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# DESCRIPTION

## FIELD OF THE INVENTION

[0001] The present invention relates to lipase variants.

## BACKGROUND OF THE INVENTION

[0002] Lipases are useful, e.g., as detergent enzymes to remove lipid or fatty stains from clothes and other textiles, as additives to dough for bread and other baked products. Thus, a lipase derived from *Thermomyces lanuginosus* (synonym *Humicola lanuginosa*, EP 258 068 and EP 305 216) is sold for detergent use under the trade name Lipolase<sup>®</sup> (product of Novo Nordisk A/S). WO 0060063 describes variants of the *T. lanuginosus* lipase with a particularly good first-wash performance in a detergent solution. WO 9704079, WO 9707202 and WO 0032758 also disclose variants of the *T. lanuginosus* lipase.

[0003] In some applications, it is of interest to minimize the formation of odor-generating short-chain fatty acids. Thus, it is known that laundry detergents with lipases may sometimes leave residual odors attached to cloth soiled with milk (EP 430315). WO 02062973 discloses lipase variants where the odor generation has been reduced by attaching a C-terminal extension.

## SUMMARY OF THE INVENTION

[0004] The inventors have found that by introducing mutations in certain regions/positions of a lipase it is possible to improve the properties or characteristics of the lipase.

[0005] In a preferred embodiment, the present invention relates to lipases having improved properties for use in detergents. For example, the invention provides variants with reduced tendency to odor generation obtained by introducing mutations in one or more regions identified in the parent lipase. In another preferred embodiment, the present invention provides lipase variants which, as compared to the parent lipase, have reduced potential for odor generation without the attachment of a C-terminal extension.

[0006] In a further aspect the invention relates to a DNA sequence encoding the lipase variant of the invention, an expression vector harbouring said DNA sequence and a transformed host cell containing the DNA sequence or the expression vector.

[0007] In another aspect, the invention provides a method of producing the lipase variant of

the present invention.

## **BRIEF DESCRIPTION OF THE FIGURES**

**[0008]** Figure 1 shows the alignment of lipases.

## **SEQUENCE LISTINGS**

**[0009]**

SEQ ID NO: 1 shows the DNA sequence encoding lipase from *Thermomyces lanoginosus*.

SEQ ID NO: 2 shows the amino acid sequence of a lipase from *Thermomyces lanoginosus*.

SEQ ID NO: 3 shows the amino acid sequence of a lipase from *Absidia reflexa*.

SEQ ID NO: 4 shows the amino acid sequence of a lipase from *Absidia corymbifera*.

SEQ ID NO: 5 shows the amino acid sequence of a lipase from *Rhizomucor miehei*.

SEQ ID NO: 6 shows the amino acid sequence of a lipase from *Rhizopus oryzae*.

SEQ ID NO: 7 shows the amino acid sequence of a lipase from *Aspergillus niger*.

SEQ ID NO: 8 shows the amino acid sequence of a lipase from *Aspergillus tubingensis*.

SEQ ID NO: 9 shows the amino acid sequence of a lipase from *Fusarium oxysporum*.

SEQ ID NO: 10 shows the amino acid sequence of a lipase from *Fusarium heterosporum*.

SEQ ID NO: 11 shows the amino acid sequence of a lipase from *Aspergillus oryzae*.

SEQ ID NO: 12 shows the amino acid sequence of a lipase from *Penicillium camemberti*.

SEQ ID NO: 13 shows the amino acid sequence of a lipase from *Aspergillus foetidus*.

SEQ ID NO: 14 shows the amino acid sequence of a lipase from *Aspergillus niger*.

SEQ ID NO: 15 shows the amino acid sequence of a lipase from *Aspergillus oryzae*.

SEQ ID NO: 16 shows the amino acid sequence of a lipase from *Landerina penisapora*.

## **DETAILED DESCRIPTION OF THE INVENTION**

**Parent lipases**

**[0010]** Any suitable parent lipase may be used. In a preferred embodiment, the parent lipase may be a fungal lipase. In another preferred embodiment, the parent lipase may be a lipase with an amino acid sequence having at least 50%, 60%, 70%, 80%, 85%, 90%, 95%, 96%, 97%, 98%, 99% or even 100% homology as defined in the section "Homology and alignment" to the sequence of the *T. lanuginosus* lipase shown in SEQ ID NO: 2.

**[0011]** The parent lipase may be a yeast polypeptide such as a *Candida*, *Kluyveromyces*, *Pichia*, *Saccharomyces*, *Schizosaccharomyces*, or *Yarrowia* polypeptide; or more preferably a filamentous fungal polypeptide such as an *Acremonium*, *Aspergillus*, *Aureobasidium*, *Cryptococcus*, *Filobasidium*, *Fusarium*, *Humicola*, *Magnaporthe*, *Mucor*, *Myceliophthora*, *Neocallimastix*, *Neurospora*, *Paecilomyces*, *Penicillium*, *Piromyces*, *Schizophyllum*, *Talaromyces*, *Thermoascus*, *Thielavia*, *Tolypocladium*, or *Trichoderma* polypeptide.

**[0012]** In a preferred aspect, the parent lipase is a *Saccharomyces carlsbergensis*, *Saccharomyces cerevisiae*, *Saccharomyces diastaticus*, *Saccharomyces douglasii*, *Saccharomyces kluyveri*, *Saccharomyces norbensis*, or *Saccharomyces oviformis* polypeptide having lipase activity.

**[0013]** In another preferred aspect, the parent lipase is an *Aspergillus aculeatus*, *Aspergillus awamori*, *Aspergillus fumigatus*, *Aspergillus foetidus*, *Aspergillus japonicus*, *Aspergillus nidulans*, *Aspergillus niger*, *Aspergillus oryzae*, *Aspergillus turbigensis*, *Fusarium bactridioides*, *Fusarium cerealis*, *Fusarium crookwellense*, *Fusarium culmorum*, *Fusarium graminearum*, *Fusarium graminum*, *Fusarium heterosporum*, *Fusarium negundi*, *Fusarium oxysporum*, *Fusarium reticulatum*, *Fusarium roseum*, *Fusarium sambucinum*, *Fusarium sarcochroum*, *Fusarium sporotrichioides*, *Fusarium sulphureum*, *Fusarium torulosum*, *Fusarium trichothecioides*, *Fusarium venenatum*, *Humicola insolens*, *Thermomyces lanuginosus* (synonym: *Humicola lanuginosa*), *Mucor miehei*, *Myceliophthora thermophila*, *Neurospora crassa*, *Penicillium purpurogenum*, *Trichoderma harzianum*, *Trichoderma koningii*, *Trichoderma longibrachiatum*, *Trichoderma reesei*, or *Trichoderma viride* polypeptide.

**[0014]** In another preferred aspect, the parent lipase is a *Thermomyces* lipase.

**[0015]** In a more preferred aspect, the parent lipase is a *Thermomyces lanuginosus* lipase. In an even more preferred embodiment the parent lipase is the lipase of SEQ ID NO: 2.

**Variant lipases**

**[0016]** The lipase variants of the present invention is at least 80% identical to SEQ ID NO:2 and comprise, as compared to the parent lipase, at least three substitutions selected from the

group consisting of:

1. a) at least two substitution in Region I, and
2. b) at least one substitution in Region II, and
3. c) at least one substitution in Region III, and
4. d) at least one substitution in Region IV,

and wherein the variant has lipase activity and exhibits improved wash performance and odor and comprises the substitutions T231 R +N233R +I255Y.

**[0017]** In a preferred embodiment the variant lipase is a variant of a *Thermomyces* lipase, more preferably, a *T. lanuginosus* lipase, and even more preferably, the *T. lanuginosus* lipase shown in SEQ ID NO: 2. In a preferred embodiment. The variant lipase has at least 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to SEQ ID NO:2. The variant lipase may be a variant of a parent lipase encoded by a gene derived/obtained from one of the following parent organisms: *Candida*, *Kluyveromyces*, *Pichia*, *Saccharomyces*, *Schizosaccharomyces*, or *Yarrowia*, *Acremonium*, *Aspergillus*, *Aureobasidium*, *Cryptococcus*, *Filobasidium*, *Fusarium*, *Humicola*, *Magnaporthe*, *Mucor*, *Myceliophthora*, *Neocallimastix*, *Neurospora*, *Paecilomyces*, *Penicillium*, *Piromyces*, *Schizophyllum*, *Talaromyces*, *Thermoascus*, *Thielavia*, *Tolypocladium*, or *Trichoderma*. In a preferred embodiment, the variant lipase has at least 50%, 60%, 70%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to a parent lipase encoded by a gene derived/obtained from one of the following parent organisms: *Candida*, *Kluyveromyces*, *Pichia*, *Saccharomyces*, *Schizosaccharomyces*, or *Yarrowia*, *Acremonium*, *Aspergillus*, *Aureobasidium*, *Cryptococcus*, *Filobasidium*, *Fusarium*, *Humicola*, *Magnaporthe*, *Mucor*, *Myceliophthora*, *Neocallimastix*, *Neurospora*, *Paecilomyces*, *Penicillium*, *Piromyces*, *Schizophyllum*, *Talaromyces*, *Thermoascus*, *Thielavia*, *Tolypocladium*, or *Trichoderma*.

**[0018]** In a preferred aspect, the variant lipase is a variant *Saccharomyces carlsbergensis*, *Saccharomyces cerevisiae*, *Saccharomyces diastaticus*, *Saccharomyces douglasii*, *Saccharomyces kluyveri*, *Saccharomyces norbensis*, or *Saccharomyces oviformis*. In a preferred embodiment, the variant lipase has at least 50%, 60%, 70%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to a parent lipase encoded by a gene derived/obtained from *Saccharomyces carlsbergensis*, *Saccharomyces cerevisiae*, *Saccharomyces diastaticus*, *Saccharomyces douglasii*, *Saccharomyces kluyveri*, *Saccharomyces norbensis*, or *Saccharomyces oviformis*.

**[0019]** The variant lipase may be a variant of a parent lipase encoded by a gene derived/obtained from one of the following parent organisms: *Aspergillus aculeatus*, *Aspergillus awamori*, *Aspergillus fumigatus*, *Aspergillus foetidus*, *Aspergillus japonicus*, *Aspergillus nidulans*, *Aspergillus niger*, *Aspergillus oryzae*, *Aspergillus turbigensis*, *Fusarium bactridioides*, *Fusarium cerealis*, *Fusarium crookwellense*, *Fusarium culmorum*, *Fusarium graminearum*, *Fusarium graminum*, *Fusarium heterosporum*, *Fusarium negundi*, *Fusarium oxysporum*, *Fusarium reticulatum*, *Fusarium roseum*, *Fusarium sambucinum*, *Fusarium sarcochroum*, *Fusarium sporotrichioides*, *Fusarium sulphureum*, *Fusarium torulosum*, *Fusarium trichothecioides*, *Fusarium venenatum*, *Humicola insolens*, *Thermomyces lanuginosus*

(synonym: *Humicola lanuginosa*), *Mucor miehei*, *Myceliophthora thermophila*, *Neurospora crassa*, *Penicillium purpurogenum*, *Trichoderma harzianum*, *Trichoderma koningii*, *Trichoderma longibrachiatum*, *Trichoderma reesei*, or *Trichoderma viride*. In a preferred embodiment, the variant lipase has at least 50%, 60%, 70%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to a parent lipase encoded by a gene derived/obtained from one of the following parent organisms: *Aspergillus aculeatus*, *Aspergillus awamori*, *Aspergillus fumigatus*, *Aspergillus foetidus*, *Aspergillus japonicus*, *Aspergillus nidulans*, *Aspergillus niger*, *Aspergillus oryzae*, *Aspergillus turbigensis*, *Fusarium bactridioides*, *Fusarium cerealis*, *Fusarium crookwellense*, *Fusarium culmorum*, *Fusarium graminearum*, *Fusarium graminum*, *Fusarium heterosporum*, *Fusarium negundi*, *Fusarium oxysporum*, *Fusarium reticulatum*, *Fusarium roseum*, *Fusarium sambucinum*, *Fusarium sarcochroum*, *Fusarium sporotrichioides*, *Fusarium sulphureum*, *Fusarium torulosum*, *Fusarium trichothecioides*, *Fusarium venenatum*, *Humicola insolens*, *Thermomyces lanuginosus* (synonym: *Humicola lanuginosa*), *Mucor miehei*, *Myceliophthora thermophila*, *Neurospora crassa*, *Penicillium purpurogenum*, *Trichoderma harzianum*, *Trichoderma koningii*, *Trichoderma longibrachiatum*, *Trichoderma reesei*, or *Trichoderma viride*.

**[0020]** In another preferred aspect, the variant is a variant of a *Thermomyces* lipase.

**[0021]** In a more preferred aspect, the parent lipase is a *Thermomyces lanuginosus* lipase. In an even more preferred embodiment the parent lipase is the lipase of SEQ ID NO: 2.

#### **Identification of regions and substitutions**

**[0022]** The positions referred to in Region I through Region IV below are the positions of the amino acid residues in SEQ ID NO:2. To find the corresponding (or homologous) positions in a different lipase, the procedure described in "Homology and alignment" is used.

#### **Substitutions in Region I**

**[0023]** Region I consists of amino acid residues surrounding the N-terminal residue E1. In this region, it is preferred to substitute an amino acid of the parent lipase with a more positive amino acid. The lipase variant comprise at least two substitutions in Region I, such as three, four, five or six substitutions in Region I, and include substitutions T231 R and N233R.

**[0024]** Amino acid residues corresponding to the following positions are comprised by Region I: 1, 2 to 11 and 223-239. The following positions are of particular interest: 1, 4, 8, 11, 223, 227, 229, 231, 233, 234, 236.

**[0025]** In particular the following additional substitutions have been identified: X1N/\* X4V and X227G.

**[0026]** In a preferred embodiment the variant lipase has at least 85%, 90%, 95%, 96%, 97%, 98%, 99% identity to SEQ ID NO:2

**[0027]** In a most preferred embodiment the variant lipase is a variant of the lipase having the amino acid sequence of SEQ ID NO: 2.

#### **Substitutions in Region II**

**[0028]** Region II consists of amino acid residues in contact with substrate on one side of the acyl chain and one side of the alcohol part. In this region it is preferred to substitute an amino acid of the parent lipase with a more positive amino acid or with a less hydrophobic amino acid.

**[0029]** The lipase variant may comprise at least one substitution in Region II, such as two, three, four, five or six substitutions in Region II and include substitution I255Y.

**[0030]** Amino acid residues corresponding to the following positions are comprised by Region II: 202 to 211 and 249 to 269. The following positions are of particular interest : 202, 210, 211, 253, 254, 255, 256.

**[0031]** In particular the following substitutions have been identified: X202G, X210K/W/A, and X256K/R and X259G/M/Q/V.

**[0032]** In a preferred embodiment the variant lipase has at least 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to SEQ ID NO:2

**[0033]** In a most preferred embodiment the variant lipase is a variant of the lipase having the amino acid sequence of SEQ ID NO: 2.

#### **Substitutions in Region III**

**[0034]** Region III consists of amino acid residues that forms a flexible structure and thus allowing the substrate to get into the active site. In this region it is preferred to substitute an amino acid of the parent lipase with a more positive amino acid or a less hydrophobic amino acid.

**[0035]** The lipase variant may comprise at least one substitution in Region III, such as two, three, four, five or six substitutions in Region III.

**[0036]** Amino acid residues corresponding to the following positions are comprised by Region II 82 to 102. The following positions are of particular interest: 83, 86, 87, 90, 91, 95, 96, 99.

**[0037]** In particular the following substitutions have been identified: X83T, X86V and X90A/R.

**[0038]** In a preferred embodiment the variant lipase has at least 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to SEQ ID NO:2

**[0039]** In a most preferred embodiment the variant lipase is a variant of the lipase having the amino acid sequence of SEQ ID NO: 2.

#### **Substitutions in Region IV**

**[0040]** Region IV consists of amino acid residues that binds electrostatically to a surface. In this region it is preferred to substitute an amino acid of the parent lipase with a more positive amino acid.

**[0041]** The lipase variant may comprise at least one substitution in Region IV, such as two, three, four, five or six substitutions in Region IV.

**[0042]** Amino acid residues corresponding to the following positions are comprised by Region IV: 27 and 54 to 62. The following positions are of particular interest: 27, 56, 57, 58, 60.

**[0043]** In particular the following substitutions have been identified: X27R, X58N/AG/T/P and X60V/S/G/N/R/K/A/L.

**[0044]** In a preferred embodiment the variant lipase has at least 85%, 90%, 95%, 96%, 97%, 98% or 99% identity to SEQ ID NO:2

**[0045]** In a most preferred embodiment the variant lipase is a variant of the lipase having the amino acid sequence of SEQ ID NO: 2.

#### **Amino acids at other positions**

**[0046]** The parent lipase may optionally comprise additional alterations, e.g, substitution of other amino acids, particularly less than 10, less than 9, less than 8, less than 7, less than 6, less than 5 alterations as compared to a parent lipase. Examples are substitutions corresponding to one or more of the positions 24, 37, 38, 46, 74, 81, 83, 115, 127, 131, 137, 143, 147, 150, 199, 200, 203, 206, 211, 263, 264, 265, 267 and 269 of the parent lipase. In a particular embodiment there is a substitution in at least one of the positions corresponding to position 81, 147, 150, 227 and 249. In a preferred embodiment the at least one substitution is selected from the group consisting of X38R, X81Q/E, X143S/C/N/D/A, X147M/Y, X150G/K, X227G and X249R/I/L.

**[0047]** The variant may comprise substitutions outside the defined Region I to IV; the number of substitutions outside the defined Region I to IV is preferably less than six, such as five, four, three, two or one substitution.

**[0048]** Further substitutions may, e.g., be made according to principles known in the art, e.g. substitutions described in WO 92/05249, WO 94/25577, WO 95/22615, WO 97/04079 and WO 97/07202.

#### Parent lipase variants

**[0049]** Variant lipases include parent lipases identical to SEQ ID NO:2 having the substitutions listed in the below table (using SEQ ID NO:2 for numbering).

| Region I       | Region II | Region III | Region IV | Outside regions |
|----------------|-----------|------------|-----------|-----------------|
| T231 R + N233R | I255Y     |            |           |                 |

#### Nomenclature for amino acid modifications

**[0050]** In describing lipase variants according to the invention, the following nomenclature is used for ease of reference:

Original amino acid(s):position(s):substituted amino acid(s)

**[0051]** According to this nomenclature, for instance the substitution of glutamic acid for glycine in position 195 is shown as G195E. A deletion of glycine in the same position is shown as G195\*, and insertion of an additional amino acid residue such as lysine is shown as G195GK.

**[0052]** Where a specific lipase contains a "deletion" in comparison with other lipases and an insertion is made in such a position this is indicated as \*36D for insertion of an aspartic acid in position 36.

**[0053]** Multiple mutations are separated by pluses, i.e.: R170Y+G195E, representing mutations in positions 170 and 195 substituting tyrosine and glutamic acid for arginine and glycine, respectively.

**[0054]** X231 indicates the amino acid in a parent polypeptide corresponding to position 231, when applying the described alignment procedure. X231 R indicates that the amino acid is replaced with R. For SEQ ID NO:2 X is T, and X231 R thus indicates a substitution of T in position 231 with R. Where the amino acid in a position (e.g. 231) may be substituted by

another amino acid selected from a group of amino acids, e.g. the group consisting of R and P and Y, this will be indicated by X231R/P/Y.

**[0055]** In all cases, the accepted IUPAC single letter or triple letter amino acid abbreviation is employed.

#### **Amino acid grouping**

**[0056]** In this specification, amino acids are classified as negatively charged, positively charged or electrically neutral according to their electric charge at pH 10. Thus, negative amino acids are E, D, C (cysteine) and Y, particularly E and D. Positive amino acids are R, K and H, particularly R and K. Neutral amino acids are G, A, V, L, I, P, F, W, S, T, M, N, Q and C when forming part of a disulfide bridge. A substitution with another amino acid in the same group (negative, positive or neutral) is termed a conservative substitution.

**[0057]** The neutral amino acids may be divided into hydrophobic or non-polar (G, A, V, L, I, P, F, W and C as part of a disulfide bridge) and hydrophilic or polar (S, T, M, N, Q).

#### **Amino acid identity**

**[0058]** The relatedness between two amino acid sequences or between two nucleotide sequences is described by the parameter "identity".

**[0059]** For purposes of the present invention, the alignment of two amino acid sequences is determined by using the Needle program from the EMBOSS package (<http://emboss.org>) version 2.8.0. The Needle program implements the global alignment algorithm described in Needleman, S. B. and Wunsch, C. D. (1970) J. Mol. Biol. 48, 443-453. The substitution matrix used is BLOSUM62, gap opening penalty is 10, and gap extension penalty is 0.5.

**[0060]** The degree of identity between an amino acid sequence of the present invention ("invention sequence"; e.g. amino acids 1 to 269 of SEQ ID NO:2) and a different amino acid sequence ("foreign sequence") is calculated as the number of exact matches in an alignment of the two sequences, divided by the length of the "invention sequence" or the length of the "foreign sequence", whichever is the shortest. The result is expressed in percent identity.

**[0061]** An exact match occurs when the "invention sequence" and the "foreign sequence" have identical amino acid residues in the same positions of the overlap. The length of a sequence is the number of amino acid residues in the sequence (e.g. the length of SEQ ID NO:2 is 269).

**[0062]** The above procedure may be used for calculation of identity as well as homology and

for alignment. In the context of the present invention homology and alignment has been calculated as described below.

### Homology and alignment

**[0063]** For purposes of the present invention, the degree of homology may be suitably determined by means of computer programs known in the art, such as GAP provided in the GCG program package (Program Manual for the Wisconsin Package, Version 8, August 1994, Genetics Computer Group, 575 Science Drive, Madison, Wisconsin, USA 53711) (Needleman, S.B. and Wunsch, C.D., (1970), Journal of Molecular Biology, 48, 443-45), using GAP with the following settings for polypeptide sequence comparison: GAP creation penalty of 3.0 and GAP extension penalty of 0.1.

**[0064]** In the present invention, corresponding (or homologous) positions in the lipase sequences of *Absidia reflexa*, *Absidia corymbifera*, *Rhizomucor miehei*, *Rhizopus delemar*, *Aspergillus niger*, *Aspergillus tubigensis*, *Fusarium oxysporum*, *Fusarium heterosporum*, *Aspergillus oryzae*, *Penicillium camembertii*, *Aspergillus foetidus*, *Aspergillus niger*, *Thermomyces lanuginosus* (synonym: *Humicola lanuginosa*) and *Landerina penisapora* are defined by the alignment shown in Figure 1.

**[0065]** To find the homologous positions in lipase sequences not shown in the alignment, the sequence of interest is aligned to the sequences shown in Figure 1. The new sequence is aligned to the present alignment in Figure 1 by using the GAP alignment to the most homologous sequence found by the GAP program. GAP is provided in the GCG program package (Program Manual for the Wisconsin Package, Version 8, August 1994, Genetics Computer Group, 575 Science Drive, Madison, Wisconsin, USA 53711) (Needleman, S.B. and Wunsch, C.D., (1970), Journal of Molecular Biology, 48, 443-45). The following settings are used for polypeptide sequence comparison: GAP creation penalty of 3.0 and GAP extension penalty of 0.1.

### Hybridization

**[0066]** The present invention also relates to isolated polypeptides having lipase activity which are encoded by polynucleotides which hybridize under very low stringency conditions, preferably low stringency conditions, more preferably medium stringency conditions, more preferably medium-high stringency conditions, even more preferably high stringency conditions, and most preferably very high stringency conditions with (i) nucleotides 178 to 660 of SEQ ID NO: 1, (ii) the cDNA sequence contained in nucleotides 178 to 660 of SEQ ID NO: 1, (iii) a subsequence of (i) or (ii), or (iv) a complementary strand of (i), (ii), or (iii) (J. Sambrook, E.F. Fritsch, and T. Maniatis, 1989, Molecular Cloning, A Laboratory Manual, 2d edition, Cold Spring Harbor, New York). A subsequence of SEQ ID NO: 1 contains at least 100 contiguous

nucleotides or preferably at least 200 contiguous nucleotides. Moreover, the subsequence may encode a polypeptide fragment which has lipase activity.

**[0067]** For long probes of at least 100 nucleotides in length, very low to very high stringency conditions are defined as prehybridization and hybridization at 42°C in 5X SSPE, 0.3% SDS, 200 ug/ml sheared and denatured salmon sperm DNA, and either 25% formamide for very low and low stringencies, 35% formamide for medium and medium-high stringencies, or 50% formamide for high and very high stringencies, following standard Southern blotting procedures for 12 to 24 hours optimally.

**[0068]** For long probes of at least 100 nucleotides in length, the carrier material is finally washed three times each for 15 minutes using 2X SSC, 0.2% SDS preferably at least at 45°C (very low stringency), more preferably at least at 50°C (low stringency), more preferably at least at 55°C (medium stringency), more preferably at least at 60°C (medium-high stringency), even more preferably at least at 65°C (high stringency), and most preferably at least at 70°C (very high stringency).

#### **DNA sequence, Expression vector, Host cell, Production of lipase**

**[0069]** The invention provides a DNA sequence encoding the lipase of the invention, an expression vector harboring the DNA sequence, and a transformed host cell containing the DNA sequence or the expression vector. These may be obtained by methods known in the art.

**[0070]** The invention also provides a method of producing the lipase by culturing the transformed host cell under conditions conducive for the production of the lipase and recovering the lipase from the resulting broth. The method may be practiced according to principles known in the art.

#### **Lipase activity**

##### **Lipase activity on tributyrin at neutral pH (LU)**

**[0071]** A substrate for lipase is prepared by emulsifying tributyrin (glycerin tributyrate) using gum Arabic as emulsifier. The hydrolysis of tributyrin at 30 °C at pH 7 or 9 is followed in a pH-stat titration experiment. One unit of lipase activity (1 LU) equals the amount of enzyme capable of releasing 1 micro mol butyric acid/min at pH 7.

##### **Benefit Risk**

**[0072]** The Benefit Risk factor describing the performance compared to the reduced risk for odor smell is defined as:  $BR = RP_{avg} / R$ , as described below.

### Uses

**[0073]** Enzymes of the present invention may find industrial use, e.g. be included in detergent compositions for removing of fatty matter.

### EXAMPLES

**[0074]** Chemicals used as buffers and substrates were commercial products of at least reagent grade.

### Media and Solutions

**[0075]**

|           |             |
|-----------|-------------|
| Product   | Tradename   |
| LAS :     | Surfac PS   |
| Zeolite A | Wessalith P |

Other ingredients used are standard laboratory reagents.

### Materials

**[0076]**

|         |                                                                      |
|---------|----------------------------------------------------------------------|
| Product | Supplier                                                             |
| EMPA221 | EMPA St. Gallen, Lerchfeldstrasse 5, CH-9014 St. Gallen, Switzerland |

### Example 1

### Production of enzyme

**[0077]** A plasmid containing the gene encoding the lipase is constructed and transformed into a suitable host cell using standard methods of the art.

**[0078]** Fermentation is carried out as a fed-batch fermentation using a constant medium temperature of 34°C and a start volume of 1.2 liter. The initial pH of the medium is set to 6.5. Once the pH has increased to 7.0 this value is maintained through addition of 10% H<sub>3</sub>PO<sub>4</sub>. The level of dissolved oxygen in the medium is controlled by varying the agitation rate and using a fixed aeration rate of 1.0 liter air per liter medium per minute. The feed addition rate is maintained at a constant level during the entire fed-batch phase.

The batch medium contains maltose syrup as carbon source, urea and yeast extract as nitrogen source and a mixture of trace metals and salts. The feed added continuously during the fed-batch phase contains maltose syrup as carbon source whereas yeast extract and urea is added in order to assure a sufficient supply of nitrogen.

**[0079]** Purification of the lipase may be done by use of standard methods known in the art, e.g. by filtering the fermentation supernatant and subsequent hydrophobic chromatography and ion exchange chromatography, e.g. as described in EP 0 851 913 EP, Example 3.

## Example 2

### AMSA - Automated Mechanical Stress Assay - for calculation of Relative Performance (RP)

**[0080]** The enzyme variants of the present application are tested using the Automatic Mechanical Stress Assay (AMSA). With the AMSA test the wash performance of a large quantity of small volume enzyme-detergent solutions can be examined. The AMSA plate has a number of slots for test solutions and a lid firmly squeezing the textile swatch to be washed against all the slot openings. During the washing time, the plate, test solutions, textile and lid are vigorously shaken to bring the test solution in contact with the textile and apply mechanical stress. For further description see WO 02/42740 especially the paragraph "Special method embodiments" at page 23-24. The containers, which contain the detergent test solution, consist of cylindrical holes (6 mm diameter, 10 mm depth) in a metal plate. The stained fabric (test material) lies on the top of the metal plate and is used as a lid and seal on the containers. Another metal plate lies on the top of the stained fabric to avoid any spillage from each container. The two metal plate together with the stained fabric are vibrated up and down at a frequency of 30 Hz with an amplitude of 2 mm.

**[0081]** The assay is conducted under the experimental conditions specified below:

**Table 3**

|               |                                          |
|---------------|------------------------------------------|
|               | 0.5 g/l LAS                              |
| Test solution | 0.52 g/l Na <sub>2</sub> CO <sub>3</sub> |
|               | 1.07 g/l Zeolite A                       |

|                                       |                                                                                                                             |
|---------------------------------------|-----------------------------------------------------------------------------------------------------------------------------|
|                                       | 0.52 g/l Na <sub>3</sub> Citrat                                                                                             |
| Test solution volume                  | 160 micro l                                                                                                                 |
| pH                                    | As is ( $\approx 9.9$ )                                                                                                     |
| Wash time                             | 20 minutes                                                                                                                  |
| Temperature                           | 30 °C                                                                                                                       |
| Water hardness                        | 15°dH                                                                                                                       |
|                                       | Ratio of Ca <sup>2+</sup> /Mg <sup>2+</sup> /NaHCO <sub>3</sub> : 4:1:7.5                                                   |
| Enzyme concentration in test solution | 0.125, 0.25, 0.50, 1.0 mg ep / l                                                                                            |
| Drying                                | Performance: After washing the textile pieces is immediately flushed in tap water and air-dried at 85°C in 5 min            |
|                                       | Odor: After washing the textile pieces is immediately flushed in tap water and dried at room temperature (20°C) for 2 hours |
| Test material                         | Cream turmeric swatch as described below (EMPA221 used as cotton textile)                                                   |

**[0082]** Cream-turmeric swatches were prepared by mixing 5 g of turmeric (Santa Maria, Denmark) with 100 g cream (38% fat, Arla, Denmark) at 50 °C, the mixture was left at this temperature for about 20 minutes and filtered (50°C) to remove any undissolved particles. The mixture is cooled to 20°C) woven cotton swatches, EMPA221, were immersed in the cream-turmeric mixture and afterwards allowed to dry at room temperature over night and frozen until use. The preparation of cream-turmeric swatches is disclosed in the patent application WO 2006/125437.

**[0083]** The performance of the enzyme variant is measured as the brightness of the colour of the textile samples washed with that specific enzyme variant. Brightness can also be expressed as the intensity of the light reflected from the textile sample when luminated with white light. When the textile is stained the intensity of the reflected light is lower, than that of a clean textile. Therefore the intensity of the reflected light can be used to measure wash performance of an enzyme variant.

**[0084]** Color measurements are made with a professional flatbed scanner (PFU DL2400pro), which is used to capture an image of the washed textile samples. The scans are made with a resolution of 200 dpi and with an output color depth of 24 bits. In order to get accurate results, the scanner is frequently calibrated with a Kodak reflective IT8 target.

**[0085]** To extract a value for the light intensity from the scanned images, a special designed software application is used (Novozymes Color Vector Analyzer). The program retrieves the 24

bit pixel values from the image and converts them into values for red, green and blue (RGB). The intensity value (Int) is calculated by adding the RGB values together as vectors and then taking the length of the resulting vector:

$$Int = \sqrt{r^2 + g^2 + b^2}$$

**[0086]** The wash performance (P) of the variants is calculated in accordance with the below formula:

$$P = Int(v) - Int(r)$$

where

Int(v) is the light intensity value of textile surface washed with enzyme, and

Int(r) is the light intensity value of textile surface washed without enzyme.

A relative performance score is given as the result of the AMSA wash in accordance with the definition:

Relative Performance scores (RP) are summing up the performances (P) of the tested enzyme variants against the reference enzyme:

$$RP = P(\text{test enzyme}) / P(\text{reference enzyme}).$$

RPavg indicates the average relative performance compared to the reference enzyme at all four enzyme concentrations (0.125, 0.25, 0.5, 1.0 mg ep/l)

$$RP_{avg} = \text{avg}(RP(0.125), RP(0.25), RP(0.5), RP(1.0))$$

A variant is considered to exhibit improved wash performance, if it performs better than the reference.

In the context of the present invention the reference enzyme is the lipase of SEQ ID NO:2 with the substitutions T231 R + N233R.

### Example 3

#### GC - Gas Chromatograph - for calculation of risk factor

**[0087]** The butyric acid release from the lipase washed swatches were measured by Solid Phase Micro Extraction Gas Chromatography (SPME-GC) using the following method. Four textile pieces (5 mm in diameter), washed in the specified solution in Table 3 containing 1 mg/l lipase, were transferred to a Gas Chromatograph (GC) vial. The samples were analysed on a Varian 3800 GC equipped with a Stabilwax- DA w/Integra-Guard column (30m, 0.32 mm ID and 0.25 micro-m df) and a Carboxen PDMS SPME fibre (75 micro-m). Each sample was preincubated for 10 min at 40°C followed by 20 min sampling with the SPME fibre in the head-space over the textile pieces. The sample was subsequently injected onto the column (injector temperature=250°C). Column flow = 2 ml Helium/min. Column oven temperature gradient: 0 min = 40°C, 2 min = 40°C, 22 min = 240°C, 32 min = 240°C. The butyric acid was detected by

FID detection and the amount of butyric acid was calculated based on a butyric acid standard curve.

**[0088]** The Risk Performance Odour, R, of a lipase variant is the ratio between the amount of released butyric acid from the lipase variant washed swatch and the amount of released butyric acid from a swatch washed with the lipase of SEQ ID NO: 2 with the substitutions T231 R + N233R (reference enzyme), after both values have been corrected for the amount of released butyric acid from a non-lipase washed swatch. The risk (R) of the variants is calculated in accordance with the below formula:

Odour = measured in micro g butyric acid developed at 1 mg enzyme protein / l corrected for blank

$$\text{Alpha}_{\text{test enzyme}} = \text{Odour}_{\text{test enzyme}} - \text{Blank}$$

$$\text{Alpha}_{\text{reference enzyme}} = \text{Odour}_{\text{reference enzyme}} - \text{Blank}$$

$$R = \text{Alpha}_{\text{test enzyme}} / \text{Alpha}_{\text{reference enzyme}}$$

A variant is considered to exhibit reduced odor compared to the reference, if the R factor is lower than 1.

#### **Example 4**

#### **Activity (LU) relative to absorbance at 280nm**

**[0089]** The activity of a lipase relative to the absorbance at 280 nm is determined by the following assay:

#### **LU/A280:**

**[0090]** The activity of the lipase is determined as described above in the section Lipase activity. The absorbance of the lipase at 280 nm is measured (A280) and the ratio LU/A280 is calculated. The relative LU/A280 is calculated as the LU/A280 of the variant divided by the LU/A280 of a reference enzyme. In the context of the present invention the reference enzyme is the lipase of SEQ ID NO:2 with the substitutions T231 R + N233R.

#### **Example 5**

#### **BR - Benefit Risk**

**[0091]** The Benefit Risk factor describing the performance compared to the reduced risk for odour smell is thus defined as:

$$BR = RP_{avg} / R$$

A variant is considered to exhibit improved wash performance and reduced odor, if the BR factor is higher than 1.

**[0092]** Applying the above methods the following results were obtained:

**Table 4**

| Variant   | Mutations in SEQ ID NO: 2                                                                                      | Average RP<br>( $RP_{avg}$ ) | BR             | LU/A280        |
|-----------|----------------------------------------------------------------------------------------------------------------|------------------------------|----------------|----------------|
| 1         | I202G + T231R + N233R                                                                                          | 0.84                         | 1.41           | not determined |
| 2         | I86V + L227G + T231R + N233R + P256K                                                                           | 1.08                         | 1.52           | 1700           |
| 3         | Q4V + S58N + V60S + T231R + N233R                                                                              | 0.87                         | 1.73           | 1950           |
| 4         | S58N + V60S + I90R + T231R + N233R                                                                             | 1.06                         | 1.27           | 2250           |
| 5         | I255Y + T231R + N233R                                                                                          | 1.19                         | 1.17           | 3600           |
| 6         | I90A + T231R + N233R + I255V                                                                                   | 1.13                         | 1.14           | 2700           |
| Reference | T231R + N233R                                                                                                  | 1.00                         | 1.00           | 3650           |
| 7         | G91A + E99K + T231R+N233R + Q249R + 270H + 271T + 272P + 273S + 274S + 275G + 276R + 277G + 278G + 279H + 280R | 0.43                         | not determined | 850            |
| 8         | G91A + E99K + T231 R, N233R + Q249R + 270H + 271T + 272P + 273S + 274S + 275G + 276R + 277G + 278G             | 0.13                         | not determined | 500            |

**[0093]** The reference lipase and variants 7 and 8 in Table 4 are described in WO 2000/060063.

#### Example 6

#### BR - Benefit Risk

**[0094]** The Benefit Risk was measured for the variants listed in Table 5. The Benefit Risk

factor was measured in the same way as described in Example 5 and it was found to be above 1 for all the listed variants.

**Table 5**

| Variant   | Mutations in SEQ ID NO: 2                                                    |
|-----------|------------------------------------------------------------------------------|
| Reference | T231R + N233R                                                                |
| 9         | L97V+ T231R+N233R                                                            |
| 10        | A150G+T231R+N233R                                                            |
| 11        | I90R+T231R+N233R                                                             |
| 12        | I202V+T231R+N233R                                                            |
| 13        | L227G+ T231R+ N233R+ P256K                                                   |
| 14        | I90A+ T231R+ N233R                                                           |
| 15        | T231R+N233R+I255P                                                            |
| 16        | I90V+I255V+T231R+N233R                                                       |
| 17        | F211L+ L227G+ T231R+ N233R+ I255L+ P256K                                     |
| 18        | S58N+ V60S+ T231R+ N233R+ Q249L                                              |
| 19        | S58N+ V60S+ T231R+ N233R+ Q249I                                              |
| 20        | A150G+ L227G+ T231R+ N233R+ P256K                                            |
| 21        | K46L+ S58N+ V60S+ T231R+ N233R+ Q249L+ D254I                                 |
| 22        | Q4L+ E43T+ K46I+ S58N+ V60S+ T231R+ N233R+ Q249L+ D254I                      |
| 23        | Q4L+ S58N+ V60S+ T231R+ N233R+ Q249L+ D254I                                  |
| 24        | K46I+ S58N+ V60S+ T231R+ N233R+ Q249L+ D254L                                 |
| 25        | K46L+ S58N+ V60S+ K223I+ T231R+ N233R+ D254I                                 |
| 26        | E43T+ K46I+ S58N+ V60S+ T231R+ N233R+ Q249L+ D254I                           |
| 27        | S58N+ V60S+ I86V+ A150G+ L227G+ T231 R+ N233R+ P256K                         |
| 28        | K24R+ K46R+ K74R+ I86V+ K98R+ K127R+ D137K+ A150G+ K223R+ T231R+ N233R       |
| 29        | S58A+V60A+ I86V+T231R+N233R                                                  |
| 30        | K24R+ K46R+ S58N+ V60S+ K74R+ I86V+ K98R+ K127R+ D137K+ K223R+ T231 R+ N233R |
| 31        | S58A+ V60A+ I86V+ A150G+ T231R+ N233R                                        |
| 32        | S58N+ V60V+ D62G+ T231R+ N233R                                               |
| 33        | Q4V+ S58N+ V60S+ I86V+ T231R+ N233R+ Q249L                                   |
| 34        | Q4V+ S58N+ V60S+ I86V+ A150G+ T231R+ N233R+ I255V                            |
| 35        | Q4V+ S58N+ V60S+ I90A+ A150G+ T231R+ N233R+ I255V                            |
| 36        | Y53A+ S58N+ V60S+ T231R+ N233R+ P256L                                        |
| 37        | I202L+ T231 R+ N233R+ I255A                                                  |
| 38        | S58A+ V60S+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K                          |

| Variant | Mutations in SEQ ID NO: 2                                                 |
|---------|---------------------------------------------------------------------------|
| 39      | D27R+ S58N+ V60S+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K                 |
| 40      | V60K+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K                             |
| 41      | Q4V+ S58A+ V60S+ S83T+ I86V+ A150G+ E210K+ L227G+ T231R+ N233R+ P256K     |
| 42      | Q4V+ V60K+ S83T+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K                  |
| 43      | D27R+ V60K+ I86V+ A150G+ L227G+ T231 R+ N233R+ P256K                      |
| 44      | Q4N+ L6S+ S58N+ V60S+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K             |
| 45      | E1 N+ V60K+ I86V+ A150G+ L227G+ T231R+ N233R+ P256K                       |
| 46      | V60K+ I86V+ A150G+ K223N+ G225S+ T231R+ N233R+ P256K                      |
| 47      | E210V+ T231R+ N233R+ Q249R                                                |
| 48      | S58N+ V60S+ E210V+ T231R+ N233R+ Q249R                                    |
| 49      | Q4V+ V60K+ I90R+ T231R+ N233R+ I255V                                      |
| 50      | Q4V+ V60K+ A150G+ T231R+ N233R                                            |
| 51      | V60K+ S83T+ T231R+ N233R                                                  |
| 52      | V60K+ A150G+ T231R+ N233R+ I255V                                          |
| 53      | T231R+ N233G+ D234G                                                       |
| 54      | S58N+ V60S+ I86V+ A150G+ E210K+ L227G+ T231R+ N233R+ Q249R+ P256K         |
| 55      | S58N+ V60S+ I86V+ A150G+ E210K+ L227G+ T231R+ N233R+ I255A+ P256K         |
| 56      | S58N+ V60S+ I86V+ A150G+ G156R+ E210K+ L227G+ T231 R+ N233R+ I255A+ P256K |
| 57      | S58T+ V60K+ I86V+ N94K+ A150G+ E210V+ L227G+ T231R+ N233R+ P256K          |
| 58      | S58T+ V60K+ I86V+ D102A+ A150G+ L227G+ T231R+ N233R+ P256K                |
| 59      | S58T+ V60K+ I86V+ D102A+ A150G+ E210V+ L227G+ T231R+ N233R+ P256K         |
| 60      | S58T+ V60K+ S83T+ I86V+ N94K+ A150G+ E210V+ L227G+ T231R+ N233R+ P256K    |
| 61      | S58A+ V60S+ I86V+ T143S+ A150G+ L227G+ T231R+ N233R+ P256K                |
| 62      | G91S+ D96V+ D254R                                                         |
| 63      | V60L+ G91M+ T231W+ Q249L                                                  |
| 64      | T37A+ D96A+ T231R+ N233R+ Q249G                                           |
| 65      | E56G+E87D+T231R+N233R+D254A                                               |
| 66      | E210K+T231R+N233R                                                         |

| Variant | Mutations in SEQ ID NO: 2                                         |
|---------|-------------------------------------------------------------------|
| 67      | D27H+E87Q+D96N+T231R+N233R+D254V                                  |
| 68      | F181L+E210V+T231R+N233R                                           |
| 69      | D27N+ D96G+ T231R+ N233R                                          |
| 70      | D96N+ T231R+ N233R                                                |
| 71      | T231 R+ N233I+ D234G                                              |
| 72      | S58K+ V60L+ E210V+ Q249R                                          |
| 73      | S58H+ V60L+ E210V+ Q249R                                          |
| 74      | Q4V+ F55V+ I86V+ T231R+ N233R+ I255V                              |
| 75      | Q4V+ S58T+ V60K+ T199L+ N200A+ E210K+ T231 R+ N233R+ I255A+ P256K |
| 76      | Q4V+ D27N+ V60K+ T231 R+ N233R                                    |
| 77      | I90F+ I202P+ T231R+ N233R+ I255L                                  |
| 78      | S58N+ V60S+ D158N+ T231R+ N233R                                   |
| 79      | S58N+ V60S+ S115K+ T231R+ N233R                                   |
| 80      | S58N+ V60S+ L147M+ A150G+ F211L+ T231 R+ N233R                    |
| 81      | V60K+ A150G+ T231 R+ N233R                                        |
| 82      | I90V+L227G+T231 R+N233R+ P256K                                    |
| 83      | T231R+N233R+I255S                                                 |
| 84      | I86G+ T231 R+ N233R                                               |
| 85      | V60K+ I202V+ E210K+ T231R+ N233R+ I255A+ P256K                    |
| 86      | I90G+ I202L+ T231 R+ N233R+ I255S                                 |
| 87      | S58G+ V60G+ T231 R+ N233R                                         |

[0095] The reference lipase is described in WO 2000/060063.

#### SEQUENCE LISTING

[0096]

<110> Novozymes A/S

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 Bjornvad, Mads Eskelund  
 Hansen, Peter Kamp  
 Ge, Haiyan  
 Lamsa, Michael

<120> Lipase variants

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| 1 5 10 15                                                       |     |
| tct gca gcc gca tac tgc gga aaa aac aat gat gcc cca gct ggt aca | 96  |
| Ser Ala Ala Ala Tyr Cys Gly Lys Asn Asn Asp Ala Pro Ala Gly Thr |     |
| 20 25 30                                                        |     |
| aac att acg tgc acg gga aat gcc tgc ccc gag gta gag aag gcg gat | 144 |
| Asn Ile Thr Cys Thr Gly Asn Ala Cys Pro Glu Val Glu Lys Ala Asp |     |
| 35 40 45                                                        |     |
| gca acg ttt ctc tac tcg ttt gaa gac tct gga gtg ggc gat gtc acc | 192 |
| Ala Thr Phe Leu Tyr Ser Phe Glu Asp Ser Gly Val Gly Asp Val Thr |     |
| 50 55 60                                                        |     |
| ggc ttc ctt gct ctc gac aac acg aac aaa ttg atc gtc ctc tct ttc | 240 |
| Gly Phe Leu Ala Leu Asp Asn Thr Asn Lys Leu Ile Val Leu Ser Phe |     |
| 65 70 75 80                                                     |     |
| cgt ggc tct cgt tcc ata gag aac tgg atc ggg aat ctt aac ttc gac | 288 |
| Arg Gly Ser Arg Ser Ile Glu Asn Trp Ile Gly Asn Leu Asn Phe Asp |     |
| 85 90 95                                                        |     |
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| Leu Lys Glu Ile Asn Asp Ile Cys Ser Gly Cys Arg Gly His Asp Gly |     |
| 100 105 110                                                     |     |
| ttc act tcg tcc tgg agg tct gta gcc gat acg tta agg cag aag gtg | 384 |
| Phe Thr Ser Ser Trp Arg Ser Val Ala Asp Thr Leu Arg Gln Lys Val |     |

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gag gat gct gtg agg gag cat ccc gac tat cgc gtg gtg ttt acc gga      432
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His Ser Leu Gly Gly Ala Leu Ala Thr Val Ala Gly Ala Asp Leu Arg
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Gly Asn Gly Tyr Asp Ile Asp Val Phe Ser Tyr Gly Ala Pro Arg Val
165              170              175

gga aac agg gct ttt gca gaa ttc ctg acc gta cag acc ggc gga aca      576
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ctc tac cgc att acc cac acc aat gat att gtc cct aga ctc ccg ccg      624
Leu Tyr Arg Ile Thr His Thr Asn Asp Ile Val Pro Arg Leu Pro Pro
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cgc gaa ttc ggt tac agc cat tct agc cca gag tac tgg atc aaa tct      672
Arg Glu Phe Gly Tyr Ser His Ser Ser Pro Glu Tyr Trp Ile Lys Ser
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gga acc ctt gtc ccc gtc acc cga aac gat atc gtg aag ata gaa ggc      720
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&lt;212&gt; PRT

&lt;213&gt; Thermomyces lanuginosus

&lt;400&gt; 2

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20              25              30

Asn Ile Thr Cys Thr Gly Asn Ala Cys Pro Glu Val Glu Lys Ala Asp
35              40              45

Ala Thr Phe Leu Tyr Ser Phe Glu Asp Ser Gly Val Gly Asp Val Thr
50              55              60

Gly Phe Leu Ala Leu Asp Asn Thr Asn Lys Leu Ile Val Leu Ser Phe
65              70              75              80

Arg Gly Ser Arg Ser Ile Glu Asn Trp Ile Gly Asn Leu Asn Phe Asp
85              90              95

Leu Lys Glu Ile Asn Asp Ile Cys Ser Gly Cys Arg Gly His Asp Gly
100              105              110

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His Ser Leu Gly Gly Ala Leu Ala Thr Val Ala Gly Ala Asp Leu Arg  
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Gly Asn Gly Tyr Asp Ile Asp Val Phe Ser Tyr Gly Ala Pro Arg Val  
165 170 175

Gly Asn Arg Ala Phe Ala Glu Phe Leu Thr Val Gln Thr Gly Gly Thr  
180 185 190

Leu Tyr Arg Ile Thr His Thr Asn Asp Ile Val Pro Arg Leu Pro Pro  
195 200 205

Arg Glu Phe Gly Tyr Ser His Ser Ser Pro Glu Tyr Trp Ile Lys Ser  
210 215 220

Gly Thr Leu Val Pro Val Thr Arg Asn Asp Ile Val Lys Ile Glu Gly  
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<212> PRT

<213> Absidia reflexa

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20 25 30

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35 40 45

Ser Asn Leu Gln Ile Thr Lys Thr Phe Ser Thr Leu Ile Thr Asp Thr  
50 55 60

Asn Val Leu Val Ala Val Gly Glu Lys Glu Lys Thr Ile Tyr Val Val  
65 70 75 80

Phe Arg Gly Thr Ser Ser Ile Arg Asn Ala Ile Ala Asp Ile Val Phe  
85 90 95

Val Pro Val Asn Tyr Pro Pro Val Asn Gly Ala Lys Val His Lys Gly  
100 105 110

Phe Leu Asp Ser Tyr Asn Glu Val Gln Asp Lys Leu Val Ala Glu Val  
115 120 125

Lys Ala Gln Leu Asp Arg His Pro Gly Tyr Lys Ile Val Val Thr Gly  
130 135 140

His Ser Leu Gly Gly Ala Thr Ala Val Leu Ser Ala Leu Asp Leu Tyr  
145 150 155 160

His His Gly His Ala Asn Ile Glu Ile Tyr Thr Gln Gly Gln Pro Arg  
165 170 175

Ile Gly Thr Pro Ala Phe Ala Asn Tyr Val Ile Gly Thr Lys Ile Pro  
180 185 190

Tyr Gln Arg Leu Val His Glu Arg Asp Ile Val Pro His Leu Pro Pro  
195 200 205

Gly Ala Phe Gly Phe Leu His Ala Gly Glu Glu Phe Trp Ile Met Lys  
210 215 220

Asp Ser Ser Leu Arg Val Cys Pro Asn Gly Ile Glu Thr Asp Asn Cys  
225 230 235 240

Ser Asn Ser Ile Val Pro Phe Thr Ser Val Ile Asp His Leu Ser Tyr  
245 250 255

Leu Asp Met Asn Thr Gly Leu Cys Leu  
260 265

<210> 4

<211> 264

<212> PRT

<213> Absidia corymbifera

<400> 4

Ser Ser Ser Thr Gln Asp Tyr Arg Ile Ala Ser Glu Ala Glu Ile Lys  
1 5 10 15

Ala His Thr Phe Tyr Thr Ala Leu Ser Ala Asn Ala Tyr Cys Arg Thr  
20 25 30

Val Ile Pro Gly Gly Gln Trp Ser Cys Pro His Cys Asp Val Ala Pro  
35 40 45

Asn Leu Asn Ile Thr Lys Thr Phe Thr Thr Leu Ile Thr Asp Thr Asn  
50 55 60

Val Leu Val Ala Val Gly Glu Asn Glu Lys Thr Ile Tyr Val Val Phe  
65 70 75 80

Arg Gly Thr Ser Ser Ile Arg Asn Ala Ile Ala Asp Ile Val Phe Val  
85 90 95

Pro Val Asn Tyr Pro Pro Val Asn Gly Ala Lys Val His Lys Gly Phe  
100 105 110

Leu Asp Ser Tyr Asn Glu Val Gln Asp Lys Leu Val Ala Glu Val Lys  
115 120 125

Ala Gln Leu Asp Arg His Pro Gly Tyr Lys Ile Val Val Thr Gly His  
130 135 140

Ser Leu Gly Gly Ala Thr Ala Val Leu Ser Ala Leu Asp Leu Tyr His  
145 150 155 160

His Gly His Asp Asn Ile Glu Ile Tyr Thr Gln Gly Gln Pro Arg Ile  
165 170 175

Gly Thr Pro Glu Phe Ala Asn Tyr Val Ile Gly Thr Lys Ile Pro Tyr  
180 185 190

Gln Arg Leu Val Asn Glu Arg Asp Ile Val Pro His Leu Pro Pro Gly  
195 200 205

Ala Phe Gly Phe Leu His Ala Gly Glu Glu Phe Trp Ile Met Lys Asp  
210 215 220

Ser Ser Leu Arg Val Cys Pro Asn Gly Ile Glu Thr Asp Asn Cys Ser  
225 230 235 240

Asn Ser Ile Val Pro Phe Thr Ser Val Ile Asp His Leu Ser Tyr Leu  
245 250 255

Asp Met Asn Thr Gly Leu Cys Leu  
260

<210> 5

<211> 269

<212> PRT

<213> Rhizomucor miehei

<400> 5

Ser Ile Asp Gly Gly Ile Arg Ala Ala Thr Ser Gln Glu Ile Asn Glu  
1 5 10 15

Leu Thr Tyr Tyr Thr Thr Leu Ser Ala Asn Ser Tyr Cys Arg Thr Val  
20 25 30

Ile Pro Gly Ala Thr Trp Asp Cys Ile His Cys Asp Ala Thr Glu Asp  
35 40 45

Leu Lys Ile Ile Lys Thr Trp Ser Thr Leu Ile Tyr Asp Thr Asn Ala  
50 55 60

Met Val Ala Arg Gly Asp Ser Glu Lys Thr Ile Tyr Ile Val Phe Arg  
65 70 75 80

Glu Ser Ser Ser Ile Asp Asp Thr Ile Ala Asn Val Thr Phe Val Pro

Gly Ser Ser Ser Ile Arg Asn Trp Ile Ala Asp Leu Thr Phe Val Pro  
85 90 95

Val Ser Tyr Pro Pro Val Ser Gly Thr Lys Val His Lys Gly Phe Leu  
100 105 110

Asp Ser Tyr Gly Glu Val Gln Asn Glu Leu Val Ala Thr Val Leu Asp  
115 120 125

Gln Phe Lys Gln Tyr Pro Ser Tyr Lys Val Ala Val Thr Gly His Ser  
130 135 140

Leu Gly Gly Ala Thr Ala Leu Leu Cys Ala Leu Asp Leu Tyr Gln Arg  
145 150 155 160

Glu Glu Gly Leu Ser Ser Ser Asn Leu Phe Leu Tyr Thr Gln Gly Gln  
165 170 175

Pro Arg Val Gly Asp Pro Ala Phe Ala Asn Tyr Val Val Ser Thr Gly  
180 185 190

Ile Pro Tyr Arg Arg Thr Val Asn Glu Arg Asp Ile Val Pro His Leu  
195 200 205

Pro Pro Ala Ala Phe Gly Phe Leu His Ala Gly Glu Glu Tyr Trp Ile  
210 215 220

Thr Asp Asn Ser Pro Glu Thr Val Gln Val Cys Thr Ser Asp Leu Glu  
225 230 235 240

Thr Ser Asp Cys Ser Asn Ser Ile Val Pro Phe Thr Ser Val Leu Asp  
245 250 255

His Leu Ser Tyr Phe Gly Ile Asn Thr Gly Leu Cys Thr  
260 265

<210> 6

<211> 271

<212> PRT

<213> Rhizopus oryzae

<400> 6

Ser Ala Ser Asp Gly Gly Lys Val Val Ala Ala Thr Thr Ala Gln Ile  
1 5 10 15

Gln Glu Phe Thr Lys Tyr Ala Gly Ile Ala Ala Thr Ala Tyr Cys Arg  
20 25 30

Ser Val Val Pro Gly Asn Lys Trp Asp Cys Val Gln Cys Gln Lys Trp  
35 40 45

Val Pro Asp Gly Lys Ile Ile Thr Thr Phe Thr Ser Leu Leu Ser Asp  
50 55 60

Thr Asn Gly Tyr Val Leu Arg Ser Asp Lys Gln Lys Thr Ile Tyr Leu  
65 70 75 80

Val Phe Arg Gly Thr Asn Ser Phe Arg Ser Ala Ile Thr Asp Ile Val  
85 90 95

Phe Asn Phe Ser Asp Tyr Lys Pro Val Lys Gly Ala Lys Val His Ala  
100 105 110

Gly Phe Leu Ser Ser Tyr Glu Gln Val Val Asn Asp Tyr Phe Pro Val  
115 120 125

Val Gln Glu Gln Leu Thr Ala His Pro Thr Tyr Lys Val Ile Val Thr  
130 135 140

Gly His Ser Leu Gly Gly Ala Gln Ala Leu Leu Ala Gly Met Asp Leu  
145 150 155 160

Tyr Gln Arg Glu Pro Arg Leu Ser Pro Lys Asn Leu Ser Ile Phe Thr  
165 170 175

Val Gly Gly Pro Arg Val Gly Asn Pro Thr Phe Ala Tyr Tyr Val Glu  
180 185 190

Ser Thr Gly Ile Pro Phe Gln Arg Thr Val His Lys Arg Asp Ile Val  
195 200 205

Pro His Val Pro Pro Gln Ser Phe Gly Phe Leu His Pro Gly Val Glu  
210 215 220

Ser Trp Ile Lys Ser Gly Thr Ser Asn Val Gln Ile Cys Thr Ser Glu  
225 230 235 240

Ile Glu Thr Lys Asp Cys Ser Asn Ser Ile Val Pro Phe Thr Ser Ile  
245 250 255

Leu Asp His Leu Ser Tyr Phe Asp Ile Asn Glu Gly Ser Cys Leu  
260 265 270

<210> 7

<211> 267

<212> PRT

<213> Aspergillus niger

<400> 7

Thr Ala Gly His Ala Leu Ala Ala Ser Thr Gln Gly Ile Ser Glu Asp  
1 5 10 15

Leu Tyr Ser Arg Leu Val Glu Met Ala Thr Ile Ser Gln Ala Ala Tyr  
20 25 30

Ala Asp Leu Cys Asn Ile Pro Ser Thr Ile Ile Lys Gly Glu Lys Ile  
35 40 45

Tyr Asn Ser Gln Thr Asp Ile Asn Gly Trp Ile Leu Arg Asp Asp Ser  
50 55 60

Ser Lys Glu Ile Ile Thr Val Phe Arg Gly Thr Gly Ser Asp Thr Asn  
65 70 75 80

Leu Gln Leu Asp Thr Asn Tyr Thr Leu Thr Pro Phe Asp Thr Leu Pro  
85 90 95

Gln Cys Asn Gly Cys Glu Val His Gly Gly Tyr Tyr Ile Gly Trp Val  
100 105 110

Ser Val Gln Asp Gln Val Glu Ser Leu Val Lys Gln Gln Val Ser Gln  
115 120 125

Tyr Pro Asp Tyr Ala Leu Thr Val Thr Gly His Ser Leu Gly Ala Ser  
130 135 140

Leu Ala Ala Leu Thr Ala Ala Gln Leu Ser Ala Thr Tyr Asp Asn Ile  
145 150 155 160

Arg Leu Tyr Thr Phe Gly Glu Pro Arg Ser Gly Asn Gln Ala Phe Ala  
165 170 175

Ser Tyr Met Asn Asp Ala Phe Gln Ala Ser Ser Pro Asp Thr Thr Gln  
180 185 190

Tyr Phe Arg Val Thr His Ala Asn Asp Gly Ile Pro Asn Leu Pro Pro  
195 200 205

Val Glu Gln Gly Tyr Ala His Gly Gly Val Glu Tyr Trp Ser Val Asp  
210 215 220

Pro Tyr Ser Ala Gln Asn Thr Phe Val Cys Thr Gly Asp Glu Val Gln  
225 230 235 240

Cys Cys Glu Ala Gln Gly Gly Gln Gly Val Asn Asn Ala His Thr Thr  
245 250 255

Tyr Phe Gly Met Thr Ser Gly Ala Cys Thr Trp  
260 265

<210> 8

<211> 266

<212> PRT

<213> Aspergillus tubingensis

<400> 8

Thr Ala Gly His Ala Leu Ala Ala Ser Thr Gln Gly Ile Ser Glu Asp  
1 5 10 15

Leu Tyr Ser Arg Leu Val Glu Met Ala Thr Ile Ser Gln Ala Ala Tyr  
20 25 30

Ala Asp Leu Cys Asn Ile Pro Ser Thr Ile Ile Lys Gly Glu Lys Ile  
35 40 45

Tyr Asn Ser Gln Thr Asp Ile Asn Gly Trp Ile Leu Arg Asp Asp Ser  
50 55 60

Ser Lys Glu Ile Ile Thr Val Phe Arg Gly Thr Gly Ser Asp Thr Asn  
65 70 75 80

Leu Gln Leu Asp Thr Asn Tyr Thr Leu Thr Pro Phe Asp Thr Leu Pro  
85 90 95

Gln Cys Asn Ser Cys Glu Val His Gly Gly Tyr Tyr Ile Gly Trp Ile  
100 105 110

Ser Val Gln Asp Gln Val Glu Ser Leu Val Gln Gln Gln Val Ser Gln  
115 120 125

Phe Pro Asp Tyr Ala Leu Thr Val Thr Gly His Ser Leu Gly Ala Ser  
130 135 140

Leu Ala Ala Leu Thr Ala Ala Gln Leu Ser Ala Thr Tyr Asp Asn Ile  
145 150 155 160

Arg Leu Tyr Thr Phe Gly Glu Pro Arg Ser Asn Gln Ala Phe Ala Ser  
165 170 175

Tyr Met Asn Asp Ala Phe Gln Ala Ser Ser Pro Asp Thr Thr Gln Tyr  
180 185 190

Phe Arg Val Thr His Ala Asn Asp Gly Ile Pro Asn Leu Pro Pro Ala  
195 200 205

Asp Glu Gly Tyr Ala His Gly Val Val Glu Tyr Trp Ser Val Asp Pro  
210 215 220

Tyr Ser Ala Gln Asn Thr Phe Val Cys Thr Gly Asp Glu Val Gln Cys  
225 230 235 240

Cys Glu Ala Gln Gly Gly Gln Gly Val Asn Asn Ala His Thr Thr Tyr  
245 250 255

Phe Gly Met Thr Ser Gly His Cys Thr Trp  
260 265

<210> 9

<211> 276

<212> PRT

<213> Fusarium oxysporum

<400> 9

Ala Val Gly Val Thr Thr Thr Asp Phe Ser Asn Phe Lys Phe Tyr Ile  
1 5 10 15

Gln His Gly Ala Ala Ala Tyr Cys Asn Ser Glu Ala Ala Ala Gly Ser  
20 25 30

Lys Ile Thr Cys Ser Asn Asn Gly Cys Pro Thr Val Gln Gly Asn Gly  
35 40 45

Ala Thr Ile Val Thr Ser Phe Val Glv Ser Lvs Thr Glv Ile Glv Glv

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50              55              60
Tyr Val Ala Thr Asp Ser Ala Arg Lys Glu Ile Val Val Ser Phe Arg
65              70              75              80
Gly Ser Ile Asn Ile Arg Asn Trp Leu Thr Asn Leu Asp Phe Gly Gln
85              90              95
Glu Asp Cys Ser Leu Val Ser Gly Cys Gly Val His Ser Gly Phe Gln
100             105             110
Arg Ala Trp Asn Glu Ile Ser Ser Gln Ala Thr Ala Ala Val Ala Ser
115             120             125
Ala Arg Lys Ala Asn Pro Ser Phe Asn Val Ile Ser Thr Gly His Ser
130             135             140
Leu Gly Gly Ala Val Ala Val Leu Ala Ala Ala Asn Leu Arg Val Gly
145             150             155             160
Gly Thr Pro Val Asp Ile Tyr Thr Tyr Gly Ser Pro Arg Val Gly Asn
165             170             175
Ala Gln Leu Ser Ala Phe Val Ser Asn Gln Ala Gly Gly Glu Tyr Arg
180             185             190
Val Thr His Ala Asp Asp Pro Val Pro Arg Leu Pro Pro Leu Ile Phe
195             200             205
Gly Tyr Arg His Thr Thr Pro Glu Phe Trp Leu Ser Gly Gly Gly Gly
210             215             220
Asp Lys Val Asp Tyr Thr Ile Ser Asp Val Lys Val Cys Glu Gly Ala
225             230             235             240
Ala Asn Leu Gly Cys Asn Gly Gly Thr Leu Gly Leu Asp Ile Ala Ala
245             250             255
His Leu His Tyr Phe Gln Ala Thr Asp Ala Cys Asn Ala Gly Gly Phe
260             265             270
Ser Trp Arg Arg
275
<210> 10
<211> 273
<212> PRT
<213> Fusarium heterosporum
<400> 10
Thr Val Thr Thr Gln Asp Leu Ser Asn Phe Arg Phe Tyr Leu Gln His
1              5              10             15
Ala Asp Ala Ala Tyr Cys Asn Phe Asn Thr Ala Val Gly Lys Pro Val
20             25             30

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His Cys Ser Ala Gly Asn Cys Pro Asp Ile Glu Lys Asp Ala Ala Ile  
35 40 45

Val Val Gly Ser Val Val Gly Thr Lys Thr Gly Ile Gly Ala Tyr Val  
50 55 60

Ala Thr Asp Asn Ala Arg Lys Glu Ile Val Val Ser Val Arg Gly Ser  
65 70 75 80

Ile Asn Val Arg Asn Trp Ile Thr Asn Phe Asn Phe Gly Gln Lys Thr  
85 90 95

Cys Asp Leu Val Ala Gly Cys Gly Val His Thr Gly Phe Leu Asp Ala  
100 105 110

Trp Glu Glu Val Ala Ala Asn Val Lys Ala Ala Val Ser Ala Ala Lys  
115 120 125

Thr Ala Asn Pro Thr Phe Lys Phe Val Val Thr Gly His Ser Leu Gly  
130 135 140

Gly Ala Val Ala Thr Ile Ala Ala Ala Tyr Leu Arg Lys Asp Gly Phe  
145 150 155 160

Pro Phe Asp Leu Tyr Thr Tyr Gly Ser Pro Arg Val Gly Asn Asp Phe  
165 170 175

Phe Ala Asn Phe Val Thr Gln Gln Thr Gly Ala Glu Tyr Arg Val Thr  
180 185 190

His Gly Asp Asp Pro Val Pro Arg Leu Pro Pro Ile Val Phe Gly Tyr  
195 200 205

Arg His Thr Ser Pro Glu Tyr Trp Leu Asn Gly Gly Pro Leu Asp Lys  
210 215 220

Asp Tyr Thr Val Thr Glu Ile Lys Val Cys Glu Gly Ile Ala Asn Val  
225 230 235 240

Met Cys Asn Gly Gly Thr Ile Gly Leu Asp Ile Leu Ala His Ile Thr  
245 250 255

Tyr Phe Gln Ser Met Ala Thr Cys Ala Pro Ile Ala Ile Pro Trp Lys  
260 265 270

Arg

<210> 11

<211> 278

<212> PRT

<213> Aspergillus oryzae

<400> 11

Asp Ile Pro Thr Thr Gln Leu Glu Asp Phe Lys Phe Trp Val Gln Tyr

<211> 278

<211> 278

&lt;212&gt; PRT

<213> *Penicillium camemberti*

&lt;400&gt; 12

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Asp Val Ser Thr Ser Glu Leu Asp Gln Phe Glu Phe Trp Val Gln Tyr
1          5          10          15

Ala Ala Ala Ser Tyr Tyr Glu Ala Asp Tyr Thr Ala Gln Val Gly Asp
20          25          30

Lys Leu Ser Cys Ser Lys Gly Asn Cys Pro Glu Val Glu Ala Thr Gly
35          40          45

Ala Thr Val Ser Tyr Asp Phe Ser Asp Ser Thr Ile Thr Asp Thr Ala
50          55          60

Gly Tyr Ile Ala Val Asp His Thr Asn Ser Ala Val Val Leu Ala Phe
65          70          75          80

Arg Gly Ser Tyr Ser Val Arg Asn Trp Val Ala Asp Ala Thr Phe Val
85          90          95

His Thr Asn Pro Gly Leu Cys Asp Gly Cys Leu Ala Glu Leu Gly Phe
100         105         110

Trp Ser Ser Trp Lys Leu Val Arg Asp Asp Ile Ile Lys Glu Leu Lys
115         120         125

Glu Val Val Ala Gln Asn Pro Asn Tyr Glu Leu Val Val Val Gly His
130         135         140

Ser Leu Gly Ala Ala Val Ala Thr Leu Ala Ala Thr Asp Leu Arg Gly
145         150         155         160

Lys Gly Tyr Pro Ser Ala Lys Leu Tyr Ala Tyr Ala Ser Pro Arg Val
165         170         175

Gly Asn Ala Ala Leu Ala Lys Tyr Ile Thr Ala Gln Gly Asn Asn Phe
180         185         190

Arg Phe Thr His Thr Asn Asp Pro Val Pro Lys Leu Pro Leu Leu Ser
195         200         205

Met Gly Tyr Val His Val Ser Pro Glu Tyr Trp Ile Thr Ser Pro Asn
210         215         220

Asn Ala Thr Val Ser Thr Ser Asp Ile Lys Val Ile Asp Gly Asp Val
225         230         235         240

Ser Phe Asp Gly Asn Thr Gly Thr Gly Leu Pro Leu Leu Thr Asp Phe
245         250         255

Glu Ala His Ile Trp Tyr Phe Val Gln Val Asp Ala Gly Lys Gly Pro
260         265         270

Gly Leu Pro Phe Lys Arg

```

275

&lt;210&gt; 13

&lt;211&gt; 270

&lt;212&gt; PRT

&lt;213&gt; Aspergillus foetidus

&lt;400&gt; 13

Ser Val Ser Thr Ser Thr Leu Asp Glu Leu Gln Leu Phe Ala Gln Trp  
1 5 10 15

Ser Ala Ala Ala Tyr Cys Ser Asn Asn Ile Asp Ser Lys Asp Ser Asn  
20 25 30

Leu Thr Cys Thr Ala Asn Ala Cys Pro Ser Val Glu Glu Ala Ser Thr  
35 40 45

Thr Met Leu Leu Glu Phe Asp Leu Thr Asn Asp Phe Gly Gly Thr Ala  
50 55 60

Gly Phe Leu Ala Ala Asp Asn Thr Asn Lys Arg Leu Val Val Ala Phe  
65 70 75 80

Arg Gly Ser Ser Thr Ile Glu Asn Trp Ile Ala Asn Leu Asp Phe Ile  
85 90 95

Leu Glu Asp Asn Asp Asp Leu Cys Thr Gly Cys Lys Val His Thr Gly  
100 105 110

Phe Trp Lys Ala Trp Glu Ser Ala Ala Asp Glu Leu Thr Ser Lys Ile  
115 120 125

Lys Ser Ala Met Ser Thr Tyr Ser Gly Tyr Thr Leu Tyr Phe Thr Gly  
130 135 140

His Ser Leu Gly Gly Ala Leu Ala Thr Leu Gly Ala Thr Val Leu Arg  
145 150 155 160

Asn Asp Gly Tyr Ser Val Glu Leu Tyr Thr Tyr Gly Cys Pro Arg Ile  
165 170 175

Gly Asn Tyr Ala Leu Ala Glu His Ile Thr Ser Gln Gly Ser Gly Ala  
180 185 190

Asn Phe Arg Val Thr His Leu Asn Asp Ile Val Pro Arg Val Pro Pro  
195 200 205

Met Asp Phe Gly Phe Ser Gln Pro Ser Pro Glu Tyr Trp Ile Thr Ser

210

215

220

Gly Asn Gly Ala Ser Val Thr Ala Ser Asp Ile Glu Val Ile Glu Gly  
225 230 235 240

Ile Asn Ser Thr Ala Gly Asn Ala Gly Glu Ala Thr Val Ser Val Leu  
245 250 255

Ala His Leu Trp Tyr Phe Phe Ala Ile Ser Glu Cys Leu Leu  
260 265 270

<210> 14

<211> 270

<212> PRT

<213> Aspergillus niger

<400> 14

Ser Val Ser Thr Ser Thr Leu Asp Glu Leu Gln Leu Phe Ser Gln Trp  
1 5 10 15

Ser Ala Ala Ala Tyr Cys Ser Asn Asn Ile Asp Ser Asp Asp Ser Asn  
20 25 30

Val Thr Cys Thr Ala Asp Ala Cys Pro Ser Val Glu Glu Ala Ser Thr  
35 40 45

Lys Met Leu Leu Glu Phe Asp Leu Thr Asn Asn Phe Gly Gly Thr Ala  
50 55 60

Gly Phe Leu Ala Ala Asp Asn Thr Asn Lys Arg Leu Val Val Ala Phe  
65 70 75 80

Arg Gly Ser Ser Thr Ile Lys Asn Trp Ile Ala Asp Leu Asp Phe Ile  
85 90 95

Leu Gln Asp Asn Asp Asp Leu Cys Thr Gly Cys Lys Val His Thr Gly  
100 105 110

Phe Trp Lys Ala Trp Glu Ala Ala Ala Asp Asn Leu Thr Ser Lys Ile  
115 120 125

Lys Ser Ala Met Ser Thr Tyr Ser Gly Tyr Thr Leu Tyr Phe Thr Gly  
130 135 140

His Ser Leu Gly Gly Ala Leu Ala Thr Leu Gly Ala Thr Val Leu Arg  
145 150 155 160

Asn Asp Gly Tyr Ser Val Glu Leu Tyr Thr Tyr Gly Cys Pro Arg Val  
165 170 175

Gly Asn Tyr Ala Leu Ala Glu His Ile Thr Ser Gln Gly Ser Gly Ala  
180 185 190

Asn Phe Pro Val Thr His Leu Asn Asp Ile Val Pro Arg Val Pro Pro  
195 200 205

Met Asp Phe Gly Phe Ser Gln Pro Ser Pro Glu Tyr Trp Ile Thr Ser  
210 215 220

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      ---          ---          ---
Gly Thr Gly Ala Ser Val Thr Ala Ser Asp Ile Glu Leu Ile Glu Gly
225                               230                               235                               240

Ile Asn Ser Thr Ala Gly Asn Ala Gly Glu Ala Thr Val Asp Val Leu
                               245                               250                               255

Ala His Leu Trp Tyr Phe Phe Ala Ile Ser Glu Cys Leu Leu
                               260                               265                               270

<210> 15
<211> 269
<212> PRT
<213> Aspergillus oryzae

<400> 15
Asp Val Ser Ser Ser Leu Leu Asn Asn Leu Asp Leu Phe Ala Gln Tyr
1          5          10          15

Ser Ala Ala Ala Tyr Cys Asp Glu Asn Leu Asn Ser Thr Gly Thr Lys
          20          25          30

Leu Thr Cys Ser Val Gly Asn Cys Pro Leu Val Glu Ala Ala Ser Thr
          35          40          45

Gln Ser Leu Asp Glu Phe Asn Glu Ser Ser Ser Tyr Gly Asn Pro Ala
50          55          60

Gly Tyr Leu Ala Ala Asp Glu Thr Asn Lys Leu Leu Val Leu Ser Phe
65          70          75          80

Arg Gly Ser Ala Asp Leu Ala Asn Trp Val Ala Asn Leu Asn Phe Gly
          85          90          95

Leu Glu Asp Ala Ser Asp Leu Cys Ser Gly Cys Glu Val His Ser Gly
          100          105          110

Phe Trp Lys Ala Trp Ser Glu Ile Ala Asp Thr Ile Thr Ser Lys Val
          115          120          125

Glu Ser Ala Leu Ser Asp His Ser Asp Tyr Ser Leu Val Leu Thr Gly
130          135          140

His Ser Tyr Gly Ala Ala Leu Ala Ala Leu Ala Ala Thr Ala Leu Arg
145          150          155          160

Asn Ser Gly His Ser Val Glu Leu Tyr Asn Tyr Gly Gln Pro Arg Leu
          165          170          175

Gly Asn Glu Ala Leu Ala Thr Tyr Ile Thr Asp Gln Asn Lys Gly Gly
          180          185          190

Asn Tyr Arg Val Thr His Thr Asn Asp Ile Val Pro Lys Leu Pro Pro

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195              200              205

Thr Leu Leu Gly Tyr His His Phe Ser Pro Glu Tyr Tyr Ile Ser Ser
210              215              220

Ala Asp Glu Ala Thr Val Thr Thr Thr Asp Val Thr Glu Val Thr Gly
225              230              235              240

Ile Asp Ala Thr Gly Gly Asn Asp Gly Thr Asp Gly Thr Ser Ile Asp
245              250              255

Ala His Arg Trp Tyr Phe Ile Tyr Ile Ser Glu Cys Ser
260              265

<210> 16
<211> 251
<212> PRT
<213> Landerina penisapora

<400> 16
Pro Gln Asp Ala Tyr Thr Ala Ser His Ala Asp Leu Val Lys Tyr Ala
1      5      10      15

Thr Tyr Ala Gly Leu Ala Tyr Gln Thr Thr Asp Ala Trp Pro Ala Ser
20     25     30

Arg Thr Val Pro Lys Asp Thr Thr Leu Ile Ser Ser Phe Asp His Thr
35     40     45

Leu Lys Gly Ser Ser Gly Tyr Ile Ala Phe Asn Glu Pro Cys Lys Glu
50     55     60

Ile Ile Val Ala Tyr Arg Gly Thr Asp Ser Leu Ile Asp Trp Leu Thr
65     70     75     80

Asn Leu Asn Phe Asp Lys Thr Ala Trp Pro Ala Asn Ile Ser Asn Ser
85     90     95

Leu Val His Glu Gly Phe Leu Asn Ala Tyr Leu Val Ser Met Gln Gln
100    105    110

Val Gln Glu Ala Val Asp Ser Leu Leu Ala Lys Cys Pro Asp Ala Thr
115    120    125

Ile Ser Phe Thr Gly His Ser Leu Gly Gly Ala Leu Ala Cys Ile Ser
130    135    140

Met Val Asp Thr Ala Gln Arg His Arg Gly Ile Lys Met Gln Met Phe
145    150    155    160

Thr Tyr Gly Gln Pro Arg Thr Gly Asn Gln Ala Phe Ala Glu Tyr Val
165    170    175

Glu Asn Leu Glv His Pro Val Phe Arg Val Val Trp Arg His Asp Ile

```

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-----
      180              185              190
Val  Pro  Arg  Met  Pro  Pro  Met  Asp  Leu  Gly  Phe  Gln  His  His  Gly  Gln
      195              200              205

Glu  Val  Trp  Tyr  Glu  Gly  Asp  Glu  Asn  Ile  Lys  Phe  Cys  Lys  Gly  Glu
      210              215              220

Gly  Glu  Asn  Leu  Thr  Cys  Glu  Leu  Gly  Val  Pro  Phe  Ser  Glu  Leu  Asn
      225              230              235              240

Ala  Lys  Asp  His  Ser  Glu  Tyr  Pro  Gly  Met  His
      245              250

```

&lt;210&gt; 17

&lt;211&gt; 12

&lt;212&gt; PRT

&lt;213&gt; Artificial

&lt;220&gt;

&lt;223&gt; Sequence used for alignment

&lt;400&gt; 17

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Ala  Cys  Met  Ser  His  Thr  Trp  Gly  Glu  Arg  Asn  Leu
1              5              10

```

&lt;210&gt; 18

&lt;211&gt; 14

&lt;212&gt; PRT

&lt;213&gt; Artificial

&lt;220&gt;

&lt;223&gt; Sequence used for alignment

&lt;400&gt; 18

```

His  Gly  Trp  Gly  Glu  Asp  Ala  Asn  Leu  Ala  Met  Asn  Pro  Ser
1              5              10

```

## REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

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- **NEEDLEMAN, S. B.WUNSCH, C. D.**J. Mol. Biol., 1970, vol. 48, 443-453 [0059]
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**PATENTKRAV**

1. Variant af en stamlipase, hvor lipasevarianten er mindst 80% identisk med SEQ ID NO: 2 og omfatter mindst tre substitutioner valgt fra gruppen bestående af (under anvendelse af SEQ ID NO: 2 til nummerering):
  - 5 a) mindst to substitutioner i Region I og
  - b) mindst én substitution i Region II og
  - c) mindst én substitution i Region III og
  - d) mindst én substitution i Region IV;og hvor varianten har lipaseaktivitet og udviser forbedret vaskeevne og lugt og omfatter  
10 substitutionerne T231R +N233R +I255Y.
2. Lipasevariant ifølge krav 1, hvor lipasevarianten er mindst 85%, 90%, 95%, 96%, 97%, 98% eller 99% identisk med SEQ ID NO: 2.
3. Lipasevariant ifølge krav 1, hvor stamlipasen er lipasen med aminosyresekvensen ifølge SEQ ID NO: 2.
- 15 4. Lipasevariant ifølge krav 1, hvor lipasen omfatter en yderligere substitution i positionen svarende til position 4 og/eller position 227 ifølge SEQ ID NO: 2.
5. Lipasevariant ifølge krav 4, hvor lipasen har en substitution svarende til X4V og X227G (under anvendelse af SEQ ID NO: 2 til nummerering).
6. Lipasevariant ifølge krav 1, hvor den mindst ene substitution i stamlipasens Region II  
20 omfatter substitutioner valgt fra gruppen bestående af substitutioner i positioner svarende til positionerne 202, 210, 211 og 256 (under anvendelse af SEQ ID NO: 2 til nummerering).
7. Lipasevariant ifølge krav 6, hvor den mindst ene substitution i stamlipasen er valgt fra gruppen bestående af X202G, X210K, X211L og X256K (under anvendelse af SEQ ID NO: 2 til nummerering).
- 25 8. Lipasevariant ifølge krav 1, hvor den mindst ene substitution i stamlipasens Region III omfatter substitutioner valgt fra gruppen bestående af substitutioner i positioner svarende til positionerne 83, 86 og 90 (under anvendelse af SEQ ID NO: 2 til nummerering).

9. Lipasevariant ifølge krav 8, hvor den mindst ene substitution i stamlipasen er valgt fra gruppen bestående af X83T, X86V og X90A/R (under anvendelse af SEQ ID NO: 2 til nummerering).
10. Lipasevariant ifølge krav 1, hvor den mindst ene substitution i stamlipasens Region IV  
5 omfatter substitutioner valgt fra gruppen bestående af substitutioner i positioner svarende til positionerne 27, 58 og 60 (under anvendelse af SEQ ID NO: 2 til nummerering).
11. Lipasevariant ifølge krav 10, hvor den mindst ene substitution i stamlipasens Region IV omfatter substitutioner valgt fra gruppen bestående af X27R, X58N/A/G/P/T og X60S/V/G/N/R/K/A/L (under anvendelse af SEQ ID NO: 2 til nummerering).
- 10 12. Lipasevariant ifølge krav 1, hvor lipasevarianten endvidere omfatter mindst én substitution i stamlipasen valgt fra gruppen bestående af substitutioner i positioner svarende til position 81, 147, 150, 227 og 249 (under anvendelse af SEQ ID NO: 2 til nummerering).
13. Lipasevariant ifølge krav 12, hvor lipasevarianten endvidere omfatter mindst én substitution valgt fra gruppen bestående af X81Q/E, X147M/Y, X150G og X249R/I/L (under  
15 anvendelse af SEQ ID NO: 2 til nummerering).
14. DNA-sekvens, som koder for lipasevarianten ifølge krav 1-13.
15. Ekspressionsvektor, som huser DNA-sekvensen ifølge krav 14.
16. Transformeret værtselle, som indeholder DNA-sekvensen ifølge krav 14.
17. Fremgangsmåde til fremstilling af en lipasevariant, hvilken fremgangsmåde omfatter  
20 dyrkning af den transformerede værtselle ifølge krav 16 under betingelser, som er befordrende for fremstilling af lipasevarianten, og indvinding af lipasevarianten fra det resulterende medium.

## DRAWINGS

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ID NO 1: SSSSTQDYRIASEARIKAHTFYTALSANA
ID NO 2: SSSSTQDYRIASEARIKAHTFYTALSANA
ID NO 3: SIDGGIRAAATQOEINELTYTTLTANS
ID NO 4: SASDGGKVVAAATTAQIQEFTKYAGIAATA
ID NO 5: TAGHALAASTQ GISEDLYSKL VEMATISQAA
ID NO 6: TAGHALAASTQ GISEDLYSKL VEMATISQAA
ID NO 7: AVGVTTTDSFNFXYIQHGAAA
ID NO 8: TVTTQDLNFRFYLQHADAA
ID NO 9: DLTPTQLRDEKFWVQYAAAT
ID NO 10: DVSTSELQDQEFWVQYAAAS
ID NO 11: SVSTSTLDELQLFAQWSAAA
ID NO 12: SVSTSTLDELQLFSQWSAAA
ID NO 13: DVSSLLNNLDELFAQYGAAS
ID NO 14: BVSQDLFNQFNLFPAQYSAAA
ID NO 15: PQDAYTASHADLVKYATYAGLA

ID NO 1: YCRFVIFG GRWSCPHCGVAS NLQITKTPST LITDENVLVAV
ID NO 2: YCRFVIFG GQWSCPHCDVAP NLNITKTPST LITDENVLVAV
ID NO 3: YCRFVIFG ATWDCIHCDATE DLKIKITWST LIYDTNAMVAR
ID NO 4: YCRSVVFG NKWDCVQCQKWP DGKLIITPTS LLSDTNGYVLR
ID NO 5: YADLCNIPST IIKGKIYNSQTDINGWILR
ID NO 6: YADLCNIPST IIKGKIYNSQTDINGWILR
ID NO 7: YC NSRAAA GSKITCSNKCPTVQNGCATIVTSF VGSKTGIGGYVAT
ID NO 8: YC NFNIAV GKPVHCSAGNCPDIEKDAIVVGSV VGTKTGIGGYVAT
ID NO 9: YCPNNYVAKD GERLNCVGNCPDVEAGSTVKLSFS DDTITDTAGFVAV
ID NO 10: YYEADYTAQV GDKLSCSKGNCPVEEATGATVSYDFS DSTITDTAGYIAV
ID NO 11: YCSNNID SK DSNLTCTANACPSVEEASTTMLLEFDLTDFGCTAGFLAA
ID NO 12: YCSNNID SD DSNVTCIADACPSVEEASTKMLLEFDLTNNFGGTAGFLAA
ID NO 13: YCOENLN ST GTKLTCVGNCELVEAASTQSLDEFNBSSESYPAGYLA
ID NO 14: YCGKNNDAPA GTNITCTCNACPEVEKADATFLYSE DSGVGDVGTGFLAL
ID NO 15: YQTTDAPAS RTVPADTTLISFD HTLKGSSEGYIAF

ID NO 1: GEKERTIYVV FRGTSSIRNA IADIVFVPVN YPPV NGA KVHKGFLDSY
ID NO 2: GENEXTIYVV FRGTSSIRNA IADIVFVPVN YPPV NGA KVHKGFLDSY
ID NO 3: GDSKRTIYV FRGSSIRNW IADLTFVPVS YPPV SGT KVHKGFLDSY
ID NO 4: SDKOKTIYLV FRGTNSIRSA ITDIVNFSD YKPV KGA KVHAGFLSSY
ID NO 5: DSSSKRTIYV FRGTGSDTNL QLDITNYTLTP FDTLEQCNGC EVHGGYIOW
ID NO 6: DSSSKRTIYV FRGTGSDTNL QLDITNYTLTP FDTLEQCNGC EVHGGYIOW
ID NO 7: DSARKETIYVS FRGSINIRNW LTNLDPG QE DCSL VSGC GVHSGFQRAW
ID NO 8: DNARKETIYVS VRGSINIRNW ITNFNPG QK TCCL VAGC GVHTGFILDAW
ID NO 9: DNTNKAIYVA FRGSYSIRNW VTDATPE QT DPGL CDGC KABLGFWTAW
ID NO 10: DNTNSAVVLA FRGSYSVRNW VADATFV HT NPGL CDGC LABLGFWSW
ID NO 11: DNTNKRLVVA FRGSSTLENW IANLDFILED NDDL CTGC KVHTGFWKAW
ID NO 12: DNTNKRLVVA FRGSSTLENW IADLDFILED NDDL CTGC KVHTGFWKAW
ID NO 13: DETNKLVLVS FRGSADLANW VANLNFQLED ASDL CSQC EVHSGFVKAW
ID NO 14: DNTNKLIVLS FRGSRISIRNW IGMNFDLKE INDI CSQC RGHGDTSSW
ID NO 15: NEPCKRTIYV YRGTDSLIDW LTNLNFQKTA WPAW IENS LVHGFPLNAY

ID NO 1: NEVQDKLVAE VKAQLDRHPG YKIVVTGHSI GGATAVLASALDLYHHGHA
ID NO 2: NEVQDKLVAE VKAQLDRHPG YKIVVTGHSI GGATAVLASALDLYHHGHD
ID NO 3: GEVQNELVAT VLDQFKQYPS YKIVVTGHSI GGATALLCALDLYQREBGLS
ID NO 4: EQVVDYFPV VQQLTAFTPT YKIVVTGHSI GGAQALLAGMDLYQREPRLS
ID NO 5: VSVQDQVESL VKQQVSQYPD YALTVTGHSI GASLAALITAAQL SATYD
ID NO 6: ISVQDQVESL VQQVVSQYPD YALTVTGHSI GASLAALITAAQL SATYD
ID NO 7: NEISSQATAA VASARKANPS ENVISTGHSI GGAVAVLAAANLVRGGT
ID NO 8: EVVAANVKAA VSAAKTANPT FKEVVVTGHSI GGAVATTAAAYLRKDG
ID NO 9: KVVVRITIKT LDELKEEESD YKIVVTGHSI GAAIASLAAADLRKNY
ID NO 10: KLVVRDITIKT LKEVVVAQNP YELVVVTGHSI GAAVATLAATDLRGKGY
ID NO 11: ESAADELTISK IKSAMSTYSG YTLVVTGHSI GGALATLGATVLRNDGY
ID NO 12: BAAADNLTISK IKSAMSTYSG YTLVVTGHSI GGALATLGATVLRNDGY
ID NO 13: SELADTITISK VEGALSDESD YSLVVTGHSI GAALAALAATAALRNSGH

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Figure 1 (cont.)

ID NO 14: RSVADTLRCK VEDAVREHPD YRVVFTCHSL GGALATVACADLRGNGY  
ID NO 15: LVSMQVQQA VDSLLAKCPD ATISFTCHSL GGALACISMVDTAQRHGI

ID NO 1: NIBIYTOG QPRIGTPAPA NVVIGT KIPYQRLVNERDIVPHL  
ID NO 2: NIBIYTOG QPRIGTPAPA NVVIGT KIPYQRLVNERDIVPHL  
ID NO 3: SSNLELYTOG QPRVGDPAFA NVVIST GIPYRRTVNERDIVPHL  
ID NO 4: PKNLSITFTG QPRVGNPAPA YVVEST GIPYQRTVHKRDIVPHV  
ID NO 5: NIRLYTFC EPRSNGQAPA SYMMDAFQASSPDTOYFRVTHANDGIPNL  
ID NO 6: NIRLYTFC EPRS NQAPA SYMMDAFQASSPDTOYFRVTHANDGIPNL  
ID NO 7: FVDIYTYG SPRVGNQALS AFVENQ AGGEYRVTHADDPVRL  
ID NO 8: PFDLYTYG SPRVGNDFPA NEVTQQ TGAERYVTHGDDPVRL  
ID NO 9: DAILYAYA APRVANKPLA EPITNQ GNNYRPTHNDPPVRL  
ID NO 10: SAKLYAYA SPRVGNAAALA KYITAO GNNFRPTHNDPPVRL  
ID NO 11: SVELYTYG CTRVGNAYALA EHITSQ GSGANFRVTHLNDIVPRV  
ID NO 12: SVELYTYG CTRVGNAYALA EHITSQ GSGANFRVTHLNDIVPRV  
ID NO 13: SVELYTYG QPRLGNEALA TYITDQ NKGNNYRVTHNDIVPRL  
ID NO 14: DIDVFSYG APRVGNKAPA EPLTVQ TGGTLVRIHTNDIVPRL  
ID NO 15: KMQMFTYG QPRTGNQAPA EYVENL GRPVFRVVYRHDIVPRM

ID NO 1: PPGAGFLHA GEEFWIMK DSSLRVCPNGIETDNCNSIV  
ID NO 2: PPGAGFLHA GEEFWIMK DSSLRVCPNGIETDNCNSIV  
ID NO 3: PPAAGFLHA GEEFWITD NSPETVQVCTSDLETSDCNSIV  
ID NO 4: PPQSGFLHP GVEYWSK GTSNVQICTSELETSDCNSIV  
ID NO 5: PPVQGYANG GVEYWSV DPYSAQNTFVCTGDEVQCCB AQGGQG  
ID NO 6: PPADGXYANG VVEYWSV DPYSAQNTFVCTGDEVQCCB AQGGQG  
ID NO 7: PPLIFGYRHT TDPFWLSGGGGKVDYTI SDVKVCSGAALC CNGGTL  
ID NO 8: PPIVGYRHT SPEYWLNG GFLDRDYTVTEIKVCEGIANVM CNGGTI  
ID NO 9: PLLMGYVHI SPEYVITA PDNTVTDNQVTVLDGYVNFK GNTGTS  
ID NO 10: PLLMGYVHV SPEYWITS PNNATVSTSDIKVIDGVSFD GNTGTS  
ID NO 11: PPMDFGSGP SPEYWITS GNGASVTASDIEVLEGINSTA CNAGEA  
ID NO 12: PPMDFGSGP SPEYWITS GNGASVTASDIEVLEGINSTA CNAGEA  
ID NO 13: PPTLLGYHMF SPEYVLS ADEATVTTDVTEVTGIDATG GNDGTD  
ID NO 14: PPREFGYHS SPEYWIKS GTLVPVTRNDIVKIEGIDATG GNNQPN  
ID NO 15: PPMDLGFQHI GQEVWYEG DENIKFCCKEGENLTCELVF

ID NO 1: PFT SVIDHLSYLDMENTOL CL  
ID NO 2: PFT SVIDHLSYLDMENTOL CL  
ID NO 3: PFT SVIDHLSYFGINTOL CT  
ID NO 4: PFT SVIDHLSYEDINEGS CL  
ID NO 5: VN NAHTTYF GNTSGACTW  
ID NO 6: VN NAHTTYF GNTSGACTW  
ID NO 7: CL DIAHILHYF QATDA CNAQCFWR R  
ID NO 8: CL DILAHITYF QSMAT CAPIALPWK R  
ID NO 9: GGLPDLAPFESHVWYFIHADACKGGLPLR  
ID NO 10: LPLLDGFBHIVYF VQVDA GKGPGLPK R  
ID NO 11: TV SVLAHLWYF FAISE CLL  
ID NO 12: TV DVLHLWYF FAISE CLL  
ID NO 13: CT SIDAHRWYF TYISE CS  
ID NO 14: IP DIPAHILWYF GLIGT CL  
ID NO 15: FSEL NAKDESBYF GMH

| ID NO: | Micro organism                  | SEQ ID NO: |
|--------|---------------------------------|------------|
| 1.     | <i>Absidia reflexa</i>          | 3          |
| 2.     | <i>Absidia corymbifera</i>      | 4          |
| 3.     | <i>Rhizomucor miehei</i>        | 5          |
| 4.     | <i>Rhizopus delema (oryzae)</i> | 6          |
| 5.     | <i>Aspergillus niger</i>        | 7          |
| 6.     | <i>Aspergillus tubingensis</i>  | 8          |
| 7.     | <i>Fusarium oxysporum</i>       | 9          |
| 8.     | <i>Fusarium heterosporum</i>    | 10         |
| 9.     | <i>Aspergillus oryzae</i>       | 11         |
| 10.    | <i>Penicillium camemberti</i>   | 12         |

Figure 1 (cont.)

|     |                                |    |
|-----|--------------------------------|----|
| 11. | <i>Aspergillus foetidus</i>    | 13 |
| 12. | <i>Aspergillus niger</i>       | 14 |
| 13. | <i>Aspergillus oryzae</i>      | 15 |
| 14. | <i>Thermomyces lanuginosus</i> | 2  |
| 15. | <i>Landerina penisapora</i>    | 16 |

Figure 1. Alignment of lipase sequences.

Figure 1 (cont.)