Lys Glu Pro Asn Asn Trp Gly Ala Met Phe Gly Gly
 145 150 155

tca tcc tgg gaa aaa gat cct tcg cat caa aca gac gaa tac tat ctc 584
 Ser Ser Trp Glu Lys Asp Pro Ser His Gln Thr Asp Glu Tyr Tyr Leu
 160 165 170

cac gtc tac gac gtc agc cag ccc gat ctt aat tgg aca aat cca gcc 632
 His Val Tyr Asp Val Ser Gln Pro Asp Leu Asn Trp Thr Asn Pro Ala

1001827087_1.txt

175	180	185	
gtt Val 190	cgc Arg	aac Asn	gag Glu
gtc Val	tgg Trp 195	gat Asp	atc Ile
atg Met	cgc Arg	ttt Phe 200	tgg Trp
ctt Leu	gat Asp	aag Lys	gga Gly 205
680			
tgt Cys	gat Asp	ggg Gly	ttt Phe
cgt Arg 210	atg Met	gat Asp	gtc Val
atc Ile	aat Asn 215	tgc Cys	att Ile
tca Ser	aaa Lys	gaa Glu 220	cct Pro
728			
gga Gly	ttt Phe	ccc Pro	gat Asp 225
ttt Phe	gaa Glu	gtc Val	acc Thr
gat Asp 230	ccg Pro	aga Arg	ctc Leu
gaa Glu	ttc Phe 235	cag Gln	att Ile
776			
ggt Gly	gtg Val	agg Arg 240	gga Gly
aan Xaa	nnn Xaa	nnn Xaa	nnn Xaa 245
cac His	gtc Val	aaa Lys	gag Glu
tac Tyr 250	ttg Leu	gaa Glu	gaa Glu
824			
atg Met	cat His 255	cgc Arg	gaa Glu
gtt Val	ctc Leu	tgt Cys 260	cat His
tat Tyr	ccc Pro	gaa Glu	gca Ala 265
ttt Phe	aca Thr	gtg Val	gga Gly
872			
gaa Glu 270	acc Thr	cca Pro	gga Gly
ctc Leu	aag Lys 275	aga Arg	ccc Pro
gag Glu	agt Ser	gca Ala 280	ttg Leu
ggc Gly	ctg Leu	gtc Val	caa Gln 285
920			
gga Gly	gga Gly	aaa Lys	cct Pro
ctt Leu 290	caa Gln	atg Met	atc Ile
ttc Phe	caa Gln 295	ttc Phe	ttc Phe
gag Glu	cat His	atg Met	tac Tyr 300
968			
gac Asp	atc Ile	caa Gln	ccc Pro 305
gga Gly	atg Met	aaa Lys	ccc Pro
ttc Phe 310	ttc Phe	cct Pro	cgt Arg
cct Pro	cgt Arg	cct Pro	tca Ser 315
tgg Trp	aat Asn		
1016			
ctt Leu	acc Thr	gag Glu	atc Ile
aag Lys	aaa Lys	aat Asn	ctc Leu 325
ggg Gly	gag Glu	tgg Trp	atg Met
tct Ser 330	ttc Phe	aat Asn	gca Glu
1064			
gag Glu	gtg Trp	gac Asp	ggc Gly
tgg Trp	gaa Glu	aaa Lys	gat Asp
gca Ala	aaa Lys	gat Asp	gta Val 405
gaa Glu	tcg Ser	aat Asn	ggg Gly
aaa Lys 420	gaa Glu	tcg Ser	aat Asn
1352			
act Thr 430	gat Asp	atc Ile	tcc Ser
cga Arg	gga Gly 435	atc Ile	caa Gln
ggc Gly	tac Tyr	gga Gly 440	aga Arg
gac Asp	aat Asn	tcg Ser	aga Arg 445
1400			
atg Met	ggg Gly	atg Met	caa Gln
tgg Trp	gac Asp	gac Asp	tct Ser
cca Pro	aat Asn	ggc Gly	ggt Gly
ttc Phe	aca Thr	caa Gln	ggt Gly
1448			

1001827087_1.txt

	450	455	460	
aaa ccg tgg atc aag acg aat gag gag tac aag gaa atc aac gtc gca	Lys Pro Trp Ile 465	Lys Thr Asn Glu 470	Tyr Lys Glu Ile 475	1496
gcg caa gat gga gtc aag gga agc aca tta gag ttc tgg aaa cag ata	Ala Gln Asp 480	Gly Val Lys Gly 485	Thr Leu Glu Phe Trp 490	1544
att aaa ctc cgg aaa gag aat ccg gtg ttg tgt aaa ggg gga ttt gaa	Ile Lys 495	Leu Arg Lys Glu 500	Pro Val Leu Cys Lys 505	1592
atg gtg gat cag gag aac gag gag gtt tat gcg tat gtg aga aag ggc	Met Val Asp 510	Gln Glu Asn Glu 515	Val Tyr Ala Tyr Val Arg Lys Gly 525	1640
gag gag aag gag tat ttg att gct tgc aat ttc aag gag cgg gat gtg	Glu Glu Lys Glu 530	Tyr Leu Ile Ala Cys Asn 535	Phe Lys Glu Arg Asp Val 540	1688
aaa tgg aag att ccg gtt gag act ggg gag ctt ttg ttt gga agc tat	Lys Trp Lys Ile 545	Pro Val Glu Thr Gly 550	Glu Leu Leu Phe Gly 555	1736
gag gac gga gta cag ggt gga gag aag gag gga gaa tta aac ttg aga	Glu Asp Gly Val 560	Gln Gly Gly Glu 565	Lys Glu Gly Glu Leu Asn Leu Arg 570	1784
ccg ttc gag gga aga ata tat gtt aga gat att tga	Pro Phe Glu Gly Arg Ile Tyr 580	Val Arg Asp Ile *		1820

<210> 525
 <211> 92
 <212> PRT
 <213> Fungi

<400> 525
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 Lys Glu Ala Val Tyr Gln Ile Tyr Pro Ala Ser Tyr Leu Asp Thr
 20 25 30
 Thr Gly Ser Gly Asp Gly Asp Leu Asn Gly Ile Thr Ser Lys Leu Pro
 35 40 45
 Tyr Ile Arg Ser Leu Gly Val Asp Val Val Trp Ile Ser Pro Ile Tyr
 50 55 60
 Ala Ser Pro Met Asn Asp Met Gly Tyr Asp Ile Ser Asp Tyr Arg Ala
 65 70 75 80
 Ile Asn Pro Met Phe Gly Thr Met Glu Asp Trp Glu
 85 90

<210> 526
 <211> 492
 <212> PRT
 <213> Fungi

<220>
 <221> UNSURE
 <222> (150)...(154)
 <223> Xaa = any amino acid

<400> 526
 Arg Leu Cys Ala Arg Ala His Glu Leu Gly Leu Lys Leu Val Met Asp
 1 5 10 15
 Leu Val Val Asn His Thr Ser Ser Glu His Pro Trp Phe Lys Glu Ser

1001827087_1.txt

Val	Ser	Gly	20	Gly	Pro	Asn	Gly	Pro	25	Lys	Arg	Asp	Phe	Tyr	30	Tyr	Trp	Gln
Pro	Pro	Lys	35	Asn	Gly	Lys	Glu	Pro	40	Asn	Asn	Trp	Gly	Ala	Met	Phe	Gly	
Gly	Ser	Ser	50	Trp	Glu	Lys	Asp	Pro	55	Ser	His	Gln	Thr	Asp	Glu	Tyr	Tyr	
Leu	His	Val	65	Tyr	Asp	Val	Ser	Gln	70	Pro	Asp	Leu	Asn	Trp	Thr	Asn	Pro	
Ala	Val	Arg	85	Asn	Glu	Val	Trp	Asp	90	Ile	Met	Arg	Phe	Trp	Leu	Asp	Lys	
Gly	Cys	Asp	100	Gly	Phe	Arg	Met	Asp	105	Val	Ile	Asn	Cys	Ile	Ser	Lys	Glu	
Pro	Gly	Phe	115	Pro	Asp	Phe	Glu	Val	120	Thr	Asp	Pro	Arg	Leu	Glu	Phe	Gln	
Ile	Gly	Val	130	Arg	Gly	Xaa	Xaa	Xaa	135	His	Val	Lys	Glu	Tyr	Leu	Glu		
Glu	Met	His	145	Arg	Glu	Val	Leu	Cys	150	His	Tyr	Pro	Glu	Ala	Phe	Thr	Val	
Gly	Glu	Thr	165	Pro	Gly	Leu	Lys	Arg	170	Pro	Glu	Ser	Ala	Leu	Gly	Leu	Val	
Gln	Gly	Gly	180	Lys	Pro	Leu	Gln	Met	185	Ile	Phe	Gln	Phe	Glu	His	Met	Tyr	
Phe	Asp	Ile	195	Gln	Pro	Gly	Met	Lys	200	Pro	Phe	Phe	Pro	Arg	Pro	Ser	Trp	
Asn	Leu	Thr	210	Glu	Ile	Lys	Lys	Asn	215	Leu	Gly	Glu	Trp	Met	Ser	Phe	Asn	
Ala	Ala	His	225	Asp	Gly	Trp	Asp	Ser	230	Leu	Tyr	Leu	Glu	Asn	His	Asp	Gln	
Pro	Arg	Ile	245	Leu	Gly	Arg	Trp	Ala	250	Lys	Asp	Thr	Pro	Glu	Trp	Arg	Val	
Gln	Ala	Ala	260	Lys	Met	Leu	Ala	Leu	265	Phe	His	Ala	Thr	Gly	Arg	Gly	Thr	
Leu	Phe	Val	275	Tyr	Gln	Gly	Gln	Glu	280	Ile	Gly	Thr	Ala	Asn	Ser	Lys	Trp	
Trp	Thr	Asn	290	Glu	Glu	Phe	Arg	Asp	295	Val	Glu	Glu	Ile	Asn	Phe	Phe	Ala	
Ala	Glu	Lys	305	Glu	Arg	Glu	Lys	Glu	310	Ser	Glu	Asn	Gly	Lys	Glu	Val	Asp	
Met	Thr	Asp	325	Ile	Ser	Arg	Gly	Ile	330	Gln	Gly	Tyr	Gly	Arg	Asp	Asn	Ser	
Arg	Met	Gly	340	Met	Gln	Trp	Asp	Asp	345	Ser	Pro	Asn	Gly	Gly	Phe	Thr	Gln	
Gly	Lys	Pro	355	Trp	Ile	Lys	Thr	Asn	360	Glu	Glu	Tyr	Lys	Glu	Ile	Asn	Val	
Ala	Ala	Gln	370	Asp	Gly	Val	Lys	Gly	375	Ser	Thr	Leu	Glu	Phe	Trp	Lys	Gln	
Ile	Ile	Lys	385	Leu	Arg	Lys	Glu	Asn	390	Pro	Val	Leu	Cys	Lys	Gly	Gly	Phe	
Glu	Met	Val	405	Gln	Glu	Asn	Glu	Glu	410	Val	Tyr	Ala	Tyr	Val	Arg	Lys		
Gly	Glu	Glu	420	Lys	Glu	Tyr	Leu	Ile	425	Ala	Cys	Asn	Phe	Lys	Glu	Arg	Asp	
Val	Lys	Trp	435	Lys	Ile	Pro	Val	Glu	440	Thr	Gly	Glu	Leu	Leu	Phe	Gly	Ser	
Tyr	Glu	Asp	450	Gly	Val	Gln	Gly	Gly	455	Glu	Lys	Glu	Gly	Glu	Leu	Asn	Leu	
Arg	Pro	Phe	465	Glu	Gly	Arg	Ile	Tyr	470	Val	Arg	Asp	Ile					
			485						490									

<210> 527
 <211> 1755
 <212> DNA
 <213> Fungi

<220>
 <221> misc_feature
 <222> (726)...(735)

<223> n = a, g, c or t

<400> 527

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gtatatcaaa	tctatcccg	ttcgtatctc	gacaccaccg	gatctggaga	tggtgatctc	120
aatggtatca	cctcgaaatt	gccatatatc	agatctctcg	gtgtagatgt	agtatggatt	180
tcacctatct	atgcatcccc	tatgaatgat	atgggctatg	atatttcgga	ttatcgagct	240
atcaacccta	tgtttggtac	tatggaagat	tggaacgctc	tttgcgccag	agcacacgaa	300
ctaggtttga	aactcgtgat	ggatctcgtg	gtaaatcaca	ctagttctga	acaccctgg	360
ttcaaagaat	ccgtctccgg	cggccccaac	ggctcctaaa	gagacttcta	ctactggcaa	420
ccgccccaaa	atggtaaaga	gccccataac	tggggtgcca	tgtttgagg	ttcatcctgg	480
gaaaaagatc	cttcgcatca	aacagacgaa	tactatctcc	acgtctacga	cgtcagccag	540
cccgatctta	attggacaaa	tccagccggt	cgcaacgagg	tctgggatat	catgcgcttt	600
tggtttgata	agggatgtga	tggttttcgt	atggatgtca	tcaattgcat	ttcaaaagaa	660
cttggatttc	ccgattttga	agtcaccgat	ccgagactcg	aattccagat	tggtgtgagg	720
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ccggtgttgt	gtaaaggggg	atttgaaatg	gtggatcagg	agaacgagga	ggtttatgcg	1560
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gtgaaatgga	agattccggt	tgagactggg	gagcttttgt	ttggaagcta	tgaggacgga	1680
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<211> 584

<212> PRT

<213> Fungi

<220>

<221> DOMAIN

<222> (23)...(445)

<223> Alpha amylase, catalytic domain

<221> UNSURE

<222> (242)...(245)

<223> Xaa = any amino acid

<400> 528

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Thr	Gly	Ser	Gly	Asp	Gly	Asp	Leu	Asn	Gly	Ile	Thr	Ser	Lys	Leu	Pro
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Tyr	Ile	Arg	Ser	Leu	Gly	Val	Asp	Val	Val	Trp	Ile	Ser	Pro	Ile	Tyr
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Ala	Ser	Pro	Met	Asn	Asp	Met	Gly	Tyr	Asp	Ile	Ser	Asp	Tyr	Arg	Ala
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Ile	Asn	Pro	Met	Phe	Gly	Thr	Met	Glu	Asp	Trp	Glu	Arg	Leu	Cys	Ala
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Pro	Asn	Gly	Pro	Lys	Arg	Asp	Phe	Tyr	Tyr	Trp	Gln	Pro	Pro	Lys	Asn
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Gly	Lys	Glu	Pro	Asn	Asn	Trp	Gly	Ala	Met	Phe	Gly	Gly	Ser	Ser	Trp

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Glu	Val	Trp	Asp	Ile	Met	Arg	Phe	Trp	Leu	Asp	Lys	Gly	Cys	Asp	Gly				
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Pro	Gly	Met	Lys	Pro	Phe	Phe	Pro	Arg	Pro	Ser	Trp	Asn	Leu	Thr	Glu				
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Met	Leu	Ala	Leu	Phe	His	Ala	Thr	Gly	Arg	Gly	Thr	Leu	Phe	Val	Tyr				
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Gln	Gly	Gln	Glu	Ile	Gly	Thr	Ala	Asn	Ser	Lys	Trp	Trp	Thr	Asn	Glu				
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Glu	Phe	Arg	Asp	Val	Glu	Glu	Ile	Asn	Phe	Phe	Ala	Ala	Glu	Lys	Glu				
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Arg	Arg	Glu	Lys	Glu	Ser	Asn	Gly	Lys	Glu	Val	Asp	Met	Thr	Asp	Ile				
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Ser	Arg	Gly	Ile	Gln	Gly	Tyr	Gly	Arg	Asp	Asn	Ser	Arg	Met	Gly	Met				
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Gln	Trp	Asp	Asp	Ser	Pro	Asn	Gly	Gly	Phe	Thr	Gln	Gly	Lys	Pro	Trp				
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Ile	Lys	Thr	Asn	Glu	Glu	Tyr	Lys	Glu	Ile	Asn	Val	Ala	Ala	Gln	Asp				
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Gly	Val	Lys	Gly	Ser	Thr	Leu	Glu	Phe	Trp	Lys	Gln	Ile	Ile	Lys	Leu				
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Arg	Lys	Glu	Asn	Pro	Val	Leu	Cys	Lys	Gly	Gly	Phe	Glu	Met	Val	Asp				
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Gln	Glu	Asn	Glu	Glu	Val	Tyr	Ala	Tyr	Val	Arg	Lys	Gly	Glu	Glu	Lys				
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Glu	Tyr	Leu	Ile	Ala	Cys	Asn	Phe	Lys	Glu	Arg	Asp	Val	Lys	Trp	Lys				
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Val	Gln	Gly	Gly	Glu	Lys	Glu	Gly	Glu	Leu	Asn	Leu	Arg	Pro	Phe	Glu				
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<210> 529
 <211> 1883
 <212> DNA
 <213> Cochliobolus heterostrophus ATCC 48331

<220>
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 <222> (1)...(1265)
 <223> Exon

<221> CDS
 <222> (1352)...(1883)

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<223> Exon

<400> 529

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His Arg Trp Trp Lys Glu Ala Val Val Tyr Gln Ile Tyr Pro Ala Ser	
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Phe Leu Asp Ser Asn Gly Asp Gly Trp Gly Asp Val Lys Gly Ile Thr	
35 40 45	
tcg aag ctg gat tac ctc aag gat cta ggc atc gat gtg atc tgg cta	192
Ser Lys Leu Asp Tyr Leu Lys Asp Leu Gly Ile Asp Val Ile Trp Leu	
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tcc cca atc tac aaa agc ccg caa gcc gat atg ggc tac gac att gca	240
Ser Pro Ile Tyr Lys Ser Pro Gln Ala Asp Met Gly Tyr Asp Ile Ala	
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gac tac gaa gat ata gac ccc agc tat ggt act ttg gct gat gtc gac	288
Asp Tyr Glu Asp Ile Asp Pro Ser Tyr Gly Thr Leu Ala Asp Val Asp	
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aac ctg atc cac gaa acc aag aaa cgc ggc atg aag ctt gtc atg gat	336
Asn Leu Ile His Glu Thr Lys Lys Arg Gly Met Lys Leu Val Met Asp	
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Leu Val Val Asn His Thr Ser Glu Glu His Ala Trp Phe Leu Asp Ser	
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Arg Ser Ser Lys Glu Ser Lys Lys Arg Asp Trp Tyr Ile Trp Lys Pro	
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Gln Ile Leu Gly Glu Ala Asn Ser Ala Trp Thr Trp Asp Glu Lys Thr	
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Gln Glu Tyr Tyr Leu Ser Leu Phe Thr Pro Glu Gln Pro Asp Leu Asn	
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Ile Ser Lys Val Gln Thr Phe Pro Asp Ala Pro Ile Ser Val Lys Asp	
225 230 235 240	
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Ala Lys Tyr Gln Pro Gly Asp Lys Tyr Phe Ala Asn Gly Pro Arg Leu	
245 250 255	
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Thr	Leu	Thr	Val	Gly	Glu	Met	Pro	Phe	Val	Arg	Asp	Glu	Asp	Glu	Val	
		275					280					285				
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Ile	Lys	Val	Val	Gly	Ser	Glu	Ser	Gly	Glu	Leu	Asn	Met	Ile	Phe	Asn	
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ttc	gac	ctt	gtc	gac	att	gac	aac	gtt	cct	ggc	gac	ttc	aag	tac	aca	960
Phe	Asp	Leu	Val	Asp	Ile	Asp	Asn	Val	Pro	Gly	Asp	Phe	Lys	Tyr	Thr	
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Leu	His	Pro	Trp	Asp	Ala	Arg	Asp	Leu	Lys	Lys	Ile	Val	Asn	Arg	Leu	
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cag	cgt	ctg	atg	ctt	gag	cgt	gac	ggg	tgg	aac	aca	cta	tat	gtg	gaa	1056
Gln	Arg	Leu	Met	Leu	Glu	Arg	Asp	Gly	Trp	Asn	Thr	Leu	Tyr	Val	Glu	
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aac	cac	gac	cag	ccc	cgc	agc	ata	agc	aga	tat	act	gac	gac	agc	gat	1104
Asn	His	Asp	Gln	Pro	Arg	Ser	Ile	Ser	Arg	Tyr	Thr	Asp	Asp	Ser	Asp	
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Glu	Trp	Arg	Ser	Tyr	Gly	Ala	Lys	Leu	Leu	Cys	Leu	Met	Gln	Thr	Thr	
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Leu	Ala	Gly	Thr	Leu	Tyr	Val	Tyr	Gln	Gly	Glu	Glu	Ile	Gly	Met	Arg	
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Asn	Ile	Pro	Lys	Glu	Trp	Gly	Pro	Glu	Glu	Tyr	Lys	Asp	Ile	Glu	Ser	
				405				410						415		
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Ile	Asn	Phe	Phe	Lys	Lys											
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tac	cgc	gac	ctt	tac	ccc	aac	aac	ccc	gaa	aaa	caa	gcc	ctc	gca	gca	1400
Tyr	Arg	Asp	Leu	Tyr	Pro	Asn	Asn	Pro	Glu	Lys	Gln	Ala	Leu	Ala	Ala	
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gaa	atc	atg	caa	cgc	aaa	gcc	cgc	gac	agc	gcc	cga	aca	ccc	atg	caa	1448
Glu	Ile	Met	Gln	Arg	Lys	Ala	Arg	Asp	Ser	Ala	Arg	Thr	Pro	Met	Gln	
	440					445					450					
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Trp	Asn	Ala	Asp	Ser	Gln	Ala	Gly	Phe	Thr	Ser	Gly	Thr	Pro	Trp	Met	
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cgc	gtc	aac	acg	gac	tat	ccc	gcc	atc	aac	gtc	gcc	gag	caa	ctc	gcc	1544
Arg	Val	Asn	Thr	Asp	Tyr	Pro	Ala	Ile	Asn	Val	Ala	Glu	Gln	Leu	Ala	
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Asn	Pro	Thr	Pro	Ala	Pro	Gly	Thr	Leu	Ser	Val	His	Ala	Phe	Trp	Lys	
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Arg	Ala	Leu	Glu	Tyr	Arg	Lys	Glu	Asn	Lys	Asp	Val	Phe	Val	Tyr	Gly	
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cgg	tgg	ggc	aag	gat	aag	gcg	ttt	atc	act	gtg	ctg	aat	ttt	agc	ggg	1736
Arg	Trp	Gly	Lys	Asp	Lys	Ala	Phe	Ile	Thr	Val	Leu	Asn	Phe	Ser	Gly	
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Val	Ala	Gly	Asn	Tyr	Asp	Glu	Arg	Glu	Leu	Glu	Thr	Lys	Ala	Lys	Ser	
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ggg	gcg	ttg	gag	ttg	agg	gct	tgg	gag	ggg	ttg	ctg	gga	ctt	ttg	gag	1880
Gly	Ala	Leu	Glu	Leu	Arg	Ala	Trp	Glu	Gly	Leu	Leu	Gly	Leu	Leu	Glu	
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<210> 530
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 <213> Cochliobolus heterostrophus ATCC 48331

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 Phe Leu Asp Ser Asn Gly Asp Gly Trp Gly Asp Val Lys Gly Ile Thr
 35 40 45
 Ser Lys Leu Asp Tyr Leu Lys Asp Leu Gly Ile Asp Val Ile Trp Leu
 50 55 60
 Ser Pro Ile Tyr Lys Ser Pro Gln Ala Asp Met Gly Tyr Asp Ile Ala
 65 70 75 80
 Asp Tyr Glu Asp Ile Asp Pro Ser Tyr Gly Thr Leu Ala Asp Val Asp
 85 90 95
 Asn Leu Ile His Glu Thr Lys Lys Arg Gly Met Lys Leu Val Met Asp
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 Leu Val Val Asn His Thr Ser Glu Glu His Ala Trp Phe Leu Asp Ser
 115 120 125
 Arg Ser Ser Lys Glu Ser Lys Lys Arg Asp Trp Tyr Ile Trp Lys Pro
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 Ala Lys Tyr Asp Ala Asp Gly Asn Arg Gln Pro Pro Asn Asn Trp Ala
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 Gln Glu Tyr Tyr Leu Ser Leu Phe Thr Pro Glu Gln Pro Asp Leu Asn
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 Trp Glu Asn Pro Asp Val Arg Glu Ala Val His Asn Ile Leu Arg Phe
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 Trp Leu Asp Arg Gly Val Ser Gly Phe Arg Met Asp Val Ile Asn Leu
 210 215 220
 Ile Ser Lys Val Gln Thr Phe Pro Asp Ala Pro Ile Ser Val Lys Asp
 225 230 235 240
 Ala Lys Tyr Gln Pro Gly Asp Lys Tyr Phe Ala Asn Gly Pro Arg Leu
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 His Glu Trp Leu Lys Glu Leu Arg Arg Asp Val Leu Ser Lys Tyr Asp
 260 265 270
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290 295 300
Phe Asp Leu Val Asp Ile Asp Asn Val Pro Gly Asp Phe Lys Tyr Thr
305 310 315 320
Leu His Pro Trp Asp Ala Arg Asp Leu Lys Lys Ile Val Asn Arg Leu
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340 345 350
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Glu Trp Arg Ser Tyr Gly Ala Lys Leu Leu Cys Leu Met Gln Thr Thr
370 375 380
Leu Ala Gly Thr Leu Tyr Val Tyr Gln Gly Glu Glu Ile Gly Met Arg
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Arg Val Asn Thr Asp Tyr Pro Ala Ile Asn Val Ala Glu Gln Leu Ala
50 55 60
Asn Pro Thr Pro Ala Pro Gly Thr Leu Ser Val His Ala Phe Trp Lys
65 70 75 80
Arg Ala Leu Glu Tyr Arg Lys Glu Asn Lys Asp Val Phe Val Tyr Gly
85 90 95
Asp Phe Glu Met Leu Asp Met Gly His Glu Lys Val Val Ala Phe Arg
100 105 110
Arg Trp Gly Lys Asp Lys Ala Phe Ile Thr Val Leu Asn Phe Ser Gly
115 120 125
Glu Glu Val Lys Trp Gly Gly Leu Gly Glu Ile Lys Val Lys Lys Trp
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Val Ala Gly Asn Tyr Asp Glu Arg Glu Leu Glu Thr Lys Ala Lys Ser
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Gly Ala Leu Glu Leu Arg Ala Trp Glu Gly Leu Leu Gly Leu Leu Glu
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aacataccca	aagaatgggg	cccagaagag	tacaaagaca	ttgaaagtat	caatttcttc	1260
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gcaggcttca	catccggaac	cccctggatg	cgcgtcaaca	cggactatcc	cgccatcaac	1440
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tggaagcgtg	ccctggagta	ccgcaaggaa	aacaaggacg	ttttcgtgta	cggggatttt	1560
gaaatgctgg	atatgggaca	tgagaagggt	gtggcggtta	ggcgggtggg	caaggataag	1620
gcgtttatca	ctgtgctgaa	ttttagcggg	gaggaggtta	agtggggagg	tttgggtgaa	1680
ataaaggtag	agaagtgggt	tgcggggaat	tatgatgaga	gggagttgga	aacgaaggcc	1740
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<210> 533
 <211> 598
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

<220>
 <221> DOMAIN
 <222> (27)...(450)
 <223> Alpha amylase, catalytic domain

<400> 533
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 His Arg Trp Trp Lys Glu Ala Val Val Tyr Gln Ile Tyr Pro Ala Ser
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 Phe Leu Asp Ser Asn Gly Asp Gly Trp Gly Asp Val Lys Gly Ile Thr
 35 40 45
 Ser Lys Leu Asp Tyr Leu Lys Asp Leu Gly Ile Asp Val Ile Trp Leu
 50 55 60
 Ser Pro Ile Tyr Lys Ser Pro Gln Ala Asp Met Gly Tyr Asp Ile Ala
 65 70 75 80
 Asp Tyr Glu Asp Ile Asp Pro Ser Tyr Gly Thr Leu Ala Asp Val Asp
 85 90 95
 Asn Leu Ile His Glu Thr Lys Lys Arg Gly Met Lys Leu Val Met Asp
 100 105 110
 Leu Val Val Asn His Thr Ser Glu Glu His Ala Trp Phe Leu Asp Ser
 115 120 125
 Arg Ser Lys Lys Glu Ser Lys Lys Arg Asp Trp Tyr Ile Trp Lys Pro
 130 135 140
 Ala Lys Tyr Asp Ala Asp Gly Asn Arg Gln Pro Pro Asn Asn Trp Ala
 145 150 155 160
 Gln Ile Leu Gly Glu Ala Asn Ser Ala Trp Thr Trp Asp Glu Lys Thr
 165 170 175
 Gln Glu Tyr Tyr Leu Ser Leu Phe Thr Pro Glu Gln Pro Asp Leu Asn
 180 185 190
 Trp Glu Asn Pro Asp Val Arg Glu Ala Val His Asn Ile Leu Arg Phe
 195 200 205
 Trp Leu Asp Arg Gly Val Ser Gly Phe Arg Met Asp Val Ile Asn Leu
 210 215 220
 Ile Ser Lys Val Gln Thr Phe Pro Asp Ala Pro Ile Ser Val Lys Asp
 225 230 235 240
 Ala Lys Tyr Gln Pro Gly Asp Lys Tyr Phe Ala Asn Gly Pro Arg Leu
 245 250 255
 His Glu Trp Leu Lys Glu Leu Arg Arg Asp Val Leu Ser Lys Tyr Asp
 260 265 270
 Thr Leu Thr Val Gly Glu Met Pro Phe Val Arg Asp Glu Asp Glu Val
 275 280 285
 Ile Lys Val Val Gly Ser Glu Ser Gly Glu Leu Asn Met Ile Phe Asn
 290 295 300
 phe Asp Leu Val Asp Ile Asp Asn Val Pro Gly Asp Phe Lys Tyr Thr
 305 310 315 320
 Leu His Pro Trp Asp Ala Arg Asp Leu Lys Lys Ile Val Asn Arg Leu

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Gln Arg Leu Met Leu Glu Arg Asp Gly Trp Asn Thr Leu Tyr Val Glu
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Asn His Asp Gln Pro Arg Ser Ile Ser Arg Tyr Thr Asp Asp Ser Asp
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Glu Trp Arg Ser Tyr Gly Ala Lys Leu Leu Cys Leu Met Gln Thr Thr
      370      375      380
Leu Ala Gly Thr Leu Tyr Val Tyr Gln Gly Glu Glu Ile Gly Met Arg
385      390      395      400
Asn Ile Pro Lys Glu Trp Gly Pro Glu Glu Tyr Lys Asp Ile Glu Ser
      405      410      415
Ile Asn Phe Phe Lys Lys Tyr Arg Asp Leu Tyr Pro Asn Asn Pro Glu
      420      425      430
Lys Gln Ala Leu Ala Ala Glu Ile Met Gln Arg Lys Ala Arg Asp Ser
      435      440      445
Ala Arg Thr Pro Met Gln Trp Asn Ala Asp Ser Gln Ala Gly Phe Thr
      450      455      460
Ser Gly Thr Pro Trp Met Arg Val Asn Thr Asp Tyr Pro Ala Ile Asn
465      470      475      480
Val Ala Glu Gln Leu Ala Asn Pro Thr Pro Ala Pro Gly Thr Leu Ser
      485      490      495
Val His Ala Phe Trp Lys Arg Ala Leu Glu Tyr Arg Lys Glu Asn Lys
      500      505      510
Asp Val Phe Val Tyr Gly Asp Phe Glu Met Leu Asp Met Gly His Glu
      515      520      525
Lys Val Val Ala Phe Arg Arg Trp Gly Lys Asp Lys Ala Phe Ile Thr
      530      535      540
Val Leu Asn Phe Ser Gly Glu Glu Val Lys Trp Gly Gly Leu Gly Glu
545      550      555      560
Ile Lys Val Lys Lys Trp Val Ala Gly Asn Tyr Asp Glu Arg Glu Leu
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Glu Thr Lys Ala Lys Ser Gly Ala Leu Glu Leu Arg Ala Trp Glu Gly
      580      585      590
Leu Leu Gly Leu Leu Glu
      595

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<210> 534
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 <213> Cochliobolus heterostrophus ATCC 48331

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 <223> Exon

<221> CDS
 <222> (316)...(425)
 <223> Exon

<221> CDS
 <222> (483)...(1002)
 <223> Exon

<221> CDS
 <222> (1045)...(1922)
 <223> Exon

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atc gtc tcc gcc gcc gac acc aat gct tgg aag tcc cgc agc atc tac 96
 Ile Val Ser Ala Ala Asp Thr Asn Ala Trp Lys Ser Arg Ser Ile Tyr
 20 25 30

ttt gtc ctg acg gat cgt att gcc cgc aac agc agc gac acg ggc ggc 144
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Ser	Ala	Cys	Ser	Asp	Leu	Gly	Asn	Tyr	Cys	Gly	Gly	Thr	Phe	Gln	Gly		
	50					55				60							
ctc	gag	tct	aag	ctc	gac	tac	atc	aag	gga	ctt	gga	ttc	gat	gcc	att		240
Leu	Glu	Ser	Lys	Leu	Asp	Tyr	Ile	Lys	Gly	Leu	Gly	Phe	Asp	Ala	Ile		
65					70					75					80		
tgg	att	acc	ccc	gtc	gtc	tca	a	gtatgttcat	cacggcttct	gccaacacag							292
Trp	Ile	Thr	Pro	Val	Val	Ser											
				85													
agcagtacta	atccaacttg	tag	ac	aag	gct	gct	gga	tac	cat	ggc	tac	tgg					344
				Asn	Lys	Ala	Ala	Gly	Tyr	His	Gly	Tyr	Trp				
						90					95						
gcc	gag	gac	ttg	tat	gcc	gtc	aac	tca	aac	tac	ggc	act	gct	gcc	gac		392
Ala	Glu	Asp	Leu	Tyr	Ala	Val	Asn	Ser	Asn	Tyr	Gly	Thr	Ala	Ala	Asp		
		100					105					110					
ttg	aag	agc	ttg	gtt	gcc	gct	gcc	cat	gcc	aag	gtatgtccgg	atttagtata					445
Leu	Lys	Ser	Leu	Val	Ala	Ala	Ala	His	Ala	Lys							
	115					120											
tatgttcata	gctagtagcat	ttgcttactc	atcatag	ggc	atc	tac	atg	atg	gtc								500
				Gly	Ile	Tyr	Met	Met	Val								
										125							
gac	gtt	gtc	gca	aac	cac	atg	ggc	cct	gga	gca	atc	aca	aac	aac	cgc		548
Asp	Val	Val	Ala	Asn	His	Met	Gly	Pro	Gly	Ala	Ile	Thr	Asn	Asn	Arg		
				135					140								
cct	gaa	cct	ctc	aac	cag	gct	tca	tca	tac	cac	cct	cct	tgc	aac	atc		596
Pro	Glu	Pro	Leu	Asn	Gln	Ala	Ser	Ser	Tyr	His	Pro	Pro	Cys	Asn	Ile		
			150					155					160				
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Asp	Tyr	Asn	Asn	Gln	Thr	Ser	Val	Glu	Val	Cys	Gln	Ile	Ala	Gly	Leu		
		165					170					175					
ccc	gac	atc	tac	acc	acc	aag	agc	gag	atc	cgc	acg	ctc	ctc	aac	acc		692
Pro	Asp	Ile	Tyr	Thr	Thr	Lys	Ser	Glu	Ile	Arg	Thr	Leu	Leu	Asn	Thr		
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tgg	gtc	aac	tgg	ctc	gta	aac	gag	tac	agc	ttc	gac	ggc	gtc	cgc	atc		740
Trp	Val	Asn	Trp	Leu	Val	Asn	Glu	Tyr	Ser	Phe	Asp	Gly	Val	Arg	Ile		
195					200					205					210		
gac	acc	gtc	aag	cac	gtc	gaa	aag	gac	ttt	tgg	cct	ggc	ttc	tct	gcc		788
Asp	Thr	Val	Lys	His	Val	Glu	Lys	Asp	Phe	Trp	Pro	Gly	Phe	Ser	Ala		
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gct	acc	ggc	gtc	tac	aac	att	ggc	gag	gtg	ttt	gac	gga	gac	cca	gcc		836
Ala	Thr	Gly	Val	Tyr	Asn	Ile	Gly	Glu	Val	Phe	Asp	Gly	Asp	Pro	Ala		
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tac	ctt	gcc	ccg	tac	gcc	aag	ctt	atg	ccc	ggc	ctc	ctc	aac	tac	gca		884
Tyr	Leu	Ala	Pro	Tyr	Ala	Lys	Leu	Met	Pro	Gly	Leu	Leu	Asn	Tyr	Ala		
		245					250					255					
gtc	tac	tac	ccg	atg	aac	aac	ttt	tac	cag	caa	acg	ggc	tct	tcc	cag		932
Val	Tyr	Tyr	Pro	Met	Asn	Asn	Phe	Tyr	Gln	Gln	Thr	Gly	Ser	Ser	Gln		
	260					265					270						
gcg	ctt	gta	gac	atg	atg	aac	act	gtc	agc	aac	acc	ttc	cct	gac	cca		980

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gac Asp	cag Gln	act Thr	ctg Leu 315	ctc Leu 315	aag Lys	aac Asn	gct Ala	ctt Leu	gct Ala 320	tat Tyr	gtc Val	atc Ile	ctc Leu	gca Ala 325	cgt Arg	1130
ggg Gly	atc Ile	ccc Pro	atc Ile 330	ttg Leu	tac Tyr	tat Tyr	ggg Gly	acc Thr 335	gag Glu	cag Gln	gga Gly	tac Tyr	gct Ala 340	ggg Gly	ggg Gly	1178
gac Asp	gac Asp	cca Pro 345	gct Ala	aac Asn	cga Arg	gag Glu	gat Asp 350	ctg Leu	tgg Trp	cgc Arg	agt Ser	ggc Gly 355	ttc Phe	aac Asn	acc Thr	1226
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gac Asp	acg Thr	gct Ala	tac Tyr	gcc Ala 395	tgg Trp	agc Ser	cgt Arg	gcc Ala	aac Asn 400	ggc Gly	aac Asn	ctg Leu	att Ile	gtc Val 405	ctc Leu	1370
acc Thr	acc Thr	aac Asn	gct Ala 410	ggc Gly	ggc Gly	aac Asn	tcc Ser	aac Asn 415	acc Thr	cag Gln	cac His	tgc Cys	ttc Phe 420	aac Asn	acg Thr	1418
caa Gln	aag Lys	gca Ala 425	aac Asn	ggc Gly	cgc Arg	tgg Trp	acc Thr 430	aac Asn	gtc Val	tac Tyr	ggc Gly	aac Asn 435	ggc Gly	gcc Ala	acc Thr	1466
gtc Val 440	tct Ser	gcc Ala	gat Asp	agc Ser	aac Asn	ggc Gly 445	caa Gln	atc Ile	tgc Cys	gtc Val	tcc Ser 450	gtc Val	aca Thr	aac Asn	ggc Gly	1514
gag Glu 455	ccc Pro	gtt Val	gtc Val	ctc Leu	ctc Leu 460	gcc Ala	ggc Gly	tcc Ser	gct Ala	acc Thr 465	ccc Pro	acc Thr	act Thr	ggc Gly	act Thr 470	1562
acc Thr	ctc Leu	tcc Ser	acc Thr	cgc Arg 475	acc Thr	gcc Ala	act Thr	gcc Ala	acc Thr 480	gcc Ala	aca Thr	cca Pro	acc Thr	gca Ala 485	tgc Cys	1610
ccc Pro	acc Thr	gcc Ala	gtc Val 490	tcc Ser	gtc Val	tcc Ser	ttc Phe 495	acc Thr	cac His	cgc Arg	gtc Val	acc Thr	act Thr 500	gtt Val	ccc Pro	1658
ggg Gly	gac Asp	acc Thr 505	atc Ile	aaa Lys	atc Ile	act Thr	ggc Gly 510	aac Asn	acg Thr	gcc Ala	cag Gln	cta Leu 515	ggg Gly	aac Asn	tgg Trp	1706
act Thr	ccc Pro 520	gcc Ala	aac Asn	ggc Gly	ctt Leu	gcc Ala 525	ttg Leu	tcc Ser	gca Ala	gct Ala	agc Ser 530	tac Tyr	aca Thr	tcc Ser	agc Ser	1754
aac	cct	atc	tgg	acc	att	acc	gtg	ccc	ctg	gcc	gct	gga	tcc	tcc	atc	1802

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Asn	Pro	Ile	Trp	Thr	Ile	Thr	Val	Pro	Leu	Ala	Ala	Gly	Ser	Ser	Ile		
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Ser	Tyr	Lys	Phe	Val	Lys	Ile	Asp	Ser	Gly	Gly	Thr	Val	Thr	Trp	Glu		
				555					560					565			
agt	gac	ccc	aac	agg	tca	tac	act	gcg	ccg	agc	tgc	cag	gcg	agt	gcc		1898
Ser	Asp	Pro	Asn	Arg	Ser	Tyr	Thr	Ala	Pro	Ser	Cys	Gln	Ala	Ser	Ala		
			570					575					580				
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Ser	Val	Asn	Ser	Ser	Trp	Gln	*										
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<210> 535
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 <213> Cochliobolus heterostrophus ATCC 48331

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 Phe Val Leu Thr Asp Arg Ile Ala Arg Asn Ser Ser Asp Thr Gly Gly
 35 40 45
 Ser Ala Cys Ser Asp Leu Gly Asn Tyr Cys Gly Gly Thr Phe Gln Gly
 50 55 60
 Leu Glu Ser Lys Leu Asp Tyr Ile Lys Gly Leu Gly Phe Asp Ala Ile
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 Trp Ile Thr Pro Val Val Ser
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 <211> 37
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 <213> Cochliobolus heterostrophus ATCC 48331

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 Val Asn Ser Asn Tyr Gly Thr Ala Ala Asp Leu Lys Ser Leu Val Ala
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 Ala Ala His Ala Lys
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 <211> 173
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

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 Ala Ile Thr Asn Asn Arg Pro Glu Pro Leu Asn Gln Ala Ser Ser Tyr
 20 25 30
 His Pro Pro Cys Asn Ile Asp Tyr Asn Asn Gln Thr Ser Val Glu Val
 35 40 45
 Cys Gln Ile Ala Gly Leu Pro Asp Ile Tyr Thr Thr Lys Ser Glu Ile
 50 55 60
 Arg Thr Leu Leu Asn Thr Trp Val Asn Trp Leu Val Asn Glu Tyr Ser
 65 70 75 80
 Phe Asp Gly Val Arg Ile Asp Thr Val Lys His Val Glu Lys Asp Phe
 85 90 95
 Trp Pro Gly Phe Ser Ala Ala Thr Gly Val Tyr Asn Ile Gly Glu Val
 100 105 110

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Phe Asp Gly Asp Pro Ala Tyr Leu Ala Pro Tyr Ala Lys Leu Met Pro
 115 120 125
 Gly Leu Leu Asn Tyr Ala Val Tyr Tyr Pro Met Asn Asn Phe Tyr Gln
 130 135 140
 Gln Thr Gly Ser Ser Gln Ala Leu Val Asp Met Met Asn Thr Val Ser
 145 150 155 160
 Asn Thr Phe Pro Asp Pro Ser Ala Leu Gly Thr Phe Leu
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<210> 538
 <211> 292
 <212> PRT
 <213> Cochliobolus heterostrophus ATCC 48331

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 Ile Leu Tyr Tyr Gly Thr Glu Gln Gly Tyr Ala Gly Gly Asp Asp Pro
 35 40 45
 Ala Asn Arg Glu Asp Leu Trp Arg Ser Gly Phe Asn Thr Asn Ala Asn
 50 55 60
 Leu Tyr Gln Ala Ile Lys Lys Leu Thr Ala Ala Arg Gln Ala Ala Gly
 65 70 75 80
 Gly Leu Ala Gly Asn Asp His Val His Leu Tyr Val Ala Asp Thr Ala
 85 90 95
 Tyr Ala Trp Ser Arg Ala Asn Gly Asn Leu Ile Val Leu Thr Thr Asn
 100 105 110
 Ala Gly Gly Asn Ser Asn Thr Gln His Cys Phe Asn Thr Gln Lys Ala
 115 120 125
 Asn Gly Arg Trp Thr Asn Val Tyr Gly Asn Gly Ala Thr Val Ser Ala
 130 135 140
 Asp Ser Asn Gly Gln Ile Cys Val Ser Val Thr Asn Gly Glu Pro Val
 145 150 155 160
 Val Leu Leu Ala Gly Ser Ala Thr Pro Thr Thr Gly Thr Thr Leu Ser
 165 170 175
 Thr Arg Thr Ala Thr Ala Thr Ala Thr Thr Ala Cys Pro Thr Ala
 180 185 190
 Val Ser Val Ser Phe Thr His Arg Val Thr Thr Val Pro Gly Asp Thr
 195 200 205
 Ile Lys Ile Thr Gly Asn Thr Ala Gln Leu Gly Asn Trp Thr Pro Ala
 210 215 220
 Asn Gly Leu Ala Leu Ser Ala Ala Ser Tyr Thr Ser Ser Asn Pro Ile
 225 230 235 240
 Trp Thr Ile Thr Val Pro Leu Ala Ala Gly Ser Ser Ile Ser Tyr Lys
 245 250 255
 Phe Val Lys Ile Asp Ser Gly Gly Thr Val Thr Trp Glu Ser Asp Pro
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 Asn Arg Ser Tyr Thr Ala Pro Ser Cys Gln Ala Ser Ala Ser Val Asn
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 Ser Ser Trp Gln
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 cgcaacagca gcgacacggg cggtcagcg tgcagcgacc tcggcaacta ctgcggtgga 180
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 gcccatgcca agggcatcta catgatggtc gacgttgctg caaaccacat gggtcctgga 420
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aacgagtaca gcttcgacgg tgtccgcacg gacaccgtca agcacgtcga aaaggacttt 660
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<210> 540
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 <213> Cochliobolus heterostrophus ATCC 48331

<220>
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<221> DOMAIN
 <222> (33)...(373)
 <223> Alpha amylase, catalytic domain

<221> DOMAIN
 <222> (489)...(586)
 <223> Starch binding domain

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Phe Val Leu Thr Asp Arg Ile Ala Arg Asn Ser Ser Asp Thr Gly Gly
 35     40     45
Ser Ala Cys Ser Asp Leu Gly Asn Tyr Cys Gly Thr Phe Gln Gly
 50     55     60
Leu Glu Ser Lys Leu Asp Tyr Ile Lys Gly Leu Gly Phe Asp Ala Ile
 65     70     75     80
Trp Ile Thr Pro Val Val Ser Asn Lys Ala Ala Gly Tyr His Gly Tyr
 85     90     95
Trp Ala Glu Asp Leu Tyr Ala Val Asn Ser Asn Tyr Gly Thr Ala Ala
100    105    110
Asp Leu Lys Ser Leu Val Ala Ala His Ala Lys Gly Ile Tyr Met
115    120    125
Met Val Asp Val Val Ala Asn His Met Gly Pro Gly Ala Ile Thr Asn
130    135    140
Asn Arg Pro Glu Pro Leu Asn Gln Ala Ser Ser Tyr His Pro Pro Cys
145    150    155    160
Asn Ile Asp Tyr Asn Asn Gln Thr Ser Val Glu Val Cys Gln Ile Ala
165    170    175
Gly Leu Pro Asp Ile Tyr Thr Thr Lys Ser Glu Ile Arg Thr Leu Leu
180    185    190
Asn Thr Trp Val Asn Trp Leu Val Asn Glu Tyr Ser Phe Asp Gly Val
195    200    205
Arg Ile Asp Thr Val Lys His Val Glu Lys Asp Phe Trp Pro Gly Phe

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210 215 220
 Ser Ala Ala Thr Gly Val Tyr Asn Ile Gly Glu Val Phe Asp Gly Asp
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 260 265 270
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48

gtt gat cag aga aag gat ctg atc tcc aca agt aac ggg gtg atg tcg
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ccaag Ala		cc	cat His	cac His 40	tcc Ser	cat His	ttt Phe	acc Thr	atc Ile 45	atc Ile	tct Ser	tct Ser	att Ile	ctc Leu	tcc Ser 50	ctc Leu	198
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gtc Val	tac Tyr 85	cca Pro	gcc Ala	agc Ser	ttc Phe	aaa Lys 90	gac Asp	tcc Ser	aac Asn	aac Asn	gac Asp 95	ggc Gly	tgg Trp	ggc Gly	gat Asp	342	
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Pro	Gly	Leu	Ile	Ser	Lys	Leu	Asp	Tyr	Leu	Ser	Asp	Leu	Gly	Ile	Asp	
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Val	Val	Trp	Val	Ser	Pro	Ile	Phe	Glu	Ser	Pro	Gln	Lys	Asp	Met	Gly	
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Glu	Asp	Val	Asp	Ile	Leu	Val	Lys	Glu	Cys	His	Ala	Arg	Gly	Leu	Lys	
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1001827087_1.txt

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His	Thr	Gln 195	Glu	Tyr	Tyr	Leu	His 200	Leu	Tyr	Ala	Pro	Asp 205	Gln	Pro	Asp
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<223> Alpha amylase, catalytic domain

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Asp Glu His Gln Tyr Ile Gly Lys Gln Pro Trp Trp Lys Ala Ala Ser
65     70     75     80
Phe Tyr Gln Val Tyr Pro Ala Ser Phe Lys Asp Ser Asn Asn Asp Gly
85     90     95
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Lys	Thr	Leu	Ala	Thr	Trp	Gln	Thr	Thr	Leu	Thr	Gly	Thr	Leu	Phe	Leu
		435					440					445			
Tyr	Gln	Gly	Gln	Glu	Ile	Gly	Met	Thr	Asn	Met	Pro	Lys	Ser	Trp	Gly
	450					455					460				
Ile	Glu	Glu	Tyr	Lys	Asp	Ile	Glu	Ser	Ser	Asn	Phe	Tyr	Ala	Glu	Ala
	465				470					475					480
Val	Ala	Ser	Gly	Asp	Glu	Lys	Arg	Val	Arg	Asp	Thr	Met	His	Gly	Leu
				485					490					495	
Gln	Ile	Met	Ala	Arg	Asp	His	Ser	Arg	Ile	Pro	Phe	Gln	Trp	Asp	Asp
			500					505					510		
Ser	Pro	Asn	Ala	Gly	Phe	Ala	Asp	Ala	Ser	Ala	Lys	Pro	Trp	Met	Arg
		515					520					525			
Val	His	Asp	Asp	Tyr	Arg	Asp	Ile	Asn	Val	Ala	Lys	Gln	Val	Lys	Asp
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Pro	Asn	Ser	Ile	Leu	Ser	Phe	Tyr	Lys	Ala	Met	Leu	Arg	Leu	Arg	Lys
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Glu	Tyr	Gln	Asp	Ile	Phe	Val	Phe	Gly	Ser	Phe	Lys	Leu	Leu	Asp	Pro
				565					570					575	
Glu	Asp	Glu	Ser	Leu	Phe	Cys	Tyr	Val	Lys	Glu	Ser	Val	Ala	Pro	Val
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Lys	Gly	Arg	Arg	Gly	Glu	Lys	Arg	Lys	Ala	Val	Val	Val	Leu	Asn	Met
		595					600					605			
Ser	Arg	Glu	Asp	Arg	Leu	Gly	Pro	Asp	Val	Pro	Thr	Val	Leu	Gly	Cys
	610					615					620				
Ser	His	Glu	Asp	Val	Arg	Leu	Met	Ala	Tyr	Thr	Arg	Glu	Thr	Ala	Gly
	625				630					635					640
Gly	Leu	Lys	Val	Gly	Arg	Pro	Met	Leu	Ala	Gly	Trp	Glu	Ser	Arg	Val
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<213> Fungi

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<223> Exon

<400> 546

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1				5					10					15		

agc	act	cag	atc	ctc	caa	gct	ggt	caa	cct	att	atg	cat	aca	ttc	gct	96
Ser	Thr	Gln	Ile	Leu	Gln	Ala	Val	Gln	Pro	Ile	Met	His	Thr	Phe	Ala	
			20					25					30			

ccc	ctc	ctt	ttg	ggt	ggc	agt	ctt	gcc	ctc	caa	act	atc	tat	gct	ttg	144
Pro	Leu	Leu	Leu	Val	Gly	Ser	Leu	Ala	Leu	Gln	Thr	Ile	Tyr	Ala	Leu	
		35					40					45				

cct	ggc	ccc	agt	aaa	ggt	gag	caa	cgc	gcc	gaa	atc	cta	aaa	cgc	tca	192
Pro	Gly	Pro	Ser	Lys	Val	Glu	Gln	Arg	Ala	Glu	Ile	Leu	Lys	Arg	Ser	
	50					55					60					

gtg	gat	tct	ttt	att	gcc	acc	gaa	agt	cct	att	gct	ttc	aga	aat	ctt	240
Val	Asp	Ser	Phe	Ile	Ala	Thr	Glu	Ser	Pro	Ile	Ala	Phe	Arg	Asn	Leu	
65					70					75					80	

tta	tgc	aat	atc	ggc	gca	gat	gga	gct	tgt	gct	tca	ggg	gca	gct	tcg	288
Leu	Cys	Asn	Ile	Gly	Ala	Asp	Gly	Ala	Cys	Ala	Ser	Gly	Ala	Ala	Ser	
				85					90					95		

ggt	att	gtg	ggt	gct	tcg	cct	gac	aaa	gtg	agc	cca	gat	t	g	tttagtatttt	338
Gly	Ile	Val	Val	Ala	Ser	Pro	Asp	Lys	Val	Ser	Pro	Asp				
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catgatgccg	tttgaataca	tggaattctg	atatgtaaac	ag	at	ttt	tat	act	tg							394
						Tyr	Phe	Tyr	Thr	Trp						
						110										

act	cga	gac	tcc	gca	ttg	gta	ttc	aaa	gcc	ctt	ggt	gat	acg	ttt	atc	442
Thr	Arg	Asp	Ser	Ala	Leu	Val	Phe	Lys	Ala	Leu	Val	Asp	Thr	Phe	Ile	
115					120					125					130	

agc	agc	agc	tat	tct	gca	tca	ctt	caa	cag	gaa	att	gaa	aat	tac	atc	490
Ser	Ser	Ser	Tyr	Ser	Ala	Ser	Leu	Gln	Gln	Glu	Ile	Glu	Asn	Tyr	Ile	
				135					140					145		

agc	tcc	caa	gct	gga	ctt	caa	acc	gtg	tca	aac	cct	tca	ggt	ggt	tta	538
Ser	Ser	Gln	Ala	Gly	Leu	Gln	Thr	Val	Ser	Asn	Pro	Ser	Gly	Gly	Leu	
			150					155					160			

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tca tcc ggt ggt ctg ggc gag ccc aaa ttt aat gcg gat aag tcg gca Ser Ser Gly Gly Leu Gly Glu Pro Lys Phe Asn Ala Asp Lys Ser Ala 165 170 175	586
ttc aca g gtacgaaaat tcgaaaactc tatcctgatg gtgactgaca tgtgaactat Phe Thr 180	643
acag gg tct tgg gga aga cct caa cga gat gga ccc gct tta cgt gct Gly Ser Trp Gly Arg Pro Gln Arg Asp Gly Pro Ala Leu Arg Ala 185 190 195	691
act gct atg att acc tat tca aaa tgg ttg att gcc aac gga tac acc Thr Ala Met Ile Thr Tyr Ser Lys Trp Leu Ile Ala Asn Gly Tyr Thr 200 205 210	739
tcc act gtg caa act att gtt tgg ccg atc atc aga aat gat ctg tct Ser Thr Val Gln Thr Ile Val Trp Pro Ile Ile Arg Asn Asp Leu Ser 215 220 225	787
tat gtt act caa tat t gtaagtcaac gggaagaaaa ccaaattagt ctaaaaatag Tyr Val Thr Gln Tyr 230	843
atgaccctct tttcttggag cttttgatta acttaattca tag gg aac caa act Trp Asn Gln Thr 235	897
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gct gct caa cat cgt gct ctt gtt gaa ggt agt gca ctg gcc gct caa Ala Ala Gln His Arg Ala Leu Val Glu Gly Ser Ala Leu Ala Ala Gln 255 260 265	993
att ggt caa agc tgt acc tat tgt gat tct ca gtaagcacat atgaaagaat Ile Gly Gln Ser Cys Thr Tyr Cys Asp Ser Gln 270 275	1045
atattcacat gcagggtacta atacattttg cag a gcc cca cag gtt ctt tgt Ala Pro Gln Val Leu Cys 280 285	1097
ttc ctc caa agc ttc tgg agt tca tcc tcg gga tac atc att tcc aac Phe Leu Gln Ser Phe Trp Ser Ser Ser Ser Gly Tyr Ile Ile Ser Asn 290 295 300	1145
atc aac caa aac agc gga aga aac gga aaa gat gca aac agt atc ttg Ile Asn Gln Asn Ser Gly Arg Asn Gly Lys Asp Ala Asn Ser Ile Leu 305 310 315	1193
agt tct atc cag acc ttt gat cct acc gct gca tgt gat gcc acc acg Ser Ser Ile Gln Thr Phe Asp Pro Thr Ala Ala Cys Asp Ala Thr Thr 320 325 330	1241
ttc caa cca tgc tca gac cgt gct ttg gca aat cac aaa gtt gtg act Phe Gln Pro Cys Ser Asp Arg Ala Leu Ala Asn His Lys Val Val Thr 335 340 345	1289
gat tct ttc aga tct atc tac agc atc aac tct ggg att gcc gaa ggt Asp Ser Phe Arg Ser Ile Tyr Ser Ile Asn Ser Gly Ile Ala Glu Gly 350 355 360 365	1337
gtc gcc gtt tct gtt gga cgc tac cca gaa gat tct tac tat ggt ggc Val Ala Val Ser Val Gly Arg Tyr Pro Glu Asp Ser Tyr Tyr Gly Gly 370 375 380	1385

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gct Ala	tta Leu	tac Tyr 400	acc Thr	tgg Trp	aat Asn	aga Arg	att Ile 405	ggc Gly	tct Ser	atc Ile	acc Thr	gta Val 410	act Thr	tcc Ser	act Thr	1481
tcc Ser	cta Leu 415	gca Ala	ttc Phe	ttc Phe	aaa Lys	gat Asp 420	ttc Phe	agc Ser	tct Ser	tcc Ser	atc Ile 425	aca Thr	gcc Ala	gga Gly	aca Thr	1529
tac Tyr 430	gca Ala	tcc Ser	tcc Ser	aca Thr	tca Ser 435	acc Thr	tac Tyr	acc Thr	act Thr	ctc Leu 440	tac Tyr	aac Asn	gcc Ala	atc Ile	aaa Lys 445	1577
acc Thr	tat Tyr	gcc Ala	gac Asp	ggc Gly 450	tac Tyr	gtc Val	aat Asn	gtc Val	gtc Val 455	gca Ala	acc Thr	tac Tyr	gca Ala	caa Gln 460	gcc Ala	1625
aac Asn	ggt Gly	tcc Ser	ctc Leu 465	tca Ser	gaa Glu	caa Gln	ttc Phe	aac Asn 470	aaa Lys	gcc Ala	ggc Gly	ggc Gly	gct Ala 475	ccc Pro	ctc Leu	1673
tca Ser	gcc Ala	tac Tyr 480	gac Asp	ctc Leu	acc Thr	tgg Trp	tcc Ser 485	tat Tyr	gct Ala	gcc Ala	ttc Phe	ctc Leu 490	acc Thr	gcc Ala	gct Ala	1721
gcc Ala	cgt Arg 495	cgc Arg	gca Ala	ggt Gly	gtc Val	gtc Val 500	cca Pro	tac Tyr	tca Ser	tgg Trp	ggt Gly 505	gag Glu	cca Pro	tcc Ser	gcc Ala	1769
tcc Ser 510	agc Ser	gtc Val	cca Pro	gga Gly	acc Thr 515	tgt Cys	tct Ser	gct Ala	acc Thr	tcg Ser 520	gcc Ala	atc Ile	gga Gly	acc Thr	tat Tyr 525	1817
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gga Gly	gga Gly	gga Gly	agt Ser 545	acc Thr	ccc Pro	aca Thr	acc Thr	agc Ser 550	aca Thr	tcc Ser	gcc Ala	acc Thr	acc Thr 555	aag Lys	aca Thr	1913
act Thr	tcc Ser	acc Thr 560	tgc Ser	acc Thr	acc Thr	act Thr	tct Ser 565	tca Ser	tgc Cys	gca Ala	aca Thr	gcc Ala 570	act Thr	tca Ser	gta Val	1961
gct Ala	gtt Val 575	act Thr	ttc Phe	aat Asn	gaa Glu	ttg Leu 580	gtc Val	act Thr	acc Thr	tcc Ser	ttt Phe 585	ggc Gly	caa Gln	acg Thr	atc Ile	2009
aaa Lys 590	ctc Leu	gct Ala	ggt Gly	tcc Ser	gtc Val 595	tct Ser	caa Gln	ttg Leu	gga Gly	agt Ser 600	tgg Trp	gca Ala	ccg Pro	gct Ala 605	tct Ser 605	2057
gcg Ala	att Ile	gca Ala	ttg Leu	agc Ser 610	gcc Ala	gcg Ala	aag Lys	tac Tyr	acc Thr 615	gct Ala	agt Ser	aat Asn	ccg Pro	ttg Leu 620	tgg Trp	2105
acc Thr	gtg Val	act Thr	gtt Val 625	aac Asn	cta Leu	cca Pro	gct Ala	ggc Gly 630	acg Thr	acg Thr	gta Val	ttg Leu	tat Tyr 635	aag Lys	ttt Phe	2153
att Ile	aac Asn	gtg Val 640	gcg Ala	agt Ser	gat Asp	ggg Gly	acg Thr 645	gtt Val	acc Thr	tgg Trp	cag Gln	gcg Ala 650	gat Asp	cca Pro	aat Asn	2201

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655 660 665

ggt agc tgg taa 2261
Gly Ser Trp *
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20 25 30
Pro Leu Leu Val Gly Ser Leu Ala Leu Gln Thr Ile Tyr Ala Leu
35 40 45
Pro Gly Pro Ser Lys Val Glu Gln Arg Ala Glu Ile Leu Lys Arg Ser
50 55 60
Val Asp Ser Phe Ile Ala Thr Glu Ser Pro Ile Ala Phe Arg Asn Leu
65 70 75 80
Leu Cys Asn Ile Gly Ala Asp Gly Ala Cys Ala Ser Gly Ala Ala Ser
85 90 95
Gly Ile Val Val Ala Ser Pro Asp Lys Val Ser Pro Asp
100 105

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1 5 10 15
Val Asp Thr Phe Ile Ser Ser Ser Tyr Ser Ala Ser Leu Gln Gln Glu
20 25 30
Ile Glu Asn Tyr Ile Ser Ser Gln Ala Gly Leu Gln Thr Val Ser Asn
35 40 45
Pro Ser Gly Gly Leu Ser Ser Gly Gly Leu Gly Glu Pro Lys Phe Asn
50 55 60
Ala Asp Lys Ser Ala Phe Thr
65 70

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<212> PRT
<213> Fungi

<400> 549
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20 25 30
Thr Val Gln Thr Ile Val Trp Pro Ile Ile Arg Asn Asp Leu Ser Tyr
35 40 45
Val Thr Gln Tyr
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<400> 550

30 May 2017

2013201807

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Asp	Ala	Asn 35	Ser	Ile	Leu	Ser	Ser 40	Ile	Gln	Thr	Phe	Asp 45	Pro	Thr	Ala
Ala	Cys 50	Asp	Ala	Thr	Thr	Phe 55	Gln	Pro	Cys	Ser	Asp 60	Arg	Ala	Leu	Ala
Asn 65	His	Lys	Val	Val	Thr 70	Asp	Ser	Phe	Arg	Ser 75	Ile	Tyr	Ser	Ile 80	Asn 80
Ser	Gly	Ile	Ala	Glu 85	Gly	Val	Ala	Val	Ser 90	Val	Gly	Arg	Tyr	Pro 95	Glu
Asp	Ser	Tyr	Tyr 100	Gly	Gly	Asn	Pro	Trp 105	Tyr	Leu	Asn	Thr	Leu 110	Ala	Ala
Ala	Glu	Gln 115	Leu	Tyr	Asp	Ala	Leu 120	Tyr	Thr	Trp	Asn	Arg 125	Ile	Gly	Ser
Ile	Thr 130	Val	Thr	Ser	Thr	Ser 135	Leu	Ala	Phe	Phe	Lys 140	Asp	Phe	Ser	Ser
Ser 145	Ile	Thr	Ala	Gly	Thr 150	Tyr	Ala	Ser	Ser	Thr 155	Ser	Thr	Tyr	Thr 160	Thr
Leu	Tyr	Asn	Ala	Ile 165	Lys	Thr	Tyr	Ala	Asp 170	Gly	Tyr	Val	Asn 175	Val 175	Val
Ala	Thr	Tyr	Ala 180	Gln	Ala	Asn	Gly	Ser 185	Leu	Ser	Glu	Gln	Phe 190	Asn	Lys
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Ala	Phe 210	Leu	Thr	Ala	Ala	Ala 215	Arg	Arg	Ala	Gly	Val 220	Val	Pro	Tyr	Ser
Trp 225	Gly	Glu	Pro	Ser	Ala 230	Ser	Ser	Val	Pro	Gly 235	Thr	Cys	Ser	Ala	Thr 240
Ser	Ala	Ile	Gly	Thr 245	Tyr	Ser	Thr	Ala	Thr 250	Ala	Thr	Ser	Trp	Pro 255	Ala
Ser	Gln	Thr	Pro 260	Ser	Gly	Gly	Gly	Gly 265	Ser	Thr	Pro	Thr	Thr 270	Ser	Thr
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Ala	Thr 290	Ala	Thr	Ser	Val	Ala 295	Val	Thr	Phe	Asn	Glu 300	Leu	Val	Thr	Thr
Ser 305	Phe	Gly	Gln	Thr	Ile 310	Lys	Leu	Ala	Gly	Ser 315	Val	Ser	Gln	Leu	Gly 320
Ser	Trp	Ala	Pro	Ala 325	Ser	Ala	Ile	Ala	Leu 330	Ser	Ala	Ala	Lys	Tyr 335	Thr
Ala	Ser	Asn	Pro 340	Leu	Trp	Thr	Val	Thr 345	Val	Asn	Leu	Pro	Ala 350	Gly	Thr
Thr	Val 355	Leu	Tyr	Lys	Phe	Ile	Asn 360	Val	Ala	Ser	Asp	Gly 365	Thr	Val	Thr
Trp	Gln 370	Ala	Asp	Pro	Asn	Lys 375	Ser	Tyr	Thr	Val	Pro 380	Val	Gly	Cys	Ala
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<210> 552
 <211> 2019
 <212> DNA
 <213> Fungi


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<210> 553
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<223> Glycosyl hydrolases family 15

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<221> DOMAIN
<222> (572)...(668)
<223> Starch binding domain

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Pro Leu Leu Leu Val Gly Ser Leu Ala Leu Gln Thr Ile Tyr Ala Leu
35     40     45
Pro Gly Pro Ser Lys Val Glu Gln Arg Ala Glu Ile Leu Lys Arg Ser
50     55     60
Val Asp Ser Phe Ile Ala Thr Glu Ser Pro Ile Ala Phe Arg Asn Leu
65     70     75     80
Leu Cys Asn Ile Gly Ala Asp Gly Ala Cys Ala Ser Gly Ala Ala Ser
85     90     95
Gly Ile Val Val Ala Ser Pro Asp Lys Val Ser Pro Asp Tyr Phe Tyr

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Tyr	130	Ile	Ser	Ser	Gln	Ala	135	Gly	Leu	Gln	Thr	Val	Ser	Asn	Pro	Ser	Gly	
145	Gly	Leu	Ser	Ser	Gly	150	Gly	Leu	Gly	Glu	Pro	155	Lys	Phe	Asn	Ala	Asp	
Ser	Ala	Phe	Thr	Gly	Ser	Trp	Gly	Arg	170	Pro	Gln	Arg	Asp	Gly	Pro	Ala		
Leu	Arg	Ala	180	Thr	Ala	Met	Ile	Thr	185	Tyr	Ser	Lys	Trp	Leu	190	Ile	Ala	
Gly	Tyr	195	Thr	Ser	Thr	Val	Gln	Thr	200	Ile	Val	Trp	Pro	205	Ile	Ile	Arg	
225	Asp	Leu	Ser	Tyr	Val	Thr	Gln	Tyr	Trp	Asn	Gln	Thr	Gly	Phe	Asp	Leu		
Trp	Glu	Glu	Val	Gln	Gly	Ser	Ser	Phe	235	Phe	Thr	Val	Ala	Ala	Gln	His		
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Cys	Thr	Tyr	Cys	Asp	Ser	Gln	Ala	Pro	265	Gln	Val	Leu	Cys	Phe	Leu	Gln		
Ser	Phe	Trp	Ser	Ser	Ser	Ser	295	Gly	Tyr	Ile	Ile	Ser	300	Asn	Ile	Asn	Gln	
Asn	Ser	Gly	Arg	Asn	Gly	Lys	Asp	Ala	Asn	Ser	315	Ile	Leu	Ser	Ser	Ile		
Gln	Thr	Phe	Asp	Pro	Thr	Ala	Ala	Cys	330	Asp	Ala	Thr	Thr	Phe	Gln	Pro		
Cys	Ser	Asp	Arg	Ala	Leu	Ala	Asn	His	345	Lys	Val	Val	Thr	Asp	Ser	Phe		
Arg	Ser	Ile	Tyr	Ser	Ile	Asn	Ser	Gly	360	Ile	Ala	Glu	Gly	365	Val	Ala	Val	
Ser	Val	Gly	Arg	Tyr	Pro	Glu	Asp	Ser	375	Tyr	Tyr	Gly	Gly	Asn	Pro	Trp		
Tyr	Leu	Asn	Thr	Leu	Ala	Ala	Ala	Glu	Gln	Leu	Tyr	Asp	Ala	Leu	Tyr			
385	Thr	Trp	Asn	Arg	Ile	Gly	Ser	Ile	Thr	Val	Thr	Ser	Thr	Ser	Leu	Ala		
Phe	Phe	Lys	Asp	Phe	Ser	Ser	Ser	Ile	Thr	Ala	Gly	Thr	Tyr	430	Ala	Ser		
Ser	Thr	Ser	Thr	Tyr	Thr	Thr	Leu	Tyr	440	Asn	Ala	Ile	Lys	Thr	Tyr	Ala		
Asp	Gly	Tyr	Val	Asn	Val	Val	Ala	Thr	Tyr	Ala	Gln	Ala	Asn	Gly	Ser			
Leu	Ser	Glu	Gln	Phe	Asn	Lys	Ala	Gly	Gly	Ala	Pro	Leu	Ser	Ala	Tyr			
465	Asp	Leu	Thr	Trp	Ser	Tyr	Ala	Ala	Phe	Leu	Thr	Ala	Ala	Ala	Arg			
Ala	Gly	Val	Val	Pro	Tyr	Ser	Trp	Gly	505	Glu	Pro	Ser	Ala	Ser	Ser	Val		
Pro	Gly	Thr	Cys	Ser	Ala	Thr	Ser	Ala	520	Ile	Gly	Thr	Tyr	Ser	Thr	Ala		
Thr	Ala	Thr	Ser	Trp	Pro	Ala	Ser	Gln	Thr	Pro	Ser	540	Gly	Gly	Gly	Gly		
Ser	Thr	Pro	Thr	Thr	Ser	Thr	Ser	Ala	Thr	Thr	Lys	Thr	Thr	Ser	Thr			
545	Ser	Thr	Thr	Thr	Ser	Ser	Cys	Ala	Thr	Ala	Thr	Ser	Val	Ala	Val	Thr		
Phe	Asn	Glu	Leu	Val	Thr	Thr	Ser	Phe	570	Gly	Gln	Thr	Ile	Lys	Leu	Ala		
Gly	Ser	Val	Ser	Gln	Leu	Gly	Ser	Trp	585	Ala	Pro	Ala	Ser	Ala	Ile	Ala		
Leu	Ser	Ala	Ala	Lys	Tyr	Thr	Ala	Ser	600	Asn	Pro	Leu	Trp	Thr	Val	Thr		
Val	610	Asn	Leu	Pro	Ala	Gly	Thr	Thr	Val	Leu	Tyr	Lys	Phe	Ile	Asn	Val		
625	Ala	Ser	Asp	Gly	Thr	Val	Thr	Trp	Gln	Ala	Asp	Pro	Asn	Lys	Ser	Tyr		

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660 665 670

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<211> 1909
<212> DNA
<213> *Cochliobolus heterostrophus* ATCC 48331

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<223> Exon

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Ala Ala Ala Met Thr Ile Thr Pro Arg Pro Trp Trp Lys Asp Ala Val
20 25 30

gtc tac cag ata tac ccc gcc tcg ttc aag gac agc aac ggc gac ggc 144
Val Tyr Gln Ile Tyr Pro Ala Ser Phe Lys Asp Ser Asn Gly Asp Gly
35 40 45

att ggc gac ctc aat ggc atc atc tcg gaa ctc gac tac atc cgc tcc 192
Ile Gly Asp Leu Asn Gly Ile Ile Ser Glu Leu Asp Tyr Ile Arg Ser
50 55 60

att ggc gtc gat gtc atc tgg gtc tgc ccc atg tac gac tcg cct cag 240
Ile Gly Val Asp Val Ile Trp Val Cys Pro Met Tyr Asp Ser Pro Gln
65 70 75 80

gtc gac atg ggc tac gac atc cgc aac tac gag gat gtc tac aga cca 288
Val Asp Met Gly Tyr Asp Ile Arg Asn Tyr Glu Asp Val Tyr Arg Pro
85 90 95

tat gga acc gtc cag gac atg cag cgg ctg att gac gag acc cac tca 336
Tyr Gly Thr Val Gln Asp Met Gln Arg Leu Ile Asp Glu Thr His Ser
100 105 110

cgg ggc atg aag atc atc ctg gat ctc gtc gtc aac cac act tca gat 384
Arg Gly Met Lys Ile Ile Leu Asp Leu Val Val Asn His Thr Ser Asp
115 120 125

cag gtatgctcgc tctagtcacg tgatttgctg tggcattggc catcacatgc 437
Gln

tcactgccta cag cac caa tgg ttc caa gaa tca cgc tcg tca aag gac 486
His Gln Trp Phe Gln Glu Ser Arg Ser Ser Lys Asp
130 135 140

aat cca aag cgt gac tgg tac att tgg cgt ccc gcc cgc tac gtc gat 534
Asn Pro Lys Arg Asp Trp Tyr Ile Trp Arg Pro Ala Arg Tyr Val Asp
145 150 155

ggc gta cgc aaa ccc ccc aac aac tgg gtc tcc aac ttc acc ggc agt 582

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Gly	Val	Arg 160	Lys	Pro	Pro	Asn	Asn 165	Trp	Val	Ser	Asn	Phe 170	Thr	Gly	Ser		
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Val	Trp	Glu	Trp	Asp	Glu	His 180	Thr	Gln	Glu	Tyr	Tyr 185	Leu	His	Leu	Phe		
tgt	cct	gag	cag	ccc	gac	ctc	aac	tgg	gaa	aat	cca	gag	aca	cgt	aaa	678	
Cys 190	Pro	Glu	Gln	Pro	Asp 195	Leu	Asn	Trp	Glu	Asn 200	Pro	Glu	Thr	Arg	Lys 205		
gct	ata	tac	gag	tcg	gcc	atg	gag	ttc	tgg	ctc	aag	cgt	ggt	gtc	gac	726	
Ala	Ile	Tyr	Glu	Ser 210	Ala	Met	Glu	Phe	Trp 215	Leu	Lys	Arg	Gly	Val 220	Asp		
ggc	ttc	cgc	gtt	gat	acc	gtc	aac	atg	tac	agt	aag	ggc	gat	atg	cgt	774	
Gly	Phe	Arg	Val 225	Asp	Thr	Val	Asn	Met 230	Tyr	Ser	Lys	Gly	Asp 235	Met	Arg		
gac	gca	ccc	atc	acc	gat	ccg	gga	tca	gag	tgg	cag	ttt	gcc	ggc	tac	822	
Asp	Ala	Pro 240	Ile	Thr	Asp	Pro	Gly 245	Ser	Glu	Trp	Gln	Phe 250	Ala	Gly	Tyr		
cag	tac	tgc	aac	ggc	ccg	cga	atg	gat	gag	ttc	ctg	ggc	gag	atg	aat	870	
Gln	Tyr 255	Cys	Asn	Gly	Pro	Arg 260	Met	Asp	Glu	Phe	Leu 265	Gly	Glu	Met	Asn		
cag	att	ctg	gaa	aag	tat	gat	gcc	atg	aca	gtc	ggc	gag	tgt	cca	cac	918	
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aca	ctt	gat	atc	aac	aga	gta	cac	caa	tac	gta	agc	gcc	aag	gag	aag	966	
Thr	Leu	Asp	Ile	Asn 290	Arg	Val	His	Gln	Tyr 295	Val	Ser	Ala	Lys	Glu 300	Lys		
cga	ctc	tcc	atg	gtg	ttt	caa	ttc	gta	agt	ggac	ctg	gcc	ctat	tgca	ataacc	1020	
Arg	Leu	Ser	Met 305	Val	Phe	Gln	Phe										
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						Asp 310	Val	Val	Asp	Val	Gly 315	Gln	Gly	Pro	Tyr	Lys 320	
ttc	caa	acc	acg	ccc	aag	aac	tgg	aca	ctg	cca	caa	ctc	aag	cgg	gca	1120	
Phe	Gln	Thr	Thr	Pro 325	Lys	Asn	Trp	Thr	Leu 330	Pro	Gln	Leu	Lys	Arg 335	Ala		
ata	gcc	cgc	act	caa	gat	ctg	att	cgg	cct	cca	tct	gac	agc	tgg	acc	1168	
Ile	Ala	Arg	Thr 340	Gln	Asp	Leu	Ile	Arg 345	Pro	Pro	Ser	Asp	Ser 350	Trp	Thr		
act	gct	ttt	ctg	gag	aac	cac	gac	caa	gct	cgg	tca	att	acg	cgc	ttc	1216	
Thr	Ala	Phe 355	Leu	Glu	Asn	His	Asp 360	Gln	Ala	Arg	Ser	Ile 365	Thr	Arg	Phe		
act	tct	gac	gcc	cca	cag	cac	cgt	gtc	gct	ggc	ggc	aag	atg	ctt	gcc	1264	
Thr	Ser 370	Asp	Ala	Pro	Gln	His 375	Arg	Val	Ala	Gly	Gly 380	Lys	Met	Leu	Ala		
ctc	atg	ctg	agc	gct	ctt	agc	ggc	acg	ctg	ttc	atc	tac	caa	ggc	cag	1312	
Leu	Met	Leu	Ser	Ala 390	Leu	Ser	Gly	Thr	Leu	Phe 395	Ile	Tyr	Gln	Gly	Gln 400		
gaa	atc	ggc	atg	aca	aac	ttc	ccc	gaa	tcg	tgg	gac	atg	agc	gaa	tac	1360	
Glu	Ile	Gly	Met	Thr 405	Asn	Phe	Pro	Glu	Ser 410	Trp	Asp	Met	Ser	Glu 415	Tyr		
aag	gac	gtc	gag	tca	tcc	aat	tac	tac	aag	atg	gtt	gcc	aag	cgc	acc	1408	

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Asn	Asn	Asp	Val	Asp	Ala	Leu	Ala	Ala	Ala	His	Lys	Ser	Leu	Gln	His		
		435					440					445					
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Leu	Ala	Arg	Asp	His	Ser	Arg	Val	Pro	Met	Ser	Trp	Ser	Thr	Ala	Pro		
	450					455					460						
tac	aat	ggc	ttc	agt	ccg	ccc	gat	gca	aaa	gac	aag	cca	tgg	atg	cgg	1552	
Tyr	Asn	Gly	Phe	Ser	Pro	Pro	Asp	Ala	Lys	Asp	Lys	Pro	Trp	Met	Arg		
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Pro	Leu	Glu	Asp	Ala	Asp	Ile	Cys	Asn	Ala	Lys	Ala	Gln	Gln	Asn	Asp		
				485					490					495			
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Lys	Thr	Ser	Val	Leu	Gly	Phe	Trp	Lys	His	Met	Leu	Gln	Leu	Arg	Lys		
			500					505					510				
gag	cac	aag	gat	ttg	ctc	gtc	ttc	gg	cag	tac	gac	gat	ctt	gac	gtt	1696	
Glu	His	Lys	Asp	Leu	Leu	Val	Phe	Gly	Gln	Tyr	Asp	Asp	Leu	Asp	Val		
		515					520					525					
gac	aat	gag	caa	ttt	tac	att	ttc	agc	aag	acg	tgg	cag	gga	aag	cgg	1744	
Asp	Asn	Glu	Gln	Phe	Tyr	Ile	Phe	Ser	Lys	Thr	Trp	Gln	Gly	Lys	Arg		
	530					535					540						
gcg	ctg	tgt	att	tgc	aac	ttt	aca	gat	gag	agc	aag	cag	ctg	gtt	ttg	1792	
Ala	Leu	Cys	Ile	Cys	Asn	Phe	Thr	Asp	Glu	Ser	Lys	Gln	Leu	Val	Leu		
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cca	gag	agc	gtg	gca	gcg	cag	aaa	atg	cag	ctg	ctg	gta	agt	agt	cgg	1840	
Pro	Glu	Ser	Val	Ala	Ala	Gln	Lys	Met	Gln	Leu	Leu	Val	Ser	Ser	Arg		
				565					570					575			
caa	gac	gag	ggc	gtc	gag	gaa	aag	acg	cta	gcg	ccg	tac	gag	gga	agg	1888	
Gln	Asp	Glu	Gly	Val	Glu	Glu	Lys	Thr	Leu	Ala	Pro	Tyr	Glu	Gly	Arg		
			580					585					590				
gtc	tat	ctt	tat	gtg	caa	tga										1909	
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 <213> Cochliobolus heterostrophus ATCC 48331

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 Val Tyr Gln Ile Tyr Pro Ala Ser Phe Lys Asp Ser Asn Gly Asp Gly
 35 40 45
 Ile Gly Asp Leu Asn Gly Ile Ser Glu Leu Asp Tyr Ile Arg Ser
 50 55 60
 Ile Gly Val Asp Val Ile Trp Val Cys Pro Met Tyr Asp Ser Pro Gln
 65 70 75 80
 Val Asp Met Gly Tyr Asp Ile Arg Asn Tyr Glu Asp Val Tyr Arg Pro
 85 90 95
 Tyr Gly Thr Val Gln Asp Met Gln Arg Leu Ile Asp Glu Thr His Ser
 100 105 110

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Arg Gly Met Lys Ile Ile Leu Asp Leu Val Val Asn His Thr Ser Asp
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 Gln

<210> 556

<211> 180

<212> PRT

<213> Cochliobolus heterostrophus ATCC 48331

<400> 556

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 20 25 30
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 35 40 45
 Asp Glu His Thr Gln Glu Tyr Tyr Leu His Leu Phe Cys Pro Glu Gln
 50 55 60
 Pro Asp Leu Asn Trp Glu Asn Pro Glu Thr Arg Lys Ala Ile Tyr Glu
 65 70 75 80
 Ser Ala Met Glu Phe Trp Leu Lys Arg Gly Val Asp Gly Phe Arg Val
 85 90 95
 Asp Thr Val Asn Met Tyr Ser Lys Gly Asp Met Arg Asp Ala Pro Ile
 100 105 110
 Thr Asp Pro Gly Ser Glu Trp Gln Phe Ala Gly Tyr Gln Tyr Cys Asn
 115 120 125
 Gly Pro Arg Met Asp Glu Phe Leu Gly Glu Met Asn Gln Ile Leu Glu
 130 135 140
 Lys Tyr Asp Ala Met Thr Val Gly Glu Cys Pro His Thr Leu Asp Ile
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 Val Phe Gln Phe
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<212> PRT

<213> Cochliobolus heterostrophus ATCC 48331

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 35 40 45
 Asn His Asp Gln Ala Arg Ser Ile Thr Arg Phe Thr Ser Asp Ala Pro
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 Gln His Arg Val Ala Gly Gly Lys Met Leu Ala Leu Met Leu Ser Ala
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 Leu Ser Gly Thr Leu Phe Ile Tyr Gln Gly Gln Glu Ile Gly Met Thr
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 Asn Phe Pro Glu Ser Trp Asp Met Ser Glu Tyr Lys Asp Val Glu Ser
 100 105 110
 Ser Asn Tyr Tyr Lys Met Val Ala Lys Arg Thr Asn Asn Asp Val Asp
 115 120 125
 Ala Leu Ala Ala Ala His Lys Ser Leu Gln His Leu Ala Arg Asp His
 130 135 140
 Ser Arg Val Pro Met Ser Trp Ser Thr Ala Pro Tyr Asn Gly Phe Ser
 145 150 155 160
 Pro Pro Asp Ala Lys Asp Lys Pro Trp Met Arg Pro Leu Glu Asp Ala
 165 170 175
 Asp Ile Cys Asn Ala Lys Ala Gln Gln Asn Asp Lys Thr Ser Val Leu
 180 185 190
 Gly Phe Trp Lys His Met Leu Gln Leu Arg Lys Glu His Lys Asp Leu
 195 200 205

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Leu Val Phe Gly Gln Tyr Asp Asp Leu Asp Val Asp Asn Glu Gln Phe
 210 215 220
 Tyr Ile Phe Ser Lys Thr Trp Gln Gly Lys Arg Ala Leu Cys Ile Cys
 225 230 235 240
 Asn Phe Thr Asp Glu Ser Lys Gln Leu Val Leu Pro Glu Ser Val Ala
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 Ala Gln Lys Met Gln Leu Leu Val Ser Ser Arg Gln Asp Glu Gly Val
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 Glu Glu Lys Thr Leu Ala Pro Tyr Glu Gly Arg Val Tyr Leu Tyr Val
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 Gln

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 ctctgtctca accacacttc agatcagcac caatggttcc aagaatcacg ctctcaaaag 420
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 aagcgactct ccatggtgtt tcaattcgac gtagtcgatg ttgggcaagg cccctacaaa 960
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 aagatgcttg ccctcatgct gagcgctctt agcggcacgc tgttcatcta ccaaggccag 1200
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<210> 559
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 <212> PRT
 <213> *Cochliobolus heterostrophus* ATCC 48331

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 <222> (1)...(19)

<221> DOMAIN
 <222> (35)...(443)
 <223> Alpha amylase, catalytic domain

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Ile	Gly	Val	Asp	Val	Ile	Trp	Val	Cys	Pro	Met	Tyr	Asp	Ser	Pro	Gln
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Val	Asp	Met	Gly	Tyr	Asp	Ile	Arg	Asn	Tyr	Glu	Asp	Val	Tyr	Arg	Pro
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Tyr	Gly	Thr	Val	Gln	Asp	Met	Gln	Arg	Leu	Ile	Asp	Glu	Thr	His	Ser
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Arg	Gly	Met	Lys	Ile	Ile	Leu	Asp	Leu	Val	Val	Asn	His	Thr	Ser	Asp
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Gln	His	Gln	Trp	Phe	Gln	Glu	Ser	Arg	Ser	Ser	Lys	Asp	Asn	Pro	Lys
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Arg	Asp	Trp	Tyr	Ile	Trp	Arg	Pro	Ala	Arg	Tyr	Val	Asp	Gly	Val	Arg
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Lys	Pro	Pro	Asn	Asn	Trp	Val	Ser	Asn	Phe	Thr	Gly	Ser	Val	Trp	Glu
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Trp	Asp	Glu	His	Thr	Gln	Glu	Tyr	Tyr	Leu	His	Leu	Phe	Cys	Pro	Glu
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Val	Asp	Thr	Val	Asn	Met	Tyr	Ser	Lys	Gly	Asp	Met	Arg	Asp	Ala	Pro
	225				230					235					240
Ile	Thr	Asp	Pro	Gly	Ser	Glu	Trp	Gln	Phe	Ala	Gly	Tyr	Gln	Tyr	Cys
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Asn	Gly	Pro	Arg	Met	Asp	Glu	Phe	Leu	Gly	Glu	Met	Asn	Gln	Ile	Leu
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Glu	Lys	Tyr	Asp	Ala	Met	Thr	Val	Gly	Glu	Cys	Pro	His	Thr	Leu	Asp
		275					280					285			
Ile	Asn	Arg	Val	His	Gln	Tyr	Val	Ser	Ala	Lys	Glu	Lys	Arg	Leu	Ser
	290					295					300				
Met	Val	Phe	Gln	Phe	Asp	Val	Val	Asp	Val	Gly	Gln	Gly	Pro	Tyr	Lys
	305				310					315					320
Phe	Gln	Thr	Thr	Pro	Lys	Asn	Trp	Thr	Leu	Pro	Gln	Leu	Lys	Arg	Ala
				325					330					335	
Ile	Ala	Arg	Thr	Gln	Asp	Leu	Ile	Arg	Pro	Pro	Ser	Asp	Ser	Trp	Thr
			340					345					350		
Thr	Ala	Phe	Leu	Glu	Asn	His	Asp	Gln	Ala	Arg	Ser	Ile	Thr	Arg	Phe
		355					360					365			
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Leu	Met	Leu	Ser	Ala	Leu	Ser	Gly	Thr	Leu	Phe	Ile	Tyr	Gln	Gly	Gln
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Glu	Ile	Gly	Met	Thr	Asn	Phe	Pro	Glu	Ser	Trp	Asp	Met	Ser	Glu	Tyr
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Lys	Asp	Val	Glu	Ser	Ser	Asn	Tyr	Tyr	Lys	Met	Val	Ala	Lys	Arg	Thr
			420					425					430		
Asn	Asn	Asp	Val	Asp	Ala	Leu	Ala	Ala	Ala	His	Lys	Ser	Leu	Gln	His
		435					440					445			
Leu	Ala	Arg	Asp	His	Ser	Arg	Val	Pro	Met	Ser	Trp	Ser	Thr	Ala	Pro
	450					455					460				
Tyr	Asn	Gly	Phe	Ser	Pro	Pro	Asp	Ala	Lys	Asp	Lys	Pro	Trp	Met	Arg
	465				470					475					480
Pro	Leu	Glu	Asp	Ala	Asp	Ile	Cys	Asn	Ala	Lys	Ala	Gln	Gln	Asn	Asp
				485					490					495	
Lys	Thr	Ser	Val	Leu	Gly	Phe	Trp	Lys	His	Met	Leu	Gln	Leu	Arg	Lys
			500					505					510		
Glu	His	Lys	Asp	Leu	Leu	Val	Phe	Gly	Gln	Tyr	Asp	Asp	Leu	Asp	Val
		515					520					525			
Asp	Asn	Glu	Gln	Phe	Tyr	Ile	Phe	Ser	Lys	Thr	Trp	Gln	Gly	Lys	Arg
	530					535					540				
Ala	Leu	Cys	Ile	Cys	Asn	Phe	Thr	Asp	Glu	Ser	Lys	Gln	Leu	Val	Leu
	545				550					555					560
Pro	Glu	Ser	Val	Ala	Gln	Lys	Met	Gln	Leu	Leu	Val	Ser	Ser	Ser	Arg
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Ser Met Leu Trp Val Val Leu Leu Thr Ser Phe Val Lys Asp Val His
20 25 30

gct gcc act tca gac gat tgg aga agc caa tct ata tat caa att atc 144
Ala Ala Thr Ser Asp Asp Trp Arg Ser Gln Ser Ile Tyr Gln Ile Ile
35 40 45

aca gat cga ttc gcc aga acg gat ggc tcg act acc tat gcg tgt aac 192
Thr Asp Arg Phe Ala Arg Thr Asp Gly Ser Thr Thr Tyr Ala Cys Asn
50 55 60

aac aca gca cag ctg tat tgc ggt gga act ttt caa gga att atc aac 240
Asn Thr Ala Gln Leu Tyr Cys Gly Gly Thr Phe Gln Gly Ile Ile Asn
65 70 75 80

cag cta gat tac atc cag gat atg gga ttt acg gct gtgagtttgt 286
Gln Leu Asp Tyr Ile Gln Asp Met Gly Phe Thr Ala
85 90

ctatctcagc ttctccttac gctcttcgca cgaacgatcc ttctctctgt tgtgtctcgt 346
acgaccgtcg cgacgaaaga tgggaaaatc caccagccta gacatcgtga actgacttaa 406
tgccaaactt agatctggat ttcaccggtg gtacag aat atc aac caa aca acg 460
Asn Ile Asn Gln Thr Thr
95

gca tat ggg caa ggc tat cat gga ttt tgg agt caa gat atc act aag 508
Ala Tyr Gly Gln Gly Tyr His Gly Phe Trp Ser Gln Asp Ile Thr Lys
100 105 110

atc aac gaa cat ttc gga aca gca gac gat ctc aaa ctc ttg tcg tca 556
Ile Asn Glu His Phe Gly Thr Ala Asp Asp Leu Lys Leu Leu Ser Ser
115 120 125 130

acc ttg cac gat cgt ggc atg gtaagctgca ttcacattcg caacccaatc 607
Thr Leu His Asp Arg Gly Met

2013201807

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135

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 Tyr Leu Met Ile Asp
 140

gcg gtc att aac gac atg gca tac gca ggg aaa ggg agc gca gtc gat 710
Ala Val Ile Asn Asp Met Ala Tyr Ala Gly Lys Gly Ser Ala Val Asp
145 150 155

tac	tcc	act	cta	gtt	ccg	ttc	aac	aaa	aaa	gaa	tac	ttc	cat	cct	ttc	758
Tyr	Ser	Thr	Leu	Val	Pro	Phe	Asn	Lys	Lys	Glu	Tyr	Phe	His	Pro	Phe	
	160					165					170					

tgt	tac	att	acc	gat	tat	tcc	aac	gca	aca	aac	ttt	caa	aac	tgt	tgg	806
Cys	Tyr	Ile	Thr	Asp	Tyr	Ser	Asn	Ala	Thr	Asn	Phe	Gln	Asn	Cys	Trp	
175					180					185					190	

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Leu Gly Asp Asp Thr Val Ser Leu Pro Asp Leu Asp Thr Glu Ser Asp
195 200 205

ttt gtc aag caa aca tgg gaa acc tgg gta aca gaa atg gtg gcc aat 902
Phe Val Lys Gln Thr Trp Glu Thr Trp Val Thr Glu Met Val Ala Asn
210 215 220

tat tct c gtatgtactt ctgatcatga atctcactca ccccttgctc tttctggaag 959
Tyr Ser

tcttggctgt	ggggaaaatc	tggggaatat	taatgcacaa	tacattccat	tgcaacaaga	1019
gtgggctaac	ataatctcga	acttttag tt	gat ggt ctt	cgt atc gat	gcc gcc	1072
		Leu	Asp Gly Leu Arg	Ile Asp Ala Ala		
		225		230		

aaa cac gtc gac aag cct ttc tgg cct ggc ttc caa cag gcc gct aat 1120
Lys His Val Asp Lys Pro Phe Trp Pro Gly Phe Gln Gln Ala Ala Asn
235 240 245

aca ttc act att ggt gaa gtt ttt gac gag ctg cct acc aat gca tgc 1168
Thr Phe Thr Ile Gly Glu Val Phe Asp Glu Leu Pro Thr Asn Ala Cys
250 255 260 265

gaa tgg gcc gtc gat gct ctc tct tct gtc cta aat tat cct gcc tgg 1216
Glu Trp Ala Val Asp Ala Leu Ser Ser Val Leu Asn Tyr Pro Ala Trp
270 275 280

tat	tac	att	acg	tct	atc	ctc	tca	aac	tca	acg	aat	ggg	atg	gga	agt	1264
Tyr	Tyr	Ile	Thr	Ser	Ile	Leu	Ser	Asn	Ser	Thr	Asn	Gly	Met	Gly	Ser	
			285					290					295			

tta	agc	tac	caa	ttc	gat	caa	acc	cag	caa	tac	tgt	tac	gat	act	aca	1312
Leu	Ser	Tyr	Gln	Phe	Asp	Gln	Thr	Gln	Gln	Tyr	Cys	Tyr	Asp	Thr	Thr	
		300					305					310				

ctt ctt ggt act ttc agt gag aat cat gat gtt gct cgt ttt gga tcc 1360
Leu Leu Gly Thr Phe Ser Glu Asn His Asp Val Ala Arg Phe Gly Ser
315 320 325

tac	acg	agt	gac	gtg	tct	caa	agg	aaa	aat	gcc	ctc	acg	ttc	gac	ttt	1408
Tyr	Thr	Ser	Asp	Val	Ser	Gln	Arg	Lys	Asn	Ala	Leu	Thr	Phe	Asp	Phe	
330					335					340					345	

ttt aca gac ggc att cct att gta tat tac gga gcc gag caa ggc ctt 1456
Phe Thr Asp Gly Ile Pro Ile Val Tyr Tyr Gly Ala Glu Gln Gly Leu
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2013201807

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<213>	Fungi

<210>	562
<211>	45
<212>	PRT
<213>	Fungi

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 35 40 45

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 <212> PRT
 <213> Fungi

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 35 40 45
 Phe Gln Asn Cys Trp Leu Gly Asp Asp Thr Val Ser Leu Pro Asp Leu
 50 55 60
 Asp Thr Glu Ser Asp Phe Val Lys Gln Thr Trp Glu Thr Trp Val Thr
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 Glu Met Val Ala Asn Tyr Ser
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 <211> 283
 <212> PRT
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 35 40 45
 Ser Ser Val Leu Asn Tyr Pro Ala Trp Tyr Tyr Ile Thr Ser Ile Leu
 50 55 60
 Ser Asn Ser Thr Asn Gly Met Gly Ser Leu Ser Tyr Gln Phe Asp Gln
 65 70 75 80
 Thr Gln Gln Tyr Cys Tyr Asp Thr Thr Leu Leu Gly Thr Phe Ser Glu
 85 90 95
 Asn His Asp Val Ala Arg Phe Gly Ser Tyr Thr Ser Asp Val Ser Gln
 100 105 110
 Arg Lys Asn Ala Leu Thr Phe Asp Phe Phe Thr Asp Gly Ile Pro Ile
 115 120 125
 Val Tyr Tyr Gly Ala Glu Gln Gly Leu Asp Gly Glu Asn Asp Pro Gln
 130 135 140
 Asn Arg Gly Ala Leu Trp Leu Asn Pro Thr Gly Tyr Asp Thr Thr Ala
 145 150 155 160
 Val Leu Tyr Gln His Ile Lys Thr Met Asn Thr Ala Arg Asn Ala Val
 165 170 175
 Asn Asn Tyr Met Ile Ala Thr Asn Tyr Ser Asn Trp Ser Pro Tyr Trp
 180 185 190
 Ala Tyr Lys Ala Ser Val Ile Lys Gln Ala Asp Asp Val Leu Val Phe
 195 200 205
 Arg Lys Gly Leu Val His Ser Ile Val Thr Ala Ile Thr Asn Val Gly
 210 215 220
 Thr Glu Gly Ala Thr Val Gly Pro Tyr Tyr Ile Glu Asp Thr Asn Phe
 225 230 235 240
 Ser Glu Gly Asn Leu Ile Glu Ile Leu Ser Cys Asn Ser Thr Val
 245 250 255
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 cagctagatt acatccagga tatgggattt acggctaata tcaaccaaac aacggcata 300
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 acagcagacg atctcaaact cttgtcgtca accttgcacg atcgtggcat gtatctaata 420
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 aacaattaca tgatagcaac caattactct aattggagtc cttactgggc atacaaagca 1260
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 gacaccaatt tcagcgaagg aaatctcttc attgagatct tgagctgtaa cagcactgtc 1440
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 Thr Asp Arg Phe Ala Arg Thr Asp Gly Ser Thr Thr Tyr Ala Cys Asn
 50 55 60
 Asn Thr Ala Gln Leu Tyr Cys Gly Gly Thr Phe Gln Gly Ile Ile Asn
 65 70 75 80
 Gln Leu Asp Tyr Ile Gln Asp Met Gly Phe Thr Ala Asn Ile Asn Gln
 85 90 95
 Thr Thr Ala Tyr Gly Gln Gly Tyr His Gly Phe Trp Ser Gln Asp Ile
 100 105 110
 Thr Lys Ile Asn Glu His Phe Gly Thr Ala Asp Asp Leu Lys Leu Leu
 115 120 125
 Ser Ser Thr Leu His Asp Arg Gly Met Tyr Leu Met Ile Asp Ala Val
 130 135 140
 Ile Asn Asp Met Ala Tyr Ala Gly Lys Gly Ser Ala Val Asp Tyr Ser
 145 150 155 160
 Thr Leu Val Pro Phe Asn Lys Lys Glu Tyr Phe His Pro Phe Cys Tyr
 165 170 175
 Ile Thr Asp Tyr Ser Asn Ala Thr Asn Phe Gln Asn Cys Trp Leu Gly

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Asp	Asp	Thr	180	Val	Ser	Leu	Pro	Asp	185	Leu	Asp	Thr	Glu	Ser	190	Asp	Phe	Val
Lys	Gln	Thr	195	Trp	Glu	Thr	Trp	Val	200	Thr	Glu	Met	Val	Ala	205	Asn	Tyr	Ser
Leu	Asp	Gly	210	Leu	Arg	Ile	Asp	Ala	215	Ala	Lys	His	Val	Asp	Lys	Pro	Phe	
225	Trp	Pro	Gly	Phe	Gln	230	Ala	Ala	235	Asn	Thr	Phe	Thr	Ile	Gly	Glu	Val	
			245	Pro	Thr	250	Asn	Ala	255	Cys	Glu	Trp	Ala	Val	Asp	Ala	Leu	
			260	Ser	Ser	Val	265	Leu	270	Trp	Tyr	Tyr	Ile	Thr	Ser	Ile	Leu	
			275	Ser	Asn	Thr	280	Asn	285	Gly	Ser	Leu	Ser	Tyr	Gln	Phe	Asp	
			290	Thr	Gln	Gln	295	Tyr	300	Thr	Thr	Leu	Leu	Gly	Thr	Phe	Ser	
			305	Asn	His	Asp	310	Val	315	Ala	Arg	Phe	Gly	Ser	Asp	Val	Ser	
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			340	Val	Tyr	Tyr	345	Gly	350	Leu	Asp	Gly	Glu	Asn	Asp	Pro	Gln	
			355	Asn	Arg	Gly	360	Ala	365	Leu	Trp	Leu	Trp	Asp	Thr	Thr	Ala	
			370	Val	Leu	Tyr	375	Gln	380	His	Ile	Lys	Thr	Met	Asn	Thr	Ala	
			385	Asn	Asn	Tyr	390	Met	395	Ile	Ala	Thr	Asn	Tyr	Ser	Asn	Trp	
			405	Ala	Tyr	Lys	410	Ala	415	Val	Thr	Ala	Ile	Thr	Asn	Val	Gly	
			420	Arg	Lys	Gly	425	Leu	430	Val	Thr	Ala	Ile	Thr	Asn	Val	Gly	
			435	Thr	Glu	Gly	440	Ala	445	Thr	Tyr	Tyr	Ile	Glu	Asp	Thr	Asn	
			450	Ser	Glu	Gly	455	Asn	460	Leu	Ile	Leu	Ser	Cys	Asn	Ser	Thr	
			465	Ala	Gly	Val	470	Gly	475	Ala	Phe	Thr	Leu	Thr	Ser	Gly	Glu	
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 <223> Alpha amylase, catalytic domain

<221> DOMAIN
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Lys Ala Glu Ala Ala Phe Val His Leu Phe Glu Trp Arg Trp Leu Asp
      50      55      60
Ile Ala Ser Glu Cys Glu Asn Phe Leu Gly Pro Lys Gly Phe Ser Ala
      65      70      75      80
Val Gln Ile Ser Pro Gln Glu His Leu Asp His His Thr Trp Trp
      85      90      95
Ala Arg Tyr Gln Pro Val Ser Leu Thr Lys Leu Thr Ser Arg Ser Gly
      100      105      110
Thr Glu Thr Glu Leu Arg Glu Met Val Ala Arg Cys Ala Ala Ala Gly
      115      120      125
Val Lys Ile Tyr Ala Asp Val Ile Asn Asn Trp Ala Thr Leu Pro
      130      135      140
Gly Ala Ser Arg Ile Gly Ser Ala Gly Ser Gln Trp Glu Ala Tyr Gln
      145      150      155      160
Tyr Pro Asp Leu Gly Pro Asp Asp Phe His Ser Pro Arg Cys Gly Ile
      165      170      175
Glu Thr Tyr Gln Asn Ala His Gln Val Trp Tyr Cys Glu Leu Tyr Gly
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Met Pro Asp Leu Tyr Thr Gly Ala Pro Gly Pro Gln Ala Tyr Val Ala
      195      200      205
Asp Tyr Ile Lys Arg Leu Thr Ala Met Gly Ile Ala Gly Phe Arg Met
      210      215      220
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      225      230      235      240
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      260      265      270
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      275      280      285
Val Asp Leu Leu Arg Asp Leu Pro Lys Ser Asp Trp Leu Pro Ser Glu
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Gln Ala Val Ile Phe Val Asp Asn His Asp Arg Glu Arg Gly His Gly
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355 360
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370 380
Ala Asn Met Leu Gly Phe Arg Ala His Thr Gln Asn Glu Pro Leu Leu
385 390
His Trp Trp Asp Asn Gly Arg Asp Gln Leu Ala Phe Ala Arg Gly Lys
400 415
Arg Gly Phe Val Ala Ile Asn Asn Ser Ser Lys Arg Leu Asn Arg Arg
420 430
Met Gln Thr Gly Leu Pro Ala Gly Arg Tyr Cys Asn Gln Leu Ala Glu
435 445
Lys Ser Ala Cys Asn Ala Pro Ile Asn Val Asp Lys Lys Gly Arg Ala
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<211> 1095

<212> DNA

<213> Thermococcus alcaliphilus AEDII12RA

<400> 569

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<210> 570

<211> 364

<212> PRT

<213> Thermococcus alcaliphilus AEDII12RA

<400> 570

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Glu Thr Leu Ile Lys Glu Glu Ile Pro Phe Gly Leu Asn Ile Thr Gly
35 40 45
Tyr Thr Leu Lys Phe Leu Pro Lys Asp Ile Ile Asp Leu Val Lys Gly
50 55 60
Gly Ile Ala Ser Asp Leu Ile Glu Ile Ile Gly Thr Ser Tyr Thr His
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Ala Ile Leu Pro Leu Leu Pro Leu Ser Arg Val Glu Ala Gln Val Gln
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Arg Asp Arg Glu Val Lys Glu Glu Leu Phe Glu Val Ser Pro Lys Gly
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Phe Trp Leu Pro Glu Leu Ala Tyr Asp Pro Ile Ile Pro Ala Ile Leu

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 Pro His Leu Ile Lys Ala Gln Arg Glu Lys Arg Phe Arg Tyr Ile Ser
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 Tyr Leu Leu Gly Leu Arg Glu Leu Arg Lys Ala Ile Lys Leu Val Phe
 180 185 190
 Glu Gly Lys Val Thr Leu Lys Ala Val Lys Asp Ile Glu Ala Val Pro
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 Val Trp Val Ala Val Asn Thr Ala Val Met Leu Gly Ile Gly Arg Leu
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 Pro Leu Met Asn Pro Lys Lys Val Ala Ser Trp Ile Glu Asp Lys Asp
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 305 310 315 320
 Tyr Asn Met Arg Gly Glu Leu Ala Phe Leu Ala Glu Asn Ser Asp Ala
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<211> 1839

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 571

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<210> 572
 <211> 612
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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 <222> (1)...(28)

<221> DOMAIN
 <222> (517)...(610)
 <223> Starch binding domain

<221> DOMAIN
 <222> (32)...(424)
 <223> Alpha amylase, catalytic domain

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 35 40 45
 Trp Leu Lys Val Thr Asp Gln Ala Asp Glu Leu Ser Ala Ala Gly Ile
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 Thr Ala Leu Trp Leu Pro Pro Ala Tyr Lys Ala Ile Asn Gly Asp Asp
 65 70 75 80
 Val Gly Tyr Gly Val Tyr Asp Leu Tyr Asp Leu Gly Glu Phe Asn Gln
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 Ala Ile Asn Ala Ala His Ala Asn Gly Leu Gln Val Tyr Gly Asp Val
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 Thr Arg Val Gln Thr Trp Asp Arg Asn Gln Glu Tyr Gly Gly Asp Leu
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 Ser Ile Gln Ala Trp Thr Gly Phe Asp Phe Pro Gly Arg Gly Asn Thr
 165 170 175
 Tyr Ser Ala Phe Lys Trp Arg Trp Tyr His Phe Asp Gly Val Asp Trp
 180 185 190
 Asp Gln Asn Leu Gln Glu Ser Gly Lys Ile Tyr Lys Phe Arg Gly Thr
 195 200 205
 Gly Lys Ala Trp Asp Trp Lys Val His Asp Glu Asn Gly Asn Tyr Asp
 210 215 220
 Tyr Leu Met Phe Ala Asp Leu Asp Met Gln His Pro Asp Val Val Asn
 225 230 235 240
 Glu Leu Lys Ser Trp Gly Val Trp Tyr Ala Asn Thr Thr Gly Val Asp
 245 250 255
 Gly Phe Arg Leu Asp Ala Leu Lys His Ile Lys Tyr Asp Phe Trp Asn
 260 265 270
 Gly Trp Leu Asp His Val Arg Asn Thr Thr Gly Lys Pro Leu Phe Thr
 275 280 285
 Val Gly Glu Leu Trp Ser Tyr Asp Val Gln Asn Leu His Asp Trp Ile
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 Thr Glu Thr Gly Gly Arg Ala Ser Leu Phe Asp Ala Pro Leu His Leu
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 Asn Phe Phe Asn Ala Ser Arg Ala Gly Gly Ser Tyr Asp Met Arg Asp
 325 330 335
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Tyr	Asp	Tyr	420	Leu	Asp	His	Trp	Asp	425	Val	Ile	Gly	Trp	Thr	430	Arg	Leu	Gly
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Pro	Thr	Trp	545	Ser	Gly	Trp	Val	His	550	Asn	Leu	Pro	Pro	Ser	555	Thr	Gln	Ile
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			610															

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 <211> 2133
 <212> DNA
 <213> Bacteria

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<210> 574
 <211> 710
 <212> PRT
 <213> Bacteria

<220>
 <221> SIGNAL
 <222> (1)...(27)

<221> DOMAIN
 <222> (437)...(521)
 <223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN
 <222> (46)...(425)
 <223> Alpha amylase, catalytic domain

<221> DOMAIN
 <222> (526)...(603)
 <223> IPT/TIG domain

<221> DOMAIN
 <222> (610)...(706)
 <223> Starch binding domain

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 Thr Asp Arg Phe Val Asp Gly Asn Thr Ser Asn Asn Pro Thr Gly Asp
 50 55 60
 Leu Tyr Asp Pro Thr His Thr Ser Leu Lys Lys Tyr Phe Gly Gly Asp
 65 70 75 80
 Trp Gln Gly Ile Ile Asn Lys Ile Asn Asp Gly Tyr Leu Thr Gly Met
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 Gly Val Thr Ala Ile Trp Ile Ser Gln Pro Val Glu Asn Ile Tyr Ala
 100 105 110
 Val Leu Pro Asp Ser Thr Phe Gly Gly Ser Thr Ser Tyr His Gly Tyr
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 Asp Phe Gln Asn Leu Ile Asn Thr Ala His Ala His Asn Ile Lys Val
 145 150 155 160
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 165 170 175
 Pro Thr Tyr Ala Glu Asn Gly Arg Leu Tyr Asp Asn Gly Thr Leu Leu
 180 185 190
 Gly Gly Tyr Thr Asn Asp Thr Asn Gly Tyr Phe His His Tyr Gly Gly
 195 200 205
 Thr Asp Phe Ser Ser Tyr Glu Asp Gly Ile Tyr Arg Asn Leu Phe Asp
 210 215 220
 Leu Ala Asp Leu Asn Gln Gln Asn Ser Thr Ile Asp Ser Tyr Leu Lys
 225 230 235 240
 Ser Ala Ile Lys Val Trp Leu Asp Met Gly Ile Asp Gly Ile Arg Leu

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Gly	Met	Ser	Leu	290	Leu	Asp	Phe	Arg	Phe	295	Ser	Gln	Lys	Val	Arg	Gln	Val	
Phe	Arg	Asp	Asn	305	Thr	Asp	Thr	Met	Tyr	310	Gly	Leu	Asp	Ser	Met	Ile	Gln	
Ser	Thr	Ala	Ser	325	Asp	Tyr	Asn	Phe	Ile	330	Asn	Asp	Met	Val	Thr	Phe	Ile	
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Gln	Phe	Lys	Phe	660	Ile	Lys	Lys	Asn	Gly	665	Asn	Thr	Ile	Thr	Trp	Glu	Gly	
Gly	Ser	Asn	His	675	Thr	Tyr	Thr	Val	Pro	680	Ser	Ser	Ser	Thr	Gly	Thr	Val	
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<210> 575
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<212> DNA
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<220>
<223> obtained from an environmental sample.

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gtgatgctga tatcgggtcg aaactggggg gtgcagcagt cgttcaccat caacagccat   1260
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ggcaactttg aatggaaaac ccgtaaaggc ccaaccggcg gcagtgggtc ggactgggaa   1620
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<210> 576

<211> 558

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<221> DOMAIN

<222> (15)...(366)

<223> Alpha amylase, catalytic domain

<400> 576

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          20          25          30
Leu Pro Ala Met Ala Ala Pro Leu Ala Glu Arg Gly Ile Thr Ser Met
 35          40          45
Trp Leu Pro Pro Ala Ala Lys Gly Met Asn Gly Thr Phe Ser Val Gly
 50          55          60
Tyr Asp Val Tyr Asp Phe Trp Asp Leu Gly Glu Phe Asn Gln Lys Gly
 65          70          75          80
Thr Thr Ala Thr Arg Tyr Gly Thr Arg Gln Gln Leu Gln Gln Ala Leu
          85          90          95
Ser Ala Leu Asp Gln Leu Gly Ile Gln Ala Tyr Phe Asp Val Val Phe
          100          105          110
Asn His Arg Met Gly Ala Asp Ala Gln Glu Asn Ile Pro Gly Phe Gly
          115          120          125
Leu Ala Trp Thr Glu Tyr His Leu Gln Gly Arg Gln Ala His Tyr Thr
          130          135          140
Gln Gln Asn Trp Gly Tyr Leu Trp His Asp Phe Asp Trp Asn Trp Thr
          145          150          155          160
Ala Phe Asn Gly Ser Asp Asn Gln Leu Tyr Pro Gly Lys Trp Trp Gly
          165          170          175
Asn Thr Phe His Phe Pro Tyr Leu Met Gly Glu Asp Val Asp Tyr Asn
          180          185          190
Arg Phe Glu Val Gln Gln Glu Met Lys Ala Trp Gly Glu Trp Ile Ile

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195 200 205
 Asn Ser Val Gly Phe Ser Gly Phe Arg Met Asp Ala Ile Ala His Val
 210 215 220
 Asp Thr Asp Phe Thr Arg Asp Trp Ile Asn His Val Gln Trp Ala Thr
 225 230 235 240
 Ser Glu Asp Val Phe Phe Val Ala Glu Ala Trp Val Ser Asp Ile Asn
 245 250 255
 Gly Tyr Leu Asp Ala Val Asn Thr Pro His Leu Arg Ala Phe Asp Phe
 260 265 270
 Asn Leu Arg Glu Asp Phe Val Ala Leu Ser Ser Gly Gly Lys Asp Met
 275 280 285
 Arg Trp Trp Gly Gly Leu Val Asn Ser Gln His Arg Asp Arg Ala Val
 290 295 300
 Thr Phe Val Asp Asn His Asp Thr Ser Arg Ala Gly Asn Pro Tyr Gly
 305 310 315 320
 Met Pro Gln Val Ile Asn Tyr Lys Asn Gln Ala Tyr Ala Tyr Ile Leu
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 Leu Arg Glu His Gly Val Pro Thr Val Phe Ala Arg Asp Tyr Asp Glu
 340 345 350
 Phe Gly Met Ala Pro Thr Leu Asp Lys Leu Ile Glu Ala Arg Arg Tyr
 355 360 365
 Phe Ala Tyr Gly Pro Gly His Glu Tyr Ser Gly Asn Thr Glu Ala Val
 370 375 380
 Tyr Ala Tyr Val Arg Glu Gly Leu Ser Thr Val Pro Gly Thr Gly Leu
 385 390 395 400
 Val Met Leu Ile Ser Gly Arg Asn Trp Gly Gly Gln Gln Ser Phe Thr
 405 410 415
 Ile Asn Ser His Gln Pro Asn Thr Thr Phe Tyr Asp Tyr Thr Gly Asn
 420 425 430
 Val Ser Gly Thr Val Thr Thr Asn Ala Gln Gly Tyr Gly Ser Phe Pro
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 Val Thr Met Thr Glu Ser Thr Gly Trp Ser Val Trp Val Pro Gln Ser
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 Ser Gly Gly Thr Gln Pro Gly Ser Ile Thr Leu Arg Met Thr Lys Asp
 465 470 475 480
 Val Gly Tyr Gly Phe Ser Leu Phe Phe Thr Gly Ser Ser Ala Glu Leu
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 Thr Asn Trp Gly Gly Ile Glu Gly Thr Trp Thr Ser Gly His Val
 500 505 510
 Trp Glu Val Thr Ile Pro Asp Pro Gly Asn Phe Glu Trp Lys Thr Arg
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 Lys Gly Pro Thr Gly Gly Ser Gly Gln Asp Trp Glu Ser Gly Ser Asn
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 His Asn Gln Thr Asn Leu His Pro Ser Phe Asn Gly Gly Phe
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<210> 577

<211> 2160

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<400> 577

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gccggaatgg	gtgtaagcgc	tttatggatc	tctcaaccgg	tcgaaaatat	ttacagcgctc	360
atcgactact	ccgggtgtcaa	cagcacatcc	tatcacgggt	attgggctcg	cgacttcaaa	420
cggacgaatc	cggcgtttgg	cagcatgagc	gattttcagg	agctcatcaa	cacggcacat	480
gccacaata	tcaagatcat	tattgatttc	gcacctaac	acacctcccc	tgcttcggag	540
acacaacctt	ccttcgccga	gaacggtcgg	ctgtacgata	atggcaacct	gatcgctggg	600
tacacggggg	acaccaacgg	aattttccat	cataatcagg	gcacgaattt	ctcctcgctc	660
gaggatggca	tctatcggaa	tctttatgat	ctggccgaca	tcaatcacca	caacaacgta	720
acggatactt	acttcaagga	cgcgattaag	ctctggctga	atatggggat	cgatggcatc	780

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<210> 578

<211> 719

<212> PRT

<213> Unknown

<220>

<223> Obtained from an environmental sample.

<221> SIGNAL

<222> (1)...(34)

<221> DOMAIN

<222> (443)...(527)

<223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN

<222> (53)...(431)

<223> Alpha amylase, catalytic domain

<221> DOMAIN

<222> (532)...(615)

<223> IPT/TIG domain

<221> DOMAIN

<222> (619)...(715)

<223> Starch binding domain

<400> 578

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Ser Ala Ala Pro Asp Thr Ser Val Leu Asn Lys Gln Gly Phe Ser Thr
35     40     45
Asp Val Ile Tyr Gln Ile Val Thr Asp Arg Phe Ala Asp Gly Asp Pro
50     55     60
Ser Asn Asn Pro Ser Gly Ala Ala Tyr Ser Pro Gly Cys Thr Asn Leu
65     70     75     80
Lys Leu Tyr Cys Gly Gly Asp Trp Arg Gly Ile Ile Asn Lys Ile Asn
85     90     95
Glu Gly Tyr Phe Ala Gly Met Gly Val Ser Ala Leu Trp Ile Ser Gln
100    105    110
Pro Val Glu Asn Ile Tyr Ser Val Ile Asp Tyr Ser Gly Val Asn Ser
115    120    125

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Ala	His	Asn	Ile	Lys	Ile	Ile	Ile	Asp	Phe	170	Ala	Pro	Asn	His	Thr
Pro	Ala	Ser	Glu	165	Thr	Gln	Pro	Ser	Phe	185	Glu	Asn	Gly	Arg	Leu
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Phe	His	His	Asn	Gln	Gly	Thr	Asn	Phe	Ser	Ser	Leu	205	Glu	Asp	Gly
210						215					220				Ile
Tyr	Arg	Asn	Leu	Tyr	Asp	Leu	Ala	Asp	Ile	Asn	His	His	Asn	Asn	Val
225					230					235					240
Thr	Asp	Thr	Tyr	Phe	Lys	Asp	Ala	Ile	Lys	Leu	Trp	Leu	Asn	Met	Gly
Ile	Asp	Gly	Ile	Arg	Val	Asp	Ala	Val	Lys	His	Met	Pro	Glu	Gly	Trp
Gln	Lys	Asn	Trp	Val	Ala	Ser	Ile	Ala	Gly	Tyr	Lys	Pro	Val	Phe	Thr
Phe	Gly	Glu	Trp	Phe	Leu	Gly	Val	Gly	Glu	Asn	Asp	Pro	Asn	Asn	Ile
Arg	Phe	Ala	Asn	Glu	Ser	Gly	Met	Ser	Leu	Leu	Asp	Phe	Gln	Phe	Gly
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Gln	Lys	Val	Arg	Gln	Val	Phe	Arg	Asp	Asn	Ser	Asp	Asn	Met	Tyr	Gly
Leu	Asn	Asn	Met	Leu	Thr	Thr	Thr	Ala	Ala	Asn	Tyr	Lys	His	Ile	His
Asp	Gln	Val	Thr	Phe	Ile	Asp	Asn	His	Asp	Met	Asp	Arg	Phe	Lys	Leu
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Ser	Arg	Gly	Val	Pro	Ala	Ile	Tyr	Tyr	Gly	Thr	Glu	Gln	Tyr	Met	Thr
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Asn	Asp	Val	Phe	Ile	Phe	Glu	Arg	Lys	Phe	Gly	Ser	Ser	Val	Ala	Leu
Val	Ala	Ile	Asn	Arg	Ser	Thr	Thr	Asn	Ala	Thr	Ser	Ile	Thr	Gly	Leu
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Thr	Val	Phe	Phe	Gly	Thr	Thr	Ala	Val	Thr	Gly	Ser	Gln	Ile	Leu	Gln
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Gln	Tyr	Asn	Ile	Lys	Val	Arg	Ser	Ala	Ser	Ser	Val	Asp	Ser	Asn	Ser
Tyr	Ser	Ser	Phe	Asn	Leu	Leu	Thr	Gly	Asp	Gln	Val	Ser	Val	Arg	Phe
Ile	Ile	Asn	Asn	Ala	Asn	Thr	Gln	Leu	Gly	Glu	Asn	Val	Tyr	Leu	Thr
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Gly	Asn	Ile	Ala	Glu	Leu	Gly	Asn	Trp	Asp	Pro	Asn	Lys	Ala	Met	Gly
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			660					665					670		

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 <211> 5007
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample.

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 <212> PRT
 <213> Unknown

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 <222> (1)...(29)

<221> DOMAIN
 <222> (390)...(821)
 <223> Alpha amylase, catalytic domain

<221> DOMAIN
 <222> (833)...(924)
 <223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN
 <222> (929)...(1009)
 <223> Fibronectin type III domain

<221> DOMAIN
 <222> (1165)...(1248)
 <223> Fibronectin type III domain

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 Thr Asp Thr Ala Pro Ala Ile Ala Asn Val Val Gly Asn Phe Gln Ser

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Thr	Tyr	50	Gly	Asn	Gly	Phe	55	Glu	Phe	Thr	Thr	60	Pro	Val	Ala	Leu
Pro	Ala	65	Gly	Asp	Tyr	Glu	70	Tyr	Val	Ala	Leu	75	Asn	His	Ser	Trp
Gly	Gly	85	Val	Pro	Ser	Gln	90	Gly	Asn	Leu	Ser	105	Phe	His	Leu	95
Asp	Ser	100	Val	Thr	Phe	Tyr	110	Tyr	Asn	Tyr	Asn	125	Thr	Ser	Ser	Ile
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Val	Gly	135	Thr	Ile	Gln	Pro	140	Ile	Gly	Ala	Gly	145	Asp	Asp	Trp	Lys
Glu	Thr	150	Ser	Thr	Ala	Ile	155	Met	Arg	Tyr	Lys	160	Phe	Asn	Asn	Val
Glu	Tyr	165	Ala	Asn	Val	Pro	170	Lys	Gly	Asn	Tyr	175	Glu	Phe	Lys	Val
Leu	Gly	180	Ser	Trp	Asp	Ile	185	Tyr	Gly	Leu	Asn	190	Gly	Glu	Gln	Asn
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Tyr	Tyr	210	Asp	Ser	Val	Ser	215	His	Asn	Ile	Trp	220	Thr	Asp	Tyr	Asn
Leu	Thr	225	Gly	Pro	Asp	Asn	230	Ile	Tyr	Tyr	Asp	235	Asp	Leu	Arg	His
Thr	His	245	Pro	Phe	Phe	Arg	250	Ser	Pro	Phe	Gly	255	Ala	Ile	Lys	Thr
Asp	Thr	260	Thr	Leu	Arg	Ile	265	Gln	Ala	Lys	Asn	270	His	Asp	Ile	Glu
Ala	Lys	275	Ile	Ser	Trp	Asp	280	Ile	Lys	Lys	Thr	285	Arg	Ile	Glu	Val
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Gln	Leu	340	Gly	Val	Gly	Lys	345	Thr	Asp	Thr	Glu	350	Asn	Lys	Asp	Phe
Glu	Leu	355	Val	Tyr	Asp	Lys	360	Asn	Leu	Asp	Thr	365	Pro	Asp	Trp	Met
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Ser	Ile	515	Tyr	Phe	Asp	Arg	520	Gly	Lys	Tyr	Leu	525	Asn	Thr	Gly	Val
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Gly	Asp	545	Trp	Tyr	Glu	Ile	550	Lys	Pro	Asp	Gly	555	Thr	Tyr	Glu	Gly
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Gly	Trp	Arg	610	Leu	Asp	Val	615	Ala	Asn	Glu	Val	Ala	His	620	Asp	Phe	Trp
His	Phe	Arg	625	Gly	Ala	Ile	630	Asn	Thr	Val	Lys	635	Pro	640	Asn	Ala	Pro
Ala	Glu	Asn	645	Trp	Asn	Asp	650	Ala	Ser	Leu	Asp	Leu	Leu	655	Gly	Asp	Ser
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Ala	Ala	Lys	690	Leu	Asp	Gln	695	Arg	Leu	Met	Ser	Ile	700	Tyr	Glu	Arg	Tyr
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Ile	Asn	Gly	830	Lys	Asp	Thr	835	Phe	Gly	Lys	Ser	Tyr	840	Pro	Asp	Ser	Val
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Asp	Leu	Ser	920	Trp	Ser	Val	925	Val	Val	Asp	Lys	Ala	930	Val	Ser	Tyr	Asn
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			1115				1120						1125				

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Ser	Pro	Ser	Ala	1170	Asp	Asp	Val	Ala	1175	Ile	Phe	Gly	Tyr	Glu	1180	Ile	Tyr
1185	Ser	Ser	Ser	1190	Glu	Thr	Gly	Pro	1195	Phe	Ile	Lys	Ile	Ala	1200	Thr	Val
Ser	Val	Tyr	Asn	1205	Tyr	Val	Asp	Thr	1210	Asp	Val	Val	Asn	Gly	1215	Asn	Val
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<213> Unknown

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<223> Obtained from an environmental sample.

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<223> Alpha amylase, catalytic domain

<221> DOMAIN
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<221> DOMAIN

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<222> (613)...(709)
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<221> DOMAIN
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Thr Asp Arg Phe Ser Asp Gly Asn Ala Ala Asn Asn Pro Thr Gly Pro
 50      55      60
Ala Phe Asp Gly Thr Cys Thr Asn Leu Arg Leu Tyr Cys Gly Gly Asp
 65      70      75      80
Trp Gln Gly Ile Ile Asn Lys Ile Asn Asp Gly Tyr Leu Thr Gly Met
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Gly Val Thr Ala Ile Trp Ile Ser Gln Pro Val Glu Asn Ile Tyr Ser
100      105      110
Val Ile Asn Tyr Ser Gly Val Ser Asn Thr Ala Tyr His Gly Tyr Trp
115      120      125
Ala Arg Asp Phe Lys Lys Thr Asn Pro Ala Tyr Gly Thr Ile Ala Asp
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Phe Gln Asn Leu Ile Ala Ala Ala His Ala Lys Asn Ile Lys Val Ile
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165      170      175
Ser Phe Ala Glu Asn Gly Lys Leu Tyr Asn Asn Gly Thr Leu Leu Gly
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195      200      205
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275      280      285
Val Asn Glu Val Ser Ala Glu Asn His Lys Phe Ala Asn Glu Ser Gly
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325      330      335
Ser Ala Thr Asp Tyr Ala Gln Val Glu Asp Gln Val Thr Phe Ile Asp
340      345      350
Asn His Asp Met Glu Arg Phe His Asp Asn Ser Ala Asn Arg Arg Lys
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385      390      395      400
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405      410      415
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Tyr Gly Thr Thr Gln Glu Arg Trp Ile Asn Asn Asp Val Leu Ile Tyr
435      440      445
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 515 520 525
 Gly His Val Gly Pro Met Met Ala Lys Pro Gly Ala Thr Val Thr Ile
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 Asp Gly Arg Gly Phe Gly Ala Thr Lys Gly Thr Val Tyr Phe Gly Ser
 545 550 555 560
 Thr Ala Val Thr Gly Ala Asn Ile Thr Ala Trp Glu Asp Thr Gln Ile
 565 570 575
 Lys Val Lys Ile Pro Ala Val Ala Gly Val Tyr Asn Ile Lys Val
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 Ala Asn Ile Ala Gly Thr Ser Ser Asn Val His Asp Asn Phe Glu Val
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 Gly Asn Trp Asp Pro Ala Lys Ala Ile Gly Pro Leu Tyr Asn Gln Val
 645 650 655
 Ile Tyr Gln Tyr Pro Thr Trp Tyr Tyr Asp Val Ser Val Pro Ala Gly
 660 665 670
 Lys Thr Ile Glu Phe Lys Phe Leu Lys Lys Gln Gly Ser Thr Val Thr
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<223> Obtained from an environmental sample.

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Gly Leu Thr Asp Asp Pro Ser Leu Ala Gly Arg Glu Ala Pro Arg Leu
1330      1335      1340
Ser Val Lys Leu Ile Val Ala Lys Ser Asp Ala Ser Glu Pro Val Ala
1345      1350      1355      1360
Glu Leu Ser Met Ala Tyr Lys Ala Asp Glu Gly Asp Ala Lys Arg Tyr
1365      1370      1375
Ala Ala Lys Phe Glu Pro Thr Glu Ala Gly Val Tyr Asn Tyr Phe Val
1380      1385      1390
Arg Val Ser Thr Asp Asn Glu Glu Thr Phe Thr Asp Ser Ala Thr Ala
1395      1400      1405
Ser Phe Glu Ala Tyr Ala Asp Pro Asn Asp Thr Met Gly Pro Glu Ala
1410      1415      1420
Pro Ile Leu Ser Glu Ile Thr Val Glu Ser Gly Gln Ala Ala Leu Asn
1425      1430      1435      1440
Trp Thr Pro Ala Gly Glu Glu Thr Ser Gly Tyr Glu Val Tyr Arg Lys
1445      1450      1455
Ala Ala Gly Glu Ala Thr Phe Arg Lys Leu Ala Thr Leu Arg Thr Ala
1460      1465      1470
Thr Ser Tyr Ile Asp Tyr Thr Val Thr Asn Gly Thr Ser Tyr Thr Tyr
1475      1480      1485
Lys Ile Ala Ala Phe Asp Ala Ala Tyr Asn Arg Gly Trp Ser Asn Glu
1490      1495      1500
Gln Thr Val Thr Pro Gln Leu Val Met Val Asp Val Thr Leu Arg Leu
1505      1510      1515      1520
His Leu Pro Asp Tyr Thr Pro Ala Thr Asp Ser Ile Tyr Ile Ala Gly
1525      1530      1535
Thr Leu Asn Asn Trp Asn Ala Ser Gly Gly Lys Leu Thr Val Pro Ser
1540      1545      1550
Gly Ala Thr Gly Arg Ser Val Val Gln Tyr Ala Phe Lys Met Met Ala
1555      1560      1565
Gly Lys Thr Ile Glu Tyr Lys Tyr Thr Arg Gly Thr Trp Ala Thr Glu
1570      1575      1580
Ala Leu Thr Ser His Ala Arg Val Pro Asn Asp Thr Thr Asp Val Ser
1585      1590      1595      1600
Asn Tyr Ala Tyr Ser Asp Glu Lys Thr Asn Met Lys Leu Thr Ile Arg
1605      1610      1615
Asn Gln Gly Gly Gly Lys Met Ile Val Asp Asp Tyr Val Leu Arg Trp
1620      1625      1630
Ala Asp Met Pro Met Ile Val Thr Met Pro Arg Ile Ser Tyr Gly Glu
1635      1640      1645
Asp Ile Ala Phe Ala Thr Ser Asp Ser Ala Phe Asp Leu Lys Ala Asn
1650      1655      1660
Val Pro Tyr Gly Val Ala Phe Thr Ile Asn Gly Gln Pro Leu Pro Lys
1665      1670      1675      1680
Gly Ala Met Asp Asp Arg Gly Asn Val Tyr Val Glu Lys Ile Pro Leu
1685      1690      1695
Lys Pro Gly Glu Asn Thr Phe Ile Leu His Ile Glu Pro Thr Ala Glu
1700      1705      1710
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1715      1720      1725
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1001827087_1.txt

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 Gly Asn Gly Gly Asp Gly Gly Asn Gly Gly Asp Gly Gly Asn Gly Gly
 1765 1770 1775
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 1780 1785 1790
 Ser Gly Ser Leu Val Ala Arg Arg Asp Thr Glu Thr Asp Gly Gln Gly
 1795 1800 1805
 Asn Thr Val Thr Val Ala Thr Val Asp Gly Glu Lys Leu Ala Ala Ala
 1810 1815 1820
 Ile Ala Ala Leu Lys Asp Gly Asn Gly Val Val Glu Ile Arg Leu Pro
 1825 1830 1835 1840
 Asp Thr Lys Gly Ser Val Arg Phe Ala Ile Pro Leu Lys Ala Ala Ala
 1845 1850 1855
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 1860 1865 1870
 Asp Gly Ser Tyr Ala Leu Pro Leu Ser Val Leu Glu Pro Glu Glu Leu
 1875 1880 1885
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 1890 1895 1900
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 1905 1910 1915 1920
 Glu Ala Ala Gly Ala Lys Val Leu Gly Gly Pro Ile Ala Tyr Thr Val
 1925 1930 1935
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 1940 1945 1950
 Val Tyr Ala Glu Arg Ser Met Thr Leu Lys Gly Val Val Asp Pro Arg
 1955 1960 1965
 His Ala Ser Val Phe Arg Val Asp Asp Glu Ala Asn Lys Leu Ser Phe
 1970 1975 1980
 Val Pro Ala Leu Phe Thr Ala Ala Gly Asn Gly Gln Thr Ala Val Ala
 1985 1990 1995 2000
 Phe Lys Arg Pro Gly Asn Ser Val Tyr Ile Val Ala Ser Ser Glu Arg
 2005 2010 2015
 Asp Phe Ala Asp Val Ala Asp His Trp Ala Arg Pro Asp Ile Leu Leu
 2020 2025 2030
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 2035 2040 2045
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 2050 2055 2060
 Ser Leu Gly Met Ala Glu Ala Asp Gly Thr Gly Gly Phe Arg Asp Val
 2065 2070 2075 2080
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 2085 2090 2095
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 2100 2105 2110
 Ile Thr Arg Glu Gln Met Ala Val Met Ala Ala Lys Ala Leu Ala Phe
 2115 2120 2125
 Ala Gly Lys Pro Ala Ala Gly Gly Ala Pro Gln Ala Gly Glu Thr Leu
 2130 2135 2140
 Ala Ala Phe Ala Asp Gly Ala Ala Val Ser Ala Trp Ala Arg Asp Ala
 2145 2150 2155 2160
 Val Ala Arg Thr Val Gln Ala Gly Leu Met Gln Gly Gly Ser Gly Gly
 2165 2170 2175
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 2180 2185 2190
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 2195 2200 2205

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 <212> DNA
 <213> Bacteria

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60
 120

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<210> 586
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 <212> PRT
 <213> Bacteria

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<221> DOMAIN
 <222> (40)...(425)
 <223> Alpha amylase, catalytic domain

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Ala Gly Ser Val Pro Val Asn Gly Thr Met Met Gln Tyr Phe Glu Trp
35     40     45
Tyr Leu Pro Asp Asp Gly Thr Leu Trp Thr Lys Val Ala Asn Asn Ala
50     55     60
Gln Ser Leu Ala Asn Leu Gly Ile Thr Ala Leu Trp Leu Pro Pro Ala
65     70     75     80
Tyr Lys Gly Thr Ser Ser Ser Asp Val Gly Tyr Gly Val Tyr Asp Leu
85     90     95
Tyr Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Val Arg Thr Lys Tyr
100    105    110
Gly Thr Lys Thr Gln Tyr Ile Gln Ala Ile Gln Ala Ala His Thr Ala
115    120    125
Gly Met Gln Val Tyr Ala Asp Val Val Phe Asn His Lys Ala Gly Ala
130    135    140
Asp Gly Thr Glu Leu Val Asp Ala Val Glu Val Asn Pro Ser Asp Arg
145    150    155    160

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Tyr	His	Phe	Asp	Gly	Thr	Asp	Trp	Asp	Glu	Ser	Arg	Lys	Leu	Asn	Arg
Ile	Tyr	Lys	Phe	Arg	Gly	Thr	Gly	Lys	Ala	Trp	Asp	Trp	Glu	Val	Asp
Thr	Glu	Asn	Gly	Asn	Tyr	Asp	Tyr	Leu	Met	Tyr	Ala	Asp	Leu	Asp	Met
Asp	His	Pro	Glu	Val	Val	Ser	Glu	Leu	Lys	Asn	Trp	Gly	Lys	Trp	Tyr
Val	Thr	Thr	Thr	Asn	Ile	Asp	Gly	Phe	Arg	Leu	Asp	Ala	Val	Lys	His
Ile	Lys	Tyr	Ser	Phe	Phe	Pro	Asp	Trp	Leu	Ser	Tyr	Val	Arg	Thr	Gln
Thr	Gln	Lys	Pro	Leu	Phe	Ser	Val	Gly	Glu	Phe	Trp	Ser	Tyr	Asp	Ile
Ser	Lys	Leu	His	Asn	Tyr	Ile	Thr	Lys	Thr	Asn	Gly	Ser	Met	Ser	Leu
Phe	Asp	Ala	Pro	Leu	His	Asn	Asn	Phe	Tyr	Ile	Ala	Ser	Lys	Ser	Gly
Gly	Tyr	Phe	Asp	Met	Arg	Thr	Leu	Leu	Asn	Asn	Thr	Leu	Met	Lys	Asp
Gln	Pro	Thr	Leu	Ala	Val	Thr	Leu	Val	Asp	Asn	His	Asp	Thr	Glu	Pro
Gly	Gln	Ser	Leu	Gln	Ser	Trp	Val	Glu	Pro	Trp	Phe	Lys	Pro	Leu	Ala
Tyr	Ala	Phe	Ile	Leu	Thr	Arg	Gln	Glu	Gly	Tyr	Pro	Cys	Val	Phe	Tyr
Gly	Asp	Tyr	Tyr	Gly	Ile	Pro	Lys	Tyr	Asn	Ile	Pro	Ala	Leu	Lys	Ser
Lys	Leu	Asp	Pro	Leu	Leu	Ile	Ala	Arg	Arg	Asp	Tyr	Ala	Tyr	Gly	Thr
Gln	His	Asp	Tyr	Ile	Asp	Ser	Ala	Asp	Ile	Ile	Gly	Trp	Thr	Arg	Glu
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Lys	Thr	Phe	Tyr	Asp	Leu	Thr	Gly	Asn	Arg	Ser	Asp	Thr	Val	Thr	Ile
Asn	Ala	Asp	Gly	Trp	Gly	Glu	Phe	Lys	Val	Asn	Gly	Gly	Ser	Val	Ser
Ile	Trp	Val	Pro	Lys	Thr	Ser	Thr	Thr	Ser	Gln	Ile	Thr	Phe	Thr	Val
Asn	Asn	Ala	Thr	Thr	Val	Trp	Gly	Gln	Asn	Val	Tyr	Val	Val	Gly	Asn
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Pro	Ser	Ser	Tyr	Pro	Thr	Trp	Val	Val	Thr	Val	Pro	Leu	Pro	Gln	Ser
Gln	Asn	Ile	Gln	Phe	Lys	Phe	Ile	Lys	Lys	Asp	Gly	Ser	Gly	Asn	Val
Ile	Trp	Glu	Asn	Ile	Ser	Asn	Arg	Thr	Tyr	Thr	Val	Pro	Thr	Ala	Ala
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 <213> Thermomyces lanuginosus ATCC 200065

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gcgttcacag agcgaagttt gtttaattga gtcataaaaa c atg tta ttc caa ccg 716
Met Leu Phe Gln Pro
1 5

act ttg tgc gcg gcc ctt gga ctc gcc gcc ttg atc gtc caa ggc gga 764
Thr Leu Cys Ala Ala Leu Gly Leu Ala Ala Leu Ile Val Gln Gly Gly
10 15 20

gaa gcc aga cct gaa aca acc gtc cca cat gca acg ggc tcg ctc gac 812
Glu Ala Arg Pro Glu Thr Thr Val Pro His Ala Thr Gly Ser Leu Asp
25 30 35

gac ttc ctc gcc gca cag agt ccg att gct ttc caa ggc atc ctg aac 860
Asp Phe Leu Ala Ala Gln Ser Pro Ile Ala Phe Gln Gly Ile Leu Asn
40 45 50

aat atc ggg cct agc gga gcg tac tcg gaa ggt gtc aat ccg ggt gtg 908
Asn Ile Gly Pro Ser Gly Ala Tyr Ser Glu Gly Val Asn Pro Gly Val
55 60 65

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Val Ile Ala Ser Pro Ser Lys Gln Asp Pro Asp
70 75 80

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Tyr

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Phe Tyr Thr Trp Val Arg Asp Ala Ala Leu Thr Val Gln Tyr Leu Val
85 90 95

gag gag ctg gtt gca gga aat gcc agt ctt cag ttc ctc att cag gac 1104
Glu Glu Leu Val Ala Gly Asn Ala Ser Leu Gln Phe Leu Ile Gln Asp
100 105 110

tac atc agc tcc cag gca cga ctg cag acg gtg gaa aat cca tcc ggc 1152
Tyr Ile Ser Ser Gln Ala Arg Leu Gln Thr Val Glu Asn Pro Ser Gly
115 120 125

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ctc cgc gcc att gcc atg att tcg ttt gcc aat tac ctg att Leu Arg Ala Ile Ala Met Ile Ser Phe Ala Asn Tyr Leu Ile 165 170 175	1290
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cgc aat gac ttg tcc tat gtc tcg cag cat tgg aac gaa aca act ttt Arg Asn Asp Leu Ser Tyr Val Ser Gln His Trp Asn Glu Thr Thr Phe 195 200 205	1447
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gct agc aag ctc ggc cat acc tgc gac aac tgc ggg tcc caa gca ccg Ala Ser Lys Leu Gly His Thr Cys Asp Asn Cys Gly Ser Gln Ala Pro 235 240 245 250	1656
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<210> 588

<211> 80

<212> PRT

<213> Thermomyces lanuginosus ATCC 200065

<400> 588

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20      25      30
Thr Gly Ser Leu Asp Asp Phe Leu Ala Ala Gln Ser Pro Ile Ala Phe
35      40      45
Gln Gly Ile Leu Asn Asn Ile Gly Pro Ser Gly Ala Tyr Ser Glu Gly
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<211> 95

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<213> Thermomyces lanuginosus ATCC 200065

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20      25      30
Asp Tyr Ile Ser Ser Gln Ala Arg Leu Gln Thr Val Glu Asn Pro Ser
35      40      45
Gly Ser Leu Ser Ser Gly Gly Leu Gly Glu Pro Lys Phe His Val Asp
50      55      60
Glu Thr Ala Phe Thr Asp Ser Trp Gly Arg Pro Gln Arg Asp Gly Pro
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Pro Leu Arg Ala Ile Ala Met Ile Ser Phe Ala Asn Tyr Leu Ile
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<210> 590

<211> 32

<212> PRT

<213> Thermomyces lanuginosus ATCC 200065

<400> 590

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20      25      30

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<213> Thermomyces lanuginosus ATCC 200065

<400> 591

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 35 40 45
 Leu Gln Ser Tyr Trp Thr Gly Ser His Ile Leu Ala Asn Thr Gly Gly
 50 55 60
 Gly Arg Ser Gly Lys Asp Val Ser Thr Ile Leu Gly Val Ile Gly Ser
 65 70 75 80
 Phe Asp Pro Asn Ala Asp Cys Asp Asp Val Thr Phe Gln Pro Cys Ser
 85 90 95
 Ala Arg Ala Leu Ala Asn His Lys Gln Val Val Asp Ser Phe Arg Ser
 100 105 110
 Ile Tyr Ala Ile Asn Ala Gly Ile Pro Ser Gly Ser Ala Val Ala Val
 115 120 125
 Gly Arg Tyr Pro Glu Asp Val Tyr Gln Gly Gly His Pro Trp Tyr Leu
 130 135 140
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 145 150 155 160
 Asn His Val Gly His Ile Asp Ile Asn Ala Val Asn Leu Asp Phe Phe
 165 170 175
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<210> 592

<211> 197

<212> PRT

<213> Thermomyces lanuginosus ATCC 200065

<400> 592

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 Thr Gln Pro Pro Glu Arg Pro Ala Cys Thr Pro Pro Ser Glu Val Thr
 85 90 95
 Leu Thr Phe Asn Ala Leu Val Asp Thr Ala Phe Gly Gln Asn Ile Tyr
 100 105 110
 Leu Val Gly Ser Ile Pro Glu Leu Gly Ser Trp Asp Pro Ala Asn Ala
 115 120 125
 Leu Leu Met Ser Ala Lys Ser Trp Thr Ser Gly Asn Pro Val Trp Thr
 130 135 140
 Leu Ser Ile Ser Leu Pro Ala Gly Thr Ser Phe Glu Tyr Lys Phe Ile
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 Arg Lys Asp Asp Gly Ser Ser Asp Val Val Trp Glu Ser Asp Pro Asn
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<212> DNA

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<213> Thermomyces lanuginosus ATCC 200065

<400> 593

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<210> 594

<211> 617

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<213> Thermomyces lanuginosus ATCC 200065

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<222> (1)...(23)

<400> 594

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160          165          170          175
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Ser	Ala	Thr	465	Ser	Ala	Thr	Gly	Pro	470	Tyr	Ala	Thr	Pro	Thr	Asn	Thr	Ala	
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<213> Unknown

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Phe	Thr	Arg	Asp	Trp	Ile	Asn	His	Val	Gln	Trp	Ala	Thr	Ser	Glu	Asp
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<212> DNA
<213> Unknown

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<210> 606
<211> 719
<212> PRT
<213> Unknown

<220>
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<221> SIGNAL
<222> (1)...(34)

<221> DOMAIN
<222> (53)...(432)
<223> Alpha amylase, catalytic domain

<221> DOMAIN
<222> (620)...(716)
<223> Starch binding domain

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<221> DOMAIN
<222> (533)...(616)
<223> IPT/TIG domain

<221> DOMAIN
<222> (444)...(528)
<223> Alpha amylase, C-terminal all-beta domain

<400> 606

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Ser Ala Ala Pro Asp Thr Ser Val Ser Asn Lys Gln Asn Phe Ser Thr
 35      40      45
Asp Val Ile Tyr Gln Ile Phe Thr Asp Arg Phe Ala Asp Gly Asn Pro
 50      55      60
Ala Asn Asn Pro Thr Gly Ser Ala Tyr Ser Pro Gly Cys Thr Asn Leu
 65      70      75
Arg Leu Tyr Cys Gly Gly Asp Trp Lys Gly Ile Thr Asp Lys Ile Asn
 85      90      95
Ala Gly Tyr Phe Thr Asn Met Gly Val Thr Ala Leu Trp Ile Ser Gln
100      105      110
Pro Val Glu Asn Ile Tyr Ser Val Ile Asn Tyr Ser Gly Val Asn Asn
115      120      125
Thr Ser Tyr His Gly Tyr Trp Ala Arg Asp Phe Lys Lys Thr Asn Pro
130      135      140
Tyr Tyr Gly Thr Phe Ala Asp Leu Gln Thr Leu Ile Asn Ala Ala His
145      150      155
Ala Lys Gly Ile Lys Ile Val Ile Asp Phe Ala Pro Asn His Thr Ser
165      170      175
Pro Ala Leu Glu Thr Asp Ser Ser Phe Ala Glu Asn Gly Arg Leu Tyr
180      185      190
Asp Asn Gly Thr Leu Leu Gly Gly Tyr Thr Asn Asp Thr Asn Gly Tyr
195      200      205
Phe His His Asn Gly Gly Thr Asp Phe Ser Ser Leu Glu Asp Gly Ile
210      215      220
Tyr Arg Asn Leu Tyr Asp Leu Ala Asp Leu Asn His Asn Val Ser Arg
225      230      235
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Ile Asp Gly Ile Arg Val Asp Ala Val Lys His Met Pro Met Gly Trp
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Gln Lys Asn Trp Met Ser Tyr Ile Tyr Gly Tyr Lys Pro Val Phe Thr
275      280      285
Phe Gly Glu Trp Phe Leu Gly Thr Asn Glu Asn Asp Pro Ser Asn Val
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Tyr Phe Ala Asn Glu Ser Gly Met Ser Leu Leu Asp Phe Arg Phe Ala
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Gln Lys Val Arg Gln Val Leu Arg Asp Asn Thr Asp Thr Met Tyr Gly
325      330      335
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340      345      350
Asp Gln Val Thr Phe Ile Asp Asn His Asp Met Glu Arg Phe Lys Ser
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Gly Ser Val Ser Asn Arg Arg Val Glu Gln Ala Leu Ala Leu Leu Leu
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Thr Ser Arg Gly Val Pro Ala Ile Tyr Tyr Gly Thr Glu Gln Tyr Met
385      390      395
Ser Gly Gly Thr Asp Pro Asp Asn Arg Ala Met Met Ser Ser Phe Ser
405      410      415
Glu Thr Thr Thr Ala Phe Asn Val Ile Lys Lys Leu Ala Pro Leu Arg
420      425      430
Lys Ser Asn Pro Ala Ile Ala Tyr Gly Ser Thr Gln Gln Arg Trp Ile
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<210> 607

<211> 3705

<212> DNA

<213> Unknown

<220>

<223> Obtained from an environmental sample

<400> 607

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<210> 608
 <211> 1235
 <212> PRT
 <213> Unknown

<220>
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<221> DOMAIN
 <222> (169)...(629)
 <223> Alpha amylase, catalytic domain

<221> DOMAIN
 <222> (727)...(801)
 <223> Fibronectin type III domain

<221> DOMAIN
 <222> (646)...(724)
 <223> Alpha amylase, C-terminal all-beta domain

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          35          40          45
Gly Ala Val Lys Ala Gly Thr Lys Val Thr Leu Arg Leu Ala Ala Lys
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Lys Gly Asp Leu Thr Lys Ala Asp Val Tyr Val Lys Asn Ala Thr Thr
          65          70          75          80
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Gly	Leu	Phe	Gln	Leu	Thr	Val	Tyr	Asp	Pro	Asn	Tyr	Lys	Thr	Pro
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Trp	Met	Lys	Glu	Ala	Val	Val	Tyr	Gln	Ile	Phe	Pro	Asp	Arg	Phe
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Leu	Ser	Glu	Glu	Ala	Arg	Lys	Gln	Val	Glu	Glu	Lys	Phe	Lys	Gln
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His	Pro	Ser	Glu	Leu	Asn	Asn	Asp	Ala	Leu	Ala	Asn	Tyr	Ile	Phe
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				1075															
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	Asn	Asp	Asp	Asp	Val	Ser	Tyr	Tyr	Gly	Tyr	Gly	Thr	Ile	Gly	Thr	Asp			
				1105															
	Leu	Lys	Val	Thr	Val	His	Asn	Glu	Gly	Asn	Asn	Thr	Met	Ile	Val	Gln			
				1125															
	Asp	Arg	Ile	Leu	Arg	Trp	Ile	Asp	Met	Pro	Val	Val	Ile	Glu	Glu	Val			
				1140															
	Gln	Lys	Gln	Gly	Ser	Gln	Val	Thr	Ile	Lys	Gly	Asn	Ala	Ile	Lys	Asn			
				1155															
	Gly	Val	Leu	Thr	Ile	Asn	Gly	Glu	Arg	Val	Pro	Ile	Asp	Gly	Arg	Met			

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1170 1175 1180
 Ala Phe Ser Tyr Thr Phe Ala Pro Ala Ser His Gln Lys Glu Val Leu
 1185 1190 1195 1200
 Ile His Ile Glu Pro Ser Ala Glu Ser Lys Thr Ala Ile Phe Asn Asn
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<210> 609
 <211> 1764
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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 caggtgaaca acaccatgta ccaggcattt tattgggatg cctaccctgg cctttgggcc 180
 aatttaccgg ccatggcggc ccctttggcc gagcgtggca ttacctgat gtggttgccg 240
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 gatctgggcg agtttaacca aaaaggcacc accgccacc gttacggtac tcgtcagcag 360
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 ttaaccacc gcatgggcg cgatgcacag gagaatattc ctggccttgg cctggcctgg 480
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<210> 610
 <211> 587
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(31)

<221> DOMAIN
 <222> (44)...(395)
 <223> Alpha amylase, catalytic domain

<400> 610
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Ala	Phe	Tyr	Trp	Asp	Ala	Tyr	Pro	Gly	Leu	Trp	Ala	Asn	Leu	Pro	Ala
Met	Ala	Ala	Pro	Leu	Ala	Glu	Arg	Gly	Ile	Thr	Ser	Met	Trp	Leu	Pro
Pro	Ala	Ala	Lys	Gly	Met	Asn	Gly	Thr	Phe	Ser	Val	Gly	Tyr	Asp	Val
Tyr	Asp	Phe	Trp	Asp	Leu	Gly	Glu	Phe	Asn	Gln	Lys	Gly	Thr	Thr	Ala
Thr	Arg	Tyr	Gly	Thr	Arg	Gln	Gln	Leu	Gln	Gln	Ala	Leu	Ser	Ala	Leu
Asp	Gln	Leu	Gly	Ile	Gln	Ala	Tyr	Phe	Asp	Val	Val	Phe	Asn	His	Arg
Met	Gly	Ala	Asp	Ala	Gln	Glu	Asn	Ile	Pro	Gly	Phe	Gly	Leu	Ala	Trp
Thr	Glu	Tyr	His	Leu	Gln	Gly	Arg	Gln	Ala	His	Tyr	Thr	Gln	Gln	Asn
Trp	Gly	Tyr	Leu	Trp	His	Asp	Phe	Asp	Trp	Asn	Trp	Thr	Ala	Phe	Asn
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His	Phe	Pro	Tyr	Leu	Met	Gly	Glu	Asp	Val	Asp	Tyr	Asn	Arg	Phe	Glu
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Gly	Gly	Leu	Val	Asn	Ser	Gln	His	Arg	Asp	Arg	Ala	Val	Thr	Phe	Val
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Val	Ile	Asn	Tyr	Lys	Asn	Gln	Ala	Tyr	Ala	Tyr	Ile	Leu	Leu	Arg	Glu
His	Gly	Val	Pro	Thr	Val	Phe	Ala	Arg	Asp	Tyr	Asp	Glu	Phe	Gly	Met
Ala	Pro	Thr	Leu	Asp	Lys	Leu	Ile	Glu	Ala	Arg	Arg	Tyr	Phe	Ala	Tyr
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His	Gln	Pro	Asn	Thr	Thr	Phe	Tyr	Asp	Tyr	Thr	Gly	Asn	Val	Ser	Gly
Thr	Val	Thr	Thr	Asn	Ala	Gln	Gly	Tyr	Gly	Ser	Phe	Pro	Val	Thr	Met
Thr	Glu	Ser	Thr	Gly	Trp	Ser	Val	Trp	Val	Pro	Gln	Ser	Ser	Gly	Gly
Thr	Gln	Pro	Gly	Ser	Ile	Thr	Leu	Arg	Ile	Thr	Lys	Asp	Val	Gly	Tyr
Gly	Phe	Ser	Leu	Phe	Phe	Thr	Gly	Ser	Ser	Ala	Glu	Leu	Thr	Asn	Trp
Gly	Gly	Gly	Ile	Glu	Gly	Thr	Trp	Thr	Ser	Gly	His	Val	Trp	Glu	Val
Thr	Ile	Pro	Asp	Pro	Gly	Asn	Phe	Glu	Trp	Lys	Thr	Arg	Lys	Gly	Pro

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Thr Gly Gly Ser Gly Gln Asp Trp Glu Ser Asp Ser Asn His Asn Gln
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 Thr Asn Leu His Pro Ser Phe Asn Gly Gly Phe
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<210> 611
 <211> 3804
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

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<210> 612
 <211> 1268
 <212> PRT
 <213> Unknown

<220>
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<221> DOMAIN
 <222> (646)...(724)
 <223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN
 <222> (727)...(801)
 <223> Fibronectin type III domain

<221> DOMAIN
 <222> (169)...(629)
 <223> Alpha amylase, catalytic domain

<400> 612

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Asp	Arg	Leu	Lys	His	Asn	Thr	Trp	Asp	Thr	Leu	Tyr	Arg	Gln	Pro	Phe
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	50				55					60					
Lys	Gly	Asp	Leu	Thr	Lys	Ala	Asp	Val	Tyr	Val	Lys	Asn	Ala	Thr	Thr
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Gly	Thr	Ala	Lys	Val	Tyr	Thr	Met	Glu	Lys	Val	Gly	Val	Leu	Gly	Asp
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Asp	Glu	Tyr	Trp	Glu	Ala	Ala	Phe	Thr	Pro	Ser	Ala	Pro	Gly	Val	Tyr
			100				105						110		
Gly	Tyr	Lys	Phe	Ile	Ala	Val	Asp	Ala	Gly	Met	Lys	Ala	Glu	Tyr	Gly
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Glu	Asp	Thr	Lys	Glu	Gly	Gln	Trp	Gly	Arg	Ala	Val	Asp	Lys	Asn	Ala
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Gly	Leu	Phe	Gln	Leu	Thr	Val	Tyr	Asp	Pro	Asn	Tyr	Lys	Thr	Pro	Asp
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Gln	Pro	Ile	Glu	His	Arg	Asp	Trp	Ser	Asp	Leu	Pro	Asp	Asn	Pro	Arg
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Gln	Ser	Leu	Gly	Val	Asn	Thr	Ile	Tyr	Leu	Asn	Pro	Ile	Ala	Asn	Ala
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Ala	Tyr	Glu	Tyr	Trp	Glu	Ala	Val	Tyr	Asp	Leu	Met	Asn	Glu	Lys	Gly
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His	Pro	Ser	Glu	Leu	Asn	Asn	Asp	Ala	Leu	Ala	Asn	Tyr	Ile	Phe	Arg
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Trp	Arg	Leu	Asp	Val	Ala	Asn	Glu	Val	Asp	Pro	Ala	Phe	Trp	Arg	Glu
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Lys	Glu	Gly	Glu	Gln	Pro	Leu	Ile	Leu	Gly	Glu	Ile	Trp	Asp	Asp	Ala
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Ser	Lys	Tyr	Phe	Leu	Gly	Asp	Gln	Tyr	Asp	Ser	Val	Met	Asn	Tyr	Arg
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Phe	Tyr	Ala	Leu	Met	Asn	Leu	Ile	Gly	Ser	His	Asp	Thr	Ala	Arg	Ala
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Ile	Lys	Thr	Val	Tyr	Ala	Gln	Gly	Asp	Val	Tyr	Val	Phe	Ala	Arg	Gln
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Thr	Val	Thr	Ala	Glu	Met	Pro	Ala	Ala	Val	Ser	Asn	Leu	Gln	Ala	Ser
				725					730					735	
Ala	Ser	Asp	Gly	Cys	Val	Thr	Leu	Thr	Trp	Glu	Gly	Asn	Ala	Ser	Arg
			740					745					750		
Tyr	Arg	Ile	Tyr	Glu	Ser	Thr	Leu	Lys	Gly	Ala	Gly	Tyr	Thr	Met	Val
		755					760					765			
Gln	Glu	Thr	Glu	Thr	Thr	Ser	Ala	Thr	Ile	Gly	Ser	Leu	Thr	Asn	Gly
	770					775					780				
Thr	Ala	Tyr	Tyr	Phe	Ala	Val	Ala	Ala	Val	Asp	Glu	Asn	Gly	Asn	Glu
785					790					795					800
Ser	Pro	Lys	Val	Glu	Thr	Asn	Arg	Val	Val	Pro	His	Tyr	Pro	Leu	Thr
				805					810					815	
Ser	Asp	Asn	Val	Gln	Phe	Val	Thr	Thr	Leu	Ser	Asp	Ala	Thr	Leu	Asp
			820					825					830		

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Leu Ser Lys Pro Gln Gln Val Asp Val His Val Asn Ile Asp Asn Val
 835 840 845
 Thr Ser Lys Gly Ala Ala Asp Gly Leu Gln Ala Val Leu Gln Val Lys
 850 855 860
 Gly Pro His Asp Glu Thr Trp Lys Glu Tyr Arg Ala Ala Tyr Gln Gly
 865 870 875 880
 Gln Asp Gly Asp Ala Asn Val Phe Arg Ala Phe Thr Pro Leu Ala
 885 890 895
 Ala Gly Thr Tyr Thr Tyr Arg Tyr Ala Leu Thr Thr Asn Leu Gly Glu
 900 905 910
 Glu Trp Val Tyr Thr Glu Glu Lys Gln Val Thr Phe Ala Ala Asp Asn
 915 920 925
 Ser Asp Gln Ile Ala Pro Ala Asp Ala Ile Glu Leu Arg Gln Pro Ala
 930 935 940
 Val Glu Ser Gly Gln Val Asn Leu Ser Trp Thr Phe Val Gly Lys Lys
 945 950 955 960
 Asp Gly Asp Ala Tyr Leu Leu Ala Ile Glu Arg Asn Gly Asp Ile Val
 965 970 975
 His Thr Thr Thr Ser Ile Gly Asp Ser Phe Thr Asp Tyr Asp Val Glu
 980 985 990
 Asn Gly Thr Glu Tyr Thr Tyr Val Val Lys Leu Tyr Asp Arg Ala Gly
 995 1000 1005
 Asn Val Val Ala Ser Asn Thr Val Lys Val Thr Pro Asp Ile Val Met
 1010 1015 1020
 Val Lys Val Ile Phe Lys Val Arg Ala Pro Asp Tyr Thr Pro Leu Asp
 1025 1030 1035 1040
 Ala Arg Ile Thr Ile Pro Asn Ser Leu Asn Gly Trp Asn Thr Gly Ala
 1045 1050 1055
 Trp Glu Met Ser Arg Asn Gly Ala Val Thr Pro Asp Trp Gln Phe Thr
 1060 1065 1070
 Val Glu Val Gln Glu Gly Glu Thr Ile Thr Tyr Lys Tyr Val Lys Gly
 1075 1080 1085
 Gly Ser Trp Asp Gln Glu Gly Leu Ala Asp His Thr Arg Glu Asp Asp
 1090 1095 1100
 Asn Asp Asp Asp Val Ser Tyr Tyr Gly Tyr Gly Thr Ile Gly Thr Asp
 1105 1110 1115 1120
 Leu Lys Val Thr Val His Asn Glu Gly Asn Asn Thr Met Ile Val Gln
 1125 1130 1135
 Asp Arg Ile Leu Arg Trp Ile Asp Met Pro Val Val Ile Glu Glu Val
 1140 1145 1150
 Gln Lys Gln Gly Ser Gln Val Thr Ile Lys Gly Asn Ala Ile Lys Asn
 1155 1160 1165
 Gly Val Leu Thr Ile Asn Gly Glu Arg Val Pro Ile Asp Gly Arg Met
 1170 1175 1180
 Ala Phe Ser Tyr Thr Phe Ala Pro Ala Ser His Gln Lys Glu Val Leu
 1185 1190 1195 1200
 Ile His Ile Glu Pro Ser Ala Glu Ser Lys Thr Ala Ile Phe Asn Asn
 1205 1210 1215
 Asp Gly Gly Ala Ile Ala Lys Asn Thr Lys Asp Tyr Val Leu Asn Leu
 1220 1225 1230
 Glu Thr Lys Gln Phe Lys Lys Leu Leu Glu Ser Thr Ser Arg Ala Ala
 1235 1240 1245
 Ala Gly Pro Ser Ile Phe His Pro Gly Gly Val Pro Gly Lys Cys Thr
 1250 1255 1260
 Gln Phe Ala Leu
 1265

<210> 613
 <211> 1764
 <212> DNA
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<400> 613
 atgcaaaaca cagcgaaaaa ctccatctgg cagagggtgc gccacagcgc cattgcctta 60
 tccgctctca gcttatcctt tggcctgcag gccagcgagt taccacaaat tccaccacag 120

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cagggtgaaca acaccatgta ccaggcattt tattgggatg cctaccctgg cttttggggc 180
aatttaccgg ccatggcggc ccctttggcc gagcgtggca ttacctcgat gtggttgccg 240
cccgccgccca aaggcatgaa tgggtacttt agtgtcggtt acgatgtata cgattttctg 300
gatctggggcg agtttaacca aaaaggcacc accgccacc gttacggtac tcgtcagcag 360
ctgcaacaag cactgagtgc tctggaccaaa ctgggtattc aggcctatct tgatgtggtg 420
tttaaccacc gcatggggcg cgatgcacag gagaatattc ctggctttgg cctggcctgg 480
accgagtatc atctgcaagg tcgtcaggcg cattataccc agcaaaaactg gggctacttg 540
tggcacgact ttgactggaa ctggaccgcg tttaatggct ccgacaatca gctctacccc 600
ggcaaatggg ggggcaatac cttccacttc ccttatttga tgggtgagga tgtcgattac 660
aaccgctttg aagtgcagca ggaaatgaaa gcctgggggg agtggatcat caacagcggt 720
ggctttagcg gctttcggat ggatgccatt gcccatgttg ataccgattt taccctgtgac 780
tggatcaatc acgtacagtg ggccaccagc gaggatgtgt tctttgtcgc agaagcctgg 840
gtcagtgata taaacggcta tctggatgca gtcaatacgc cgcattttgcg cgcttttgat 900
ttcaatttgc gcgaagactt tgttgcttta agcagtgggt gcaaagacat gcgctggtgg 960
ggtggtctgg ttaatagcca gcaccgtgat cgggcggtca cttttgtcga taaccacgat 1020
accagccggg ccggcaaccc ttatggcatg ccgcagggtg tcaactacaa gaaccaggcc 1080
tacgcttaca ttctgttgcg tgagcatggg gtgccgactg tgtttgcccg cgattacgac 1140
gaattttggca tggcgccaac gctggataaa ttgattgaag cacggcggtt ctttgccat 1200
ggtcctggcc atgagtactc cggcaatacc gagggcgtct acgcctatgt gcgcgaaggg 1260
cttagcactg tgccgggcac cggctctggt atgctgatat cgggtcgaaa ctgggggtgg 1320
cagcagtctg tcaccatcaa cagccatcag ccgaatacca ctttttacga ttataccggc 1380
aatgtcagtg gcactgtgac caccaatgcg cagggttatg gcagcttccc ggtcactatg 1440
acggaaaagta ccggttggtc agtctgggta ccacaatcca gtggtggcac tcagccggga 1500
tccattaccc tgccgataac caaggatgtt ggctatggtt tttcactggt cttcactggc 1560
agcagtgcag agctcactaa ttggggcggc ggtattgaag gcacctggac cagtggccat 1620
gtctgggaag tgaccatccc ggtccggggc aactttgaat ggaaaaccg taaaggccca 1680
accggtggca ttgggtcagga ctgggaaagt ggcagcaacc acaaccagac caatttgcac 1740
cccagtttta atggtggggt ttaa

```

<210> 614
 <211> 587
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(31)

<221> DOMAIN
 <222> (44)...(395)
 <223> Alpha amylase, catalytic domain

```

<400> 614
Met Gln Asn Thr Ala Lys Asn Ser Ile Trp Gln Arg Val Arg His Ser
 1          5          10          15
Ala Ile Ala Leu Ser Ala Leu Ser Leu Ser Phe Gly Leu Gln Ala Ser
          20          25          30
Glu Leu Pro Gln Ile Pro Pro Gln Gln Val Asn Asn Thr Met Tyr Gln
          35          40          45
Ala Phe Tyr Trp Asp Ala Tyr Pro Gly Leu Trp Ala Asn Leu Pro Ala
          50          55          60
Met Ala Ala Pro Leu Ala Glu Arg Gly Ile Thr Ser Met Trp Leu Pro
65          70          75          80
Pro Ala Ala Lys Gly Met Asn Gly Thr Phe Ser Val Gly Tyr Asp Val
          85          90          95
Tyr Asp Phe Trp Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Thr Ala
          100          105          110
Thr Arg Tyr Gly Thr Arg Gln Gln Leu Gln Gln Ala Leu Ser Ala Leu
          115          120          125
Asp Gln Leu Gly Ile Gln Ala Tyr Phe Asp Val Val Phe Asn His Arg
          130          135          140
Met Gly Ala Asp Ala Gln Glu Asn Ile Pro Gly Phe Gly Leu Ala Trp
145          150          155          160
Thr Glu Tyr His Leu Gln Gly Arg Gln Ala His Tyr Thr Gln Gln Asn
          165          170          175
Trp Gly Tyr Leu Trp His Asp Phe Asp Trp Asn Trp Thr Ala Phe Asn

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2013201807

<210>	615
<211>	1752
<212>	DNA
<213>	Unknown

<220>
<223> obtained from an environmental sample

<400> 615							
atgtccgaaa	gaggcgctcag	gagagccgtg	cgcaccgccc	tggctcgggtt	ggcggcgggcc		60
gccaccgccg	cggtgacgct	cggcgctccc	accgcccgagg	ccgcgccggc	cggaacaag		120
gacgtcaccg	ccgtcctggt	ctcctggaac	ttcaactcca	tcgcccgcga	gtgcaccgag		180
cgactcggcc	ccgcgggcta	cggcttcgtg	caggtctcgc	cgccccagga	gcacatccag		240
ggctccgcgt	ggtggaccca	gtaccagccc	gtgagttaca	acatctccgg	cggctcggcc		300
aacgagcagc	agttccgctc	gatggtaaac	acctgtaaca	acgcgggtgt	cggcgctcctc		360
gtgggacgcg	tcatacaacca	catgtccgcc	ggctcgggca	ccggcaccgg	cggcacctcc		420

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tacagcaagt	acaactaccc	cggcctctac	agcggcaacg	acttcggttc	ctgccgcgag	480
gacatccggg	gcagcgacta	caccaacgac	cgctggcggg	tccagaactg	cgagctggtc	540
ggcctcgcg	acctggacac	cggcagcgcc	tacgtccagc	agcagatcgc	cggctacatg	600
aaccgtctgc	tggtgctggg	cgctgcgggc	ttccgggtgg	acgccgtcaa	gcacatcccg	660
gcccggcaca	tgagcagat	ccgcgcccgg	gtcaacggcg	gcgacgtctt	ctggaaccag	720
gaggtcatct	acggcgccgg	cgaggcgatc	acgcccagag	agtacctcaa	caccgggtcac	780
gtgcaggagt	tccgctacgc	gttcgacctc	aagcggatgt	tccagggcga	ccggatcgcc	840
aatctgcaga	acttcggcga	gtcatggggc	tacatgccct	ccaaccgctc	cggcgtcttc	900
gtcgacaacc	acgacaccga	gcgcaacggc	tccaccctca	gctacaagga	caacgccgcc	960
tacaccctgg	ccaacgtctt	catgctggcc	tggtccctacg	gcagcccgga	cgtgcactcc	1020
ggctacgagt	tcaccaactt	cgacgccggg	ccgcccacaa	acggcaacgt	gaccgcctgc	1080
caccagagcg	ggtggaagtg	ccagcacgcc	tggtcaggaga	tctcctccat	ggtgggcttc	1140
cgcaacgcc	cccgcggcca	ggcgtcacc	aactgggtgt	ccaacggcaa	caacgccatc	1200
gccttcggcc	ggggcaaccg	cggctacgtg	gcgatcaacc	acgagaacac	cacactgaac	1260
cggaccttcc	agacctccct	gccggccggg	gactactgca	acgtgcagaa	cggcaccacc	1320
gtcaccgtca	acggcgccgg	gcagttcacc	gcgagcctag	gcgcccgcac	cgccctcgcc	1380
ctgcacgtcg	acgcccgcaa	ttgcgccggc	ggcggcaccg	ggggcaacgg	cggcaccggg	1440
ggaggcgaga	acccggtgac	cggcggtgcc	tccttcggcg	tcgacgcgac	cacgcagatg	1500
ggccagaaca	tccacgtcgc	gggcaacatc	cccgcgctgg	gcgactggaa	caccgccaac	1560
gccccccgca	tgagcgccga	cacctaccgc	gtctggcggc	tggaactgaa	cctgccgccc	1620
ggcaccacct	tccagtacaa	gtacatccgc	agggacgcca	acggcaacgt	gacctgggaa	1680
tccgggaaca	accgcaccgc	caccgtgccc	tcctccggcc	gggtgacgct	caacgacacc	1740
tggtcgcaact	ga					1752

<210> 616
 <211> 583
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(34)

<221> DOMAIN
 <222> (390)...(466)
 <223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN
 <222> (487)...(579)
 <223> Starch binding domain

<221> DOMAIN
 <222> (41)...(381)
 <223> Alpha amylase, catalytic domain

<400> 616
 Met Ser Glu Arg Gly Val Arg Arg Ala Val Arg Thr Ala Leu Val Gly
 1 5 10 15
 Leu Ala Ala Ala Ala Thr Ala Ala Val Thr Leu Gly Ala Pro Thr Ala
 20 25 30
 Gln Ala Ala Pro Ala Gly Asn Lys Asp Val Thr Ala Val Leu Phe Ser
 35 40 45
 Trp Asn Phe Asn Ser Ile Ala Arg Glu Cys Thr Glu Arg Leu Gly Pro
 50 55 60
 Ala Gly Tyr Gly Phe Val Gln Val Ser Pro Pro Gln Glu His Ile Gln
 65 70 75 80
 Gly Ser Ala Trp Trp Thr Gln Tyr Gln Pro Val Ser Tyr Asn Ile Ser
 85 90 95
 Gly Arg Leu Gly Asn Glu Gln Gln Phe Arg Ser Met Val Asn Thr Cys
 100 105 110
 Asn Asn Ala Gly Val Gly Val Ile Val Asp Ala Val Ile Asn His Met
 115 120 125
 Ser Ala Gly Ser Gly Thr Gly Thr Gly Gly Thr Ser Tyr Ser Lys Tyr
 130 135 140
 Asn Tyr Pro Gly Leu Tyr Ser Gly Asn Asp Phe Gly Ser Cys Arg Glu
 145 150 155 160

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Asp Ile Arg Gly Ser Asp Tyr Thr Asn Asp Arg Trp Arg Val Gln Asn
 165 170 175
 Cys Glu Leu Val Gly Leu Ala Asp Leu Asp Thr Gly Ser Ala Tyr Val
 180 185 190
 Gln Gln Gln Ile Ala Gly Tyr Met Asn Arg Leu Leu Gly Trp Gly Val
 195 200 205
 Ala Gly Phe Arg Val Asp Ala Val Lys His Ile Pro Ala Arg His Met
 210 215 220
 Glu Gln Ile Arg Ala Arg Val Asn Gly Gly Asp Val Phe Trp Asn Gln
 225 230 235 240
 Glu Val Ile Tyr Gly Ala Gly Glu Ala Ile Thr Pro Glu Glu Tyr Leu
 245 250 255
 Asn Thr Gly His Val Gln Glu Phe Arg Tyr Ala Phe Asp Leu Lys Arg
 260 265 270
 Met Phe Gln Gly Asp Arg Ile Ala Asn Leu Gln Asn Phe Gly Glu Ser
 275 280 285
 Trp Gly Tyr Met Pro Ser Asn Arg Ser Gly Val Phe Val Asp Asn His
 290 295 300
 Asp Thr Glu Arg Asn Gly Ser Thr Leu Ser Tyr Lys Asp Asn Ala Ala
 305 310 315 320
 Tyr Thr Leu Ala Asn Val Phe Met Leu Ala Trp Pro Tyr Gly Ser Pro
 325 330 335
 Asp Val His Ser Gly Tyr Glu Phe Thr Asn Phe Asp Ala Gly Pro Pro
 340 345 350
 Asn Asn Gly Asn Val Thr Ala Cys His Gln Ser Gly Trp Lys Cys Gln
 355 360 365
 His Ala Trp Gln Glu Ile Ser Ser Met Val Gly Phe Arg Asn Ala Thr
 370 375 380
 Arg Gly Gln Ala Val Thr Asn Trp Trp Ser Asn Gly Asn Asn Ala Ile
 385 390 395 400
 Ala Phe Gly Arg Gly Asn Arg Gly Tyr Val Ala Ile Asn His Glu Asn
 405 410 415
 Thr Thr Leu Asn Arg Thr Phe Gln Thr Ser Leu Pro Ala Gly Asp Tyr
 420 425 430
 Cys Asn Val Gln Asn Gly Thr Thr Val Thr Val Asn Gly Ala Gly Gln
 435 440 445
 Phe Thr Ala Ser Leu Gly Ala Arg Thr Ala Leu Ala Leu His Val Asp
 450 455 460
 Ala Arg Asn Cys Ala Gly Gly Gly Thr Gly Gly Asn Gly Gly Thr Gly
 465 470 475 480
 Gly Gly Glu Asn Pro Val Thr Gly Gly Ala Ser Phe Gly Val Asp Ala
 485 490 495
 Thr Thr Gln Met Gly Gln Asn Ile His Val Ala Gly Asn Ile Pro Ala
 500 505 510
 Leu Gly Asp Trp Asn Thr Ala Asn Ala Pro Arg Met Ser Ala Asp Thr
 515 520 525
 Tyr Pro Val Trp Arg Leu Glu Leu Asn Leu Pro Pro Gly Thr Thr Phe
 530 535 540
 Gln Tyr Lys Tyr Ile Arg Asp Ala Asn Gly Asn Val Thr Trp Glu
 545 550 555 560
 Ser Gly Asn Asn Arg Thr Ala Thr Val Pro Ser Ser Gly Arg Val Thr
 565 570 575
 Leu Asn Asp Thr Trp Arg Asn
 580

<210> 617
 <211> 1668
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

<400> 617
 atgaaccgat acctgcgact ggccgcgttg acgctggccc tggcgccgct ggcctacccc 60
 tggggtaatc tggcccgcgc cgccgatgca ccgggcaaga ccgccagcgg cgtgcgtac 120
 cacggcggcg acgaaatcat cctccagggc ttctactgga acaccgtgcg tacctcgagc 180
 aactggtacg cgacgctggc cagcatggcg ccgaccctgg ccgccgatgg tttcagcgca 240

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atctggatgc cgggtgccctg ggcgcgacttc tccagctgga ggcaccccg caacggcacc 300
tcggggcggcg gcgaaggcta ctcttgacac gacttcaaca agaacggccg ctatggcagc 360
gacagcctgc tcaaacaggc cgccggcgcg ctcaatgcgg ccgggggtcaa acccatctac 420
gacgtggtgc ccaaccacat gaaccgcggc taccgggaca aggaaatcaa cctgccggcc 480
ggccaggggc tatggcgcca cgactgcaat gacccgggca actacgccaa cgactgcgac 540
gacggcgacc gcttcatggg cggcgacgcc gacctcaata ccggccaccc gcagaactac 600
gcatgttcc gcgacgagtt cgccaggctg cgcagccagt atggcgccgg gggctttcgc 660
ttcgacttcg tccggcgcta tgccggcgag cgggtcgcca actggatgag cgatgcccac 720
gacaacggct tctgcctcgg cgagctgtgg aaggcgccgg gcgaataccc aagctgggac 780
tggcgcaacg gcgccagctg gcagcagatc ctcaaggact ggtccgatcg cgccaagtgc 840
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cacggcctca acggcaaccc cgatgcgcgc tggcgcgagg tggcggtgac cttcgtcgac 960
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cgccgtgcgg cgggggtcaa ggccggcctc gcgatccagt tccactcggg tttctccgga 1200
ctggtggcga cagtcagcgg cagccagcag caactgtcta tcgccctgga ctccaacctg 1260
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gcatccgta tctggcgag cggtcagagt gacgatgagg agcagggcga tctggtcagc 1380
gtcaacttcc gctgtgacaa cggcgtagcc cagtggggcg acagcgtcta cgcggtgggc 1440
aacgtcgccc aactgggcaa ctggagcccc gccagcgccg tgcgcctgac cgataccagc 1500
gcctaccgga cctggaaagg cagcatcgcc ctgccggccg gccagcaggt gcagtggaaa 1560
tgcacgtgc gcagcgaag caaccgcacc caggtcaaga cctggcagcc gggcggaac 1620
aacgccgtga cggtgacggc aggggccagc accgcgggga gtttctaa 1668

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<210> 618
 <211> 555
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(27)

<221> DOMAIN
 <222> (458)...(554)
 <223> Starch binding domain

<221> DOMAIN
 <222> (45)...(395)
 <223> Alpha amylase, catalytic domain

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<400> 618
Met Asn Arg Tyr Leu Arg Leu Ala Ala Leu Thr Leu Ala Leu Ala Pro
 1          5          10          15
Leu Ala Tyr Pro Trp Gly Asn Leu Ala Arg Ala Ala Asp Ala Pro Gly
          20          25          30
Lys Thr Ala Ser Gly Val Arg Tyr His Gly Gly Asp Glu Ile Ile Leu
          35          40          45
Gln Gly Phe His Trp Asn Thr Val Arg Thr Ser Ser Asn Trp Tyr Ala
          50          55          60
Thr Leu Ala Ser Met Ala Pro Thr Leu Ala Ala Asp Gly Phe Ser Ala
          65          70          75          80
Ile Trp Met Pro Val Pro Trp Arg Asp Phe Ser Ser Trp Ser Asp Pro
          85          90          95
Gly Asn Gly Thr Ser Gly Gly Gly Glu Gly Tyr Phe Trp His Asp Phe
          100          105          110
Asn Lys Asn Gly Arg Tyr Gly Ser Asp Ser Leu Leu Lys Gln Ala Ala
          115          120          125
Gly Ala Leu Asn Ala Ala Gly Val Lys Pro Ile Tyr Asp Val Val Pro
          130          135          140
Asn His Met Asn Arg Gly Tyr Pro Asp Lys Glu Ile Asn Leu Pro Ala
          145          150          155          160
Gly Gln Gly Leu Trp Arg His Asp Cys Asn Asp Pro Gly Asn Tyr Ala
          165          170          175
Asn Asp Cys Asp Asp Gly Asp Arg Phe Met Gly Gly Asp Ala Asp Leu

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Asn	Thr	Gly	180	His	Pro	Gln	Asn	Tyr	185	Ala	Met	Phe	Arg	Asp	190	Glu	Phe	Ala
Arg	Leu	Arg	195	Ser	Gln	Tyr	Gly	200	Ala	Gly	Gly	Phe	Arg	Phe	205	Asp	Phe	Val
Arg	Gly	Tyr	210	Ala	Gly	Glu	Arg	215	Val	Ala	Asn	Trp	Met	Ser	220	Asp	Ala	His
Asp	Asn	Gly	225	Phe	Cys	Leu	Gly	230	Glu	Leu	Trp	Lys	Ala	Pro	235	Gly	Glu	Tyr
Pro	Ser	Trp	245	Trp	Arg	Asn	Gly	250	Ala	Ser	Trp	Gln	Gln	Ile	255	Leu	Lys	
Asp	Trp	Ser	260	Asp	Arg	Ala	Lys	265	Thr	Val	Phe	Asp	Phe	270	Ala	Leu	Lys	
Glu	Arg	Met	275	Gln	Asn	Gly	Gly	280	Ile	Ala	Asp	Trp	Arg	285	His	Gly	Leu	Asn
Gly	Asn	Pro	290	Asp	Ala	Arg	Trp	295	Arg	Glu	Val	Ala	Val	300	Thr	Phe	Val	Asp
Asn	His	Asp	305	Thr	Gly	Tyr	Ser	310	Pro	Gly	Pro	His	Gly	315	Gly	Gln	His	His
Trp	Pro	Leu	325	Asp	Ala	Arg	Leu	330	Gln	Ala	Tyr	Ala	Tyr	335	Ile	Leu		
Ser	Ser	Pro	340	Gly	Thr	Pro	Val	345	Tyr	Trp	Pro	His	Met	350	Tyr	Asp	Trp	
Gly	His	Gly	355	Asp	Phe	Ile	Arg	360	Gln	Leu	Ile	Gln	Ile	365	Arg	Arg	Ala	Ala
Gly	Val	Lys	370	Ala	Ala	Ser	375	Ala	Ile	Gln	Phe	His	Ser	380	Gly	Phe	Ser	Gly
Leu	Val	Ala	385	Thr	Val	Ser	Gly	390	Ser	Gln	Gln	Gln	Leu	395	Leu	Ile	Ala	Leu
Asp	Ser	Asn	405	Leu	Ser	Thr	Pro	410	Ser	Gln	Val	Ala	Ser	415	Gly	Asp	Phe	Thr
Gln	Ala	Leu	420	Asn	Thr	Asp	Asn	425	Ala	Ile	Arg	Ile	Trp	430	Arg	Ser	Gly	
Gln	Ser	Asp	435	Asp	Glu	Glu	Gln	440	Gly	Asp	Leu	Val	Ser	445	Val	Asn	Phe	Arg
Cys	Asp	Asn	450	Gly	Val	Thr	Gln	455	Trp	Gly	Asp	Ser	Val	460	Tyr	Ala	Val	Gly
Asn	Val	Ala	465	Gln	Leu	Gly	Asn	470	Trp	Ser	Pro	Ala	Ser	475	Ala	Val	Arg	Leu
Thr	Asp	Thr	485	Ala	Tyr	Pro	Thr	490	Trp	Lys	Gly	Ser	Ile	495	Ala	Leu	Pro	
Ala	Gly	Gln	500	Val	Gln	Trp	Lys	505	Cys	Ile	Val	Arg	Ser	510	Glu	Ser	Asn	
Pro	Thr	Gln	515	Val	Lys	Thr	Trp	520	Gln	Pro	Gly	Gly	Asn	525	Asn	Ala	Val	Thr
Val	Thr	Ala	530	Gly	Ala	Ser	Thr	535	Ala	Gly	Ser	Phe		540				
545						550						555						

<210> 619
 <211> 1716
 <212> DNA
 <213> Bacteria

<400> 619																		
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atgttcgaat	ggaacttcgc	ctccgtcgcc	cgcgagtgca	ccgaccgcct	cggccccgcc													180
ggatacggat	acgtccaggt	ctccccgccc	caggaacacc	tccagggcgg	ccagtgggtg													240
acctcctacc	agccggtcag	ctacaagatc	gccggacgcc	tcggtgaccg	caccgccttc													300
aagaagatga	tcgacacctg	ccacgcggcc	ggcgtcaagg	tcgtcgccga	ctccgtcatc													360
aaccacatgg	cgaacggctc	cggcacggga	acgggcggga	ccccgttctc	caagtacgac													420
tacccggggc	tctactccgg	cgccgacatg	gacgactgcc	gcgcgacgat	ctccacctac													480
caggaccggg	gcaacgtcca	gaactgcgaa	ctggtccagc	tccccgacct	cgacaccggc													540
gaggaccacg	tccgcgggaa	gatagccggc	tacctcaacg	acctgctctc	cctcggcgtc													600
gacggcttcc	gcgtcgacgc	cgccaagcac	atgccggccg	ccgacctcgc	cgccatcaag													660
gcgcggctca	gcaacccccg	cgcgctactg	aagcaggagg	ccatcttcgg	cgcgggcgag													720
gccgtctccc	cgagcgagta	cctcggcaac	ggcgacgtcc	aggagttccg	ctacgcctgc													780
gacctcaagc	gcgtcctgca	gaacgagaag	ctcgcctacc	tcaagaactt	cggcgaggcc													840

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```

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tacgtggcca tcaaccacga gacctcggcg ctcaccgcga cctaccagac ctcgctgccc 1260
gccggcacct actgcgacgt ccagtccaac acccccgtga cggatgaacgg ctccggccag 1320
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aacgccaccg ccaccaccgt ggtcggtcag gacatctacg tcaccggcaa ccgcgccgag 1500
ctcggcaact ggtccccggc cgccgccctc aagctcgacc ccgcgacgta ccccgctctg 1560
aagctgaccg tcggcctgcc cgccggaacc gccttcgagt acaagtacct ccgcaaggac 1620
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<210> 620
 <211> 571
 <212> PRT
 <213> Bacteria

<220>
 <221> SIGNAL
 <222> (1)...(29)

<221> DOMAIN
 <222> (475)...(567)
 <223> Starch binding domain

<221> DOMAIN
 <222> (382)...(458)
 <223> Alpha amylase, C-terminal all-beta domain

<221> DOMAIN
 <222> (36)...(373)
 <223> Alpha amylase, catalytic domain

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 20          25          30
Gly Glu Lys Asp Val Thr Ala Val Met Phe Glu Trp Asn Phe Ala Ser
 35          40          45
Val Ala Arg Glu Cys Thr Asp Arg Leu Gly Pro Ala Gly Tyr Gly Tyr
 50          55          60
Val Gln Val Ser Pro Pro Gln Glu His Leu Gln Gly Gly Gln Trp Trp
 65          70          75          80
Thr Ser Tyr Gln Pro Val Ser Tyr Lys Ile Ala Gly Arg Leu Gly Asp
 85          90          95
Arg Thr Ala Phe Lys Lys Met Ile Asp Thr Cys His Ala Ala Gly Val
100          105          110
Lys Val Val Ala Asp Ser Val Ile Asn His Met Ala Asn Gly Ser Gly
115          120          125
Thr Gly Thr Gly Gly Thr Pro Phe Ser Lys Tyr Asp Tyr Pro Gly Leu
130          135          140
Tyr Ser Gly Ala Asp Met Asp Asp Cys Arg Ala Thr Ile Ser Thr Tyr
145          150          155          160
Gln Asp Arg Gly Asn Val Gln Asn Cys Glu Leu Val Gln Leu Pro Asp
165          170          175
Leu Asp Thr Gly Glu Asp His Val Arg Gly Lys Ile Ala Gly Tyr Leu
180          185          190
Asn Asp Leu Leu Ser Leu Gly Val Asp Gly Phe Arg Val Asp Ala Ala
195          200          205
Lys His Met Pro Ala Ala Asp Leu Ala Ala Ile Lys Ala Arg Leu Ser
210          215          220
Asn Pro Gly Ala Tyr Trp Lys Gln Glu Ala Ile Phe Gly Ala Gly Glu
225          230          235          240

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Ala	Val	Ser	Pro	Ser	Glu	Tyr	Leu	Gly	Asn	Gly	Asp	Val	Gln	Glu	Phe
Arg	Tyr	Ala	Arg	Asp	Leu	Lys	Arg	Val	Leu	Gln	Asn	Glu	Lys	Leu	Ala
Tyr	Leu	Lys	Asn	Phe	Gly	Glu	Ala	Trp	Gly	Tyr	Met	Pro	Ser	Ala	Gln
Ser	Gly	Val	Phe	Val	Asp	Asn	His	Asp	Thr	Glu	Arg	Gly	Gly	Asp	Thr
Leu	Ser	Tyr	Lys	Asp	Gly	Ala	Asn	Tyr	Thr	Leu	Ala	Ser	Val	Phe	Met
Leu	Ala	Trp	Pro	Tyr	Gly	Ser	Pro	Asp	Val	His	Ser	Gly	Tyr	Glu	Trp
Thr	Asp	Lys	Asp	Ala	Gly	Pro	Pro	Ser	Gly	Gly	Gln	Val	Asn	Ala	Cys
Tyr	Thr	Asp	Gly	Trp	Lys	Cys	Gln	His	Ala	Trp	Arg	Glu	Ile	Ser	Ser
Met	Val	Ala	Phe	Arg	Asn	Thr	Ala	Arg	Gly	Gln	Ala	Val	Thr	Gly	Trp
Trp	Asp	Asn	Gly	Asn	Asn	Ala	Ile	Ala	Phe	Gly	Arg	Gly	Ser	Lys	Ala
Tyr	Val	Ala	Ile	Asn	His	Glu	Thr	Ser	Ala	Leu	Thr	Arg	Thr	Tyr	Gln
Thr	Ser	Leu	Pro	Ala	Gly	Thr	Tyr	Cys	Asp	Val	Gln	Ser	Asn	Thr	Pro
Val	Thr	Val	Asn	Gly	Ser	Gly	Gln	Phe	Thr	Ala	Thr	Leu	Gly	Ala	His
Thr	Ala	Leu	Ala	Leu	His	Val	Gly	Ala	Met	Asn	Cys	Gly	Ser	Gly	Thr
Thr	Thr	Pro	Pro	Pro	Thr	Thr	Pro	Pro	Ala	Thr	Pro	Gly	Ala	Ser	Phe
Asn	Ala	Thr	Ala	Thr	Thr	Val	Val	Gly	Gln	Asp	Ile	Tyr	Val	Thr	Gly
Asn	Arg	Ala	Glu	Leu	Gly	Asn	Trp	Ser	Pro	Ala	Ala	Ala	Leu	Lys	Leu
Asp	Pro	Ala	Thr	Tyr	Pro	Val	Trp	Lys	Leu	Thr	Val	Gly	Leu	Pro	Ala
Gly	Thr	Ala	Phe	Glu	Tyr	Lys	Tyr	Leu	Arg	Lys	Asp	Ala	Ala	Gly	Asn
Val	Thr	Trp	Glu	Ser	Gly	Ala	Asn	Arg	Thr	Ala	Thr	Val	Pro	Ala	Ser
Gly	Arg	Ile	Val	Leu	Asn	Asp	Thr	Phe	Arg	Ser					

<210> 621
 <211> 1770
 <212> DNA
 <213> Unknown

<220>
 <223> obtained from an environmental sample

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caggtgaaca	acaccatgta ccaggcattt tattgggatg cctaccctgg cctttgggcc 180
aatttaccgg	ccatggcggc ccctttggcc gaacggggca ttacctcgat gtggttgccg 240
cccgcgccca	aaggcatgaa tgggtactttc agtgtcgggtt acgatgtata cgatttctgg 300
gatctgggcg	agttaaacca aaaaggcacc accgccacc gttacggtac tcgtcagcag 360
ctgcaacaag	cactgagtgc tctggaccaaa ctgggtattc aggcctattt tgatgtggtg 420
tttaaccacc	gcatgggcgc cgaatgcacag gagaatattc ctggcctttgg cctggcctgg 480
accgagtatc	atctgcaagg tcgtcaggcg cattataccc agcaaaactg gggctacttg 540
tggcagcact	ttgactggaa ctggaccgcg tttaatggct cgcacaatca gctctacccc 600
ggcaaatggt	ggggcaatac cttccacttc ccttatttga tgggtgagga tgtcgattac 660
aaccgctttg	aagtgcagca ggaaatgaaa gcctggggcg agtggatcat caacagcggt 720
ggcttttagcg	gctttcggat ggacgccatc gcacacgtgg ataccgattt taccctgtgac 780
tggatcaatc	acgtacagtg ggccaccagc gaggatgtat tctttgtcgc agaagcctgg 840
gtcagtgata	tcaacggcta tctggatgca gtcaatacgc cgcattttgc cgcttttgat 900
ttcaatttgc	gcgaagactt tgttgcttta agcagtgggtg gcaaagacat gcgctggtgg 960

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ggtggtctgg ttaatagcca gcaccgtgat cgggcggtca cttttgtcga taaccacgat 1020
accagccggg ccggcaaccc ttatggcatg ccgcagggtga tcaactacaa gaaccaggcc 1080
tacgcttaca ttctgttgcg tgagcatggg gtgccgactg tgtttgccg cgattacgac 1140
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agcagtgtag agctcaccaa ttggggcggc ggtattgaag gcacctggac cagtggccat 1620
gtctgggaag tgaccatccc ggatccgggc aactttgaat ggaaaaccg taaaggccca 1680
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cccagtttta atggtggggt tagatctcgc 1770

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<210> 622
 <211> 590
 <212> PRT
 <213> Unknown

<220>
 <223> Obtained from an environmental sample

<221> SIGNAL
 <222> (1)...(31)

<221> DOMAIN
 <222> (44)...(395)
 <223> Alpha amylase, catalytic domain

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Ala Ile Ala Leu Ser Ala Leu Ser Leu Ser Phe Gly Leu Gln Ala Ser
          20           25           30
Glu Leu Pro Gln Ile Pro Pro Gln Gln Val Asn Asn Thr Met Tyr Gln
          35           40           45
Ala phe Tyr Trp Asp Ala Tyr Pro Gly Leu Trp Ala Asn Leu Pro Ala
          50           55           60
Met Ala Ala Pro Leu Ala Glu Arg Gly Ile Thr Ser Met Trp Leu Pro
65          70          75          80
Pro Ala Ala Lys Gly Met Asn Gly Thr Phe Ser Val Gly Tyr Asp Val
          85           90           95
Tyr Asp Phe Trp Asp Leu Gly Glu Phe Asn Gln Lys Gly Thr Thr Ala
          100          105          110
Thr Arg Tyr Gly Thr Arg Gln Gln Leu Gln Gln Ala Leu Ser Ala Leu
          115          120          125
Asp Gln Leu Gly Ile Gln Ala Tyr Phe Asp Val Val Phe Asn His Arg
          130          135          140
Met Gly Ala Asp Ala Gln Glu Asn Ile Pro Gly Phe Gly Leu Ala Trp
145          150          155          160
Thr Glu Tyr His Leu Gln Gly Arg Gln Ala His Tyr Thr Gln Gln Asn
          165          170          175
Trp Gly Tyr Leu Trp His Asp Phe Asp Trp Asn Trp Thr Ala Phe Asn
          180          185          190
Gly Ser Asp Asn Gln Leu Tyr Pro Gly Lys Trp Trp Gly Asn Thr Phe
          195          200          205
His phe Pro Tyr Leu Met Gly Glu Asp Val Asp Tyr Asn Arg Phe Glu
          210          215          220
Val Gln Gln Glu Met Lys Ala Trp Gly Glu Trp Ile Ile Asn Ser Val
225          230          235          240
Gly Phe Ser Gly Phe Arg Met Asp Ala Ile Ala His Val Asp Thr Asp
          245          250          255
phe Thr Arg Asp Trp Ile Asn His Val Gln Trp Ala Thr Ser Glu Asp
          260          265          270
Val phe phe Val Ala Glu Ala Trp Val Ser Asp Ile Asn Gly Tyr Leu
          275          280          285
Asp Ala Val Asn Thr Pro His Leu Arg Ala Phe Asp Phe Asn Leu Arg

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300

	290					295					300				
Glu 305	Asp	Phe	Val	Ala	Leu 310	Ser	Ser	Gly	Gly	Lys 315	Asp	Met	Arg	Trp	Trp 320
Gly	Gly	Leu	Val	Asn 325	Ser	Gln	His	Arg	Asp 330	Arg	Ala	Val	Thr	Phe 335	Val
Asp	Asn	His	Asp 340	Thr	Ser	Arg	Ala	Gly 345	Asn	Pro	Tyr	Gly	Met 350	Pro	Gln
Val	Ile	Asn 355	Tyr	Lys	Asn	Gln	Ala 360	Tyr	Ala	Tyr	Ile	Leu 365	Leu	Arg	Glu
His	Gly 370	Val	Pro	Thr	Val	Phe 375	Ala	Arg	Asp	Tyr	Asp 380	Glu	Phe	Gly	Met
Ala 385	Pro	Thr	Leu	Asp	Lys 390	Leu	Ile	Glu	Ala	Arg 395	Arg	Tyr	Phe	Ala	Tyr 400
Gly	Pro	Gly	His	Glu 405	Tyr	Ser	Gly	Asn	Thr 410	Glu	Ala	Val	Tyr	Ala 415	Tyr
Val	Arg	Glu	Gly 420	Leu	Ser	Thr	Val	Pro 425	Gly	Thr	Gly	Leu	Val 430	Met	Leu
Ile	Ser	Gly 435	Arg	Asn	Trp	Gly	Gly 440	Gln	Ser	Phe	Thr 445	Ile	Asn	Ser	
His	Gln 450	Pro	Asn	Thr	Thr	Phe 455	Tyr	Asp	Tyr	Thr	Gly 460	Asn	Val	Ser	Gly
Thr 465	Val	Thr	Thr	Asn	Ala 470	Gln	Gly	Tyr	Gly	Ser 475	Phe	Pro	Val	Thr	Met 480
Thr	Glu	Ser	Thr	Gly 485	Trp	Ser	Val	Trp	Val 490	Pro	Gln	Ser	Ser	Gly 495	Gly
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Gly	Phe	Ser 515	Leu	Phe	Phe	Thr	Gly 520	Ser	Ser	Ala	Glu	Leu 525	Thr	Asn	Trp
Gly	Gly 530	Gly	Ile	Glu	Gly	Thr 535	Trp	Thr	Ser	Gly	His 540	Val	Trp	Glu	Val
Thr 545	Ile	Pro	Asp	Pro	Gly 550	Asn	Phe	Glu	Trp	Lys 555	Thr	Arg	Lys	Gly	Pro 560
Thr	Gly	Gly	Ser	Gly 565	Gln	Asp	Trp	Glu	Ser 570	Gly	Ser	Asn	His	Asn 575	Gln
Thr	Asn	Leu	His 580	Pro	Ser	Phe	Asn	Gly 585	Phe	Arg	Ser	Arg 590			