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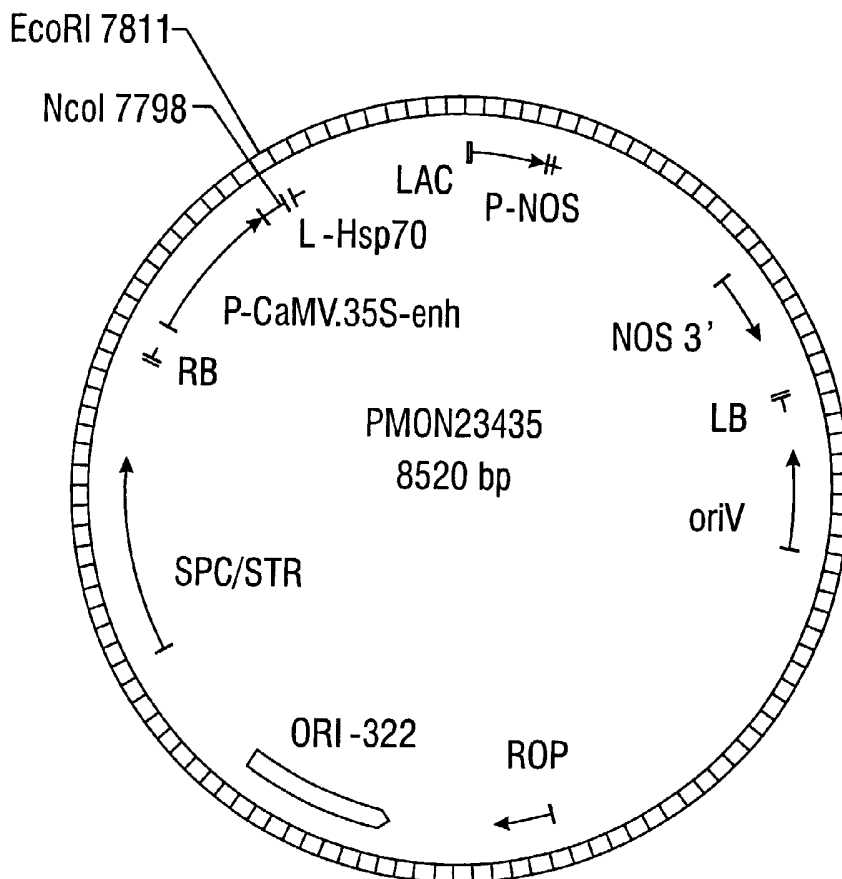
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(54) Title: METHOD OF INCREASING PLANT ORGAN AND SEED SIZE IN A PLANT



(57) Abstract: The present invention provides methods and DNA molecules useful for producing transgenic plants with agronomically desired traits based on altered sizes of plant organs and tissues.



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METHOD OF INCREASING PLANT ORGAN AND SEED SIZE IN A PLANT5 **CROSS REFERENCE TO RELATED APPLICATIONS**

This Application claims the benefit of U.S. Provisional Application No.60/381,100, filed 5/15/2002.

FIELD OF THE INVENTION

10 This invention relates to plant molecular biology. In particular, this invention relates to transgenic plants having increased or enhanced seed and organ sizes.

BACKGROUND OF THE INVENTION

Recent advances in genetic engineering have provided the prerequisite tools to transform
15 plants to contain foreign (often referred to as "heterogeneous or heterologous ") or improved endogenous genes. The introduction of such a gene in a plant can desirably lead to an improvement of an already existing pathway in plant tissues or introduction of a novel pathway to modify desired product levels, increase metabolic efficiency, and/or save on energy cost to the cell. Plants with unique physiological and biochemical traits and characteristics, such as
20 herbicide resistance and insect resistance, have already been produced. The ability to create traits that play an essential role in plant growth and development, crop yield potential and stability, and crop quality and composition are particularly desirable targets for the crop plant improvement.

Normally a plant goes through a development cycle, which includes seed germination,
25 maturation of plant, reproduction, and finally senescence that leads to death of a plant. Several biological processes are common to different stages of plant development. Desired effects such as growth of tissue organ are achieved in nature by fine-tuning of the metabolism of the organism. The final phase of growth is senescence which is a highly regulated, genetically controlled and active process (Thomas H., and Stoddart J. L., Ann. Rev. Plant Physiol (31) 83-
30 111, 1980). Senescence is mostly studied in plant leaves and is regarded as a series of events concerned with cellular disassembly and the mobilization of released material to other plant parts

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such as seeds, storage organs or developing leaves and flowers (Nooden L.D. In *Senescence and Aging in Plants*, Academic press, 391-439, 1988). Leaf senescence can be initiated by seed development in certain species of plants. This was demonstrated in soybean by surgically removing flowers or physically restricting pod growth to observe the delay in leaf senescence (Nooden L.D. In *Senescence and Aging in Plants*, Academic press, 330-368, 1988; Miceli F, Crafts-Brandner S.J., Egli D.B. *Crop Sci.* (35), 1080-1085, 1995). During senescence, partitioning of resources between vegetative and reproductive development involves a complex interplay of generative and degenerative processes, requiring differential expression of genes.

Differentially expressed genes during senescence are usually referred as "Senescence Associated Genes" or SAGs (Hensel L.L, Grbic V., Baumgarten D.A., Bleecker A.B., *The Plant Cell* (5) 553-564 1993). All SAG genes may not be functionally related, but they all are involved in similar physiological processes. In the past, senescence studies were directed towards understanding processes to generally enhance knowledge and applying this information relating to senescence in agriculture to enhancing yield and reducing post harvest losses (Hensel L.L, Grbic V., Baumgarten D.A., Bleecker A.B., *The Plant Cell* (5) 553-564 1993; Gan S., Amasino R.M., *Science*, (270) 1986-1988, 1995; Gan S., Amasino R.M., *Plant Physiol.*, (113) 313-319, 1997; Guarente L., Ruvkun G., Amasino R.M., *Proc. Natl Acad. Sci USA* (95) 1034-1036, 1998; US Patent Number 5 68 9042; PCT/US00/03494; and PCT/US00/18364, July 2000).

The SAG 13 gene was first described by Lohman et al. in 1994 (Lohman K.M., Gan S., John M.C., Amasino R.M., *Physiologia Plantarum* (92) 322-328, 1994) and then by Weaver et al. in 1998 (Weaver L.M. Gan S., Quirino B, Amasino R.M., *Plant Mol. Biol.* 455-469, 1998) as one of the genes associated with senescence. SAG genes by definition are up regulated during age-mediated senescence. SAG 13 was observed to be induced strongly shortly before visible senescence marked by yellowing of green leaves.

SUMMARY OF THE INVENTION

The present invention relates to transgenic plants with increased organ size when compared to a non transformed plant of the same species. In one preferred embodiment, the present invention provides a transgenic plant with increased seed size when compared to a non transformed plant of the same species.

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One embodiment of the invention provides a method of increasing seed and organ size of a plant by transforming the plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, where in said DNA molecule is selected from the from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, 5
SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, and SEQ ID NO: 70 and wherein 10
said promoter is heterologous to said DNA molecule. The method also includes the DNA construct operably linked to a 3' termination region and selecting a desired plant from a population of transformed plants containing said DNA construct; wherein said desired plant exhibits increased seed and organ size compared to a plant of a same plant species not transformed to contain said DNA construct. The method may also encompass various promoters 15
including a caulimovirus promoter, or a heterologous plant constitutive promoter, or a tissue specific or an organ enhanced promoter.

In another embodiment, the present invention provides a method for increasing seed and other plant organ sizes by transforming a plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, wherein said 20
protein comprises at least an N-terminal 50% portion of a polypeptide selected from the group consisting of : SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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, 25
SEQ ID NO: 67, SEQ ID NO: 69, and SEQ ID NO: 71 wherein said DNA construct is operably linked to a 3' termination region; and selecting a plant from a population of transformed plants containing said DNA construct; wherein said plant exhibits increased seed and organ size compared to a plant of a same species not transformed to contain said DNA construct. In a 30
further embodiment, the present invention provides an isolated DNA construct comprising a promoter capable of functioning in a plant cell, operably linked to a structural nucleic acid

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sequence encoding SEQ ID NO: 2 or SEQ ID NO: 4, and a 3' non translated nucleic acid sequence capable of causing transcriptional termination and the addition of polyadenylated nucleotides to the 3' end of the transcribed mRNA sequence. This nucleic acid sequence may optionally include an intron, a 5' untranslated leader sequence or another nucleic acid sequence
5 designed to enhance transcription and/or translation.

Another aspect of the present invention provides a method for improving the seed and organ size in a plant comprising the steps of:

(a) inserting into the genome of a plant a DNA construct comprising in the 5' to 3' direction:

- 10 (i) a promoter that functions in the cell of a selected plant tissue operably linked to;
- (ii) a structural nucleic acid sequence that encodes a protein product selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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
15 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71 operably linked to
- (iii) a 3' non translated nucleic acid sequences that functions in a plant cell to cause transcriptional termination and addition of polyadenylated nucleotides to the 3' end of an RNA
20 sequence;
- (b) obtaining transformed plant cells containing the nucleic acid of step (a); and
- (c) regenerating from transformed plant cells a transformed plant that over-expresses the gene product in the plant cell.

One aspect of invention provides a transgenic plant with increased seed and organ size
25 when compared to a non transformed plant of a same plant species, said transgenic plant comprising a DNA construct, wherein said DNA construct comprises a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, wherein said DNA molecule is selected from the group consisting of: SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, SEQ ID NO: 9, SEQ ID NO: 11, SEQ ID NO: 13, SEQ ID NO: 15, SEQ ID
30 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

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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: 66, SEQ ID NO: 68, and SEQ ID NO: 70 operably linked to a 3' termination region, wherein said promoter is heterologous to said DNA molecules.

5 Another aspect of invention provides a transgenic plant with increased seed and organ size when compared to a non transformed plant of a same plant species, the transgenic plant comprising a DNA construct, wherein said DNA construct encodes a protein that comprises at least an N-terminal 50% portion selected from the group consisting of: SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71.

15 Yet another aspect of the present invention provides a DNA construct comprising a promoter operably linked to an isolated nucleic acid molecule comprising a nucleotide sequence or complement thereof, wherein the nucleotide sequence encodes a polypeptide having an amino acid sequence that has at least 50% sequence identity, preferably at least 60%, more preferably at least 70% sequence identity, even more preferably at least 80% or 90% sequence identity, and most preferably at least 95% to 98% sequence identity to a polypeptide selected from group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, SEQ ID NO: 71 where in said promoter is heterologous to the said nucleic acid molecules.

BRIEF DESCRIPTION OF THE FIGURES

30 **Figure 1** shows a plasmid map for plant transformation vector pMON 23435 with its elements.

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Figure 2 shows a plasmid map for plant transformation vector pMON 57521 with its elements.

Figure 3 shows a plasmid map for plant transformation vector pMON 54955 with its elements.

5 **Figure 4** shows a plasmid map for plant transformation vector pMON 73955 with its elements.

Figure 5 shows a plasmid map for plant transformation vector pMON 73963 with its elements.

10 **Figure 6** shows a plasmid map for bacterial transformation vector pMON 63132 with its elements.

Figure 7 shows a plasmid map for bacterial transformation vector pMON 63133 with its elements.

Figure 8 shows a plasmid map for bacterial transformation vector pMON 6134 with its elements.

15 **Figure 9** shows a plasmid map for bacterial transformation vector pMON 6135 with its elements.

Figure 10 shows primers used for amplification of gene and for sequencing of amplicons.

Figure 11 shows putative products from steroid reductase activity.

Figure 12 shows a potential steroid reductase reaction.

20

DETAILED DESCRIPTION OF THE INVENTION

Isolated Nucleic Acid Molecules of the Present Invention

25 The present invention provides a method of increasing seed and organ size of a plant by transforming the plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, where in said DNA molecule is selected from the from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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
30 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

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ID NO: 51, SEQ ID NO: 53, SEQ ID NO: 66, SEQ ID NO: 68, SEQ ID NO: 70 and wherein said promoter is heterologous to said DNA molecule.

In another embodiment, the present invention provides a method for increasing seed and other plant organ sizes by transforming a plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, wherein said protein comprises at least an N-terminal 50% portion of a polypeptide selected from the group consisting of : SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, or SEQ ID NO: 71 operably linked to a 3' termination region.

The term "nucleic acid molecule" as used herein means a deoxyribonucleic acid (DNA) molecule or ribonucleic acid (RNA) molecule. Both DNA and RNA molecules are constructed from nucleotides linked end to end, wherein each of the nucleotides contains a phosphate group, a sugar moiety, and either a purine or a pyrimidine base. Nucleic acid molecules can be a single or double-stranded polymer of nucleotides read from the 5' to the 3' end. Nucleic acid molecules may also optionally contain synthetic, non-natural or altered nucleotide bases that permit correct read through by a polymerase and do not alter expression of a polypeptide encoded by that nucleic acid molecule.

The term "an isolated nucleic acid molecule" as used herein means a nucleic acid molecule that is no longer accompanied by some of materials with which it is associated in its natural state or to a nucleic acid molecule the structure of which is not identical to that of any of naturally occurring nucleic acid molecule. Examples of an isolated nucleic acid molecule include: (1) DNAs which have the sequence of part of a naturally occurring genomic DNA molecule but are not flanked by two coding sequences that flank that part of the molecule in the genome of the organism in which it naturally occurs; (2) a nucleic acid molecule incorporated into a vector or into the genomic DNA of a prokaryote or eukaryote in a manner such that the resulting molecule is not identical to any naturally occurring vector or genomic DNA; (3) a separate molecule such as a cDNA, a genomic fragment, a fragment produced by polymerase chain reaction (PCR), or a restriction fragment; (4) recombinant DNAs; and (5) synthetic DNAs.

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An isolated nucleic acid molecule may also be comprised of one or more segments of cDNA, genomic DNA or synthetic DNA.

It is also contemplated by the inventors that the isolated nucleic acid molecules of the present invention also include known types of modifications, for example, labels which are known in the art, methylation, "caps", substitution of one or more of the naturally occurring nucleotides with an analog. Other known modifications include internucleotide modifications, for example, those with uncharged linkages (methyl phosphonates, phosphotriesters, phosphoamidates, carbamates, etc.) and with charged linkages (phosphorothioates, phosphorodithioates, etc.), those containing pendant moieties, such as, proteins (including nucleases, toxins, antibodies, signal peptides, poly-L-lysine, etc.), those with intercalators (acridine, psoralen, etc.), those containing chelators (metals, radioactive metals, boron, oxidative metals, etc.), those containing alkylators, and those with modified linkages.

The term "nucleotide sequence" as used herein means the order contiguous of nucleic acid molecule of both the sense and antisense strands or as a duplex strand. It includes, but is not limited to, self-replicating plasmids, synthetic polynucleotides, chromosomal sequences, and infectious polymers of DNA or RNA.

A nucleotide sequence is said to be the "complement" of another nucleotide sequence if they exhibit complete complementarity. As used herein, molecules are said to exhibit "complete complementarity" when every nucleotide of one of the sequences is complementary to a nucleotide of the other.

As used herein both terms "a coding sequence" and "a structural nucleotide sequence" mean a nucleotide sequence, which is translated into a polypeptide, usually via mRNA, when placed under the control of appropriate regulatory sequences. The boundaries of the coding sequence are determined by a translation start codon at the 5'-terminus and a translation stop codon at the 3'-terminus. A coding sequence can include, but is not limited to, genomic DNA, cDNA, and recombinant nucleotide sequences.

The polypeptides of the invention, like other polypeptides, have different domains which perform different functions. Thus, the coding sequences need not be full length, so long as the desired functional domain of the polypeptide is expressed. The distinguishing features of polypeptides of the present invention are discussed in detail in Examples.

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The term "recombinant DNAs" as used herein means DNAs that contains a genetically engineered modification through manipulation via mutagenesis, restriction enzymes, and the like.

The term "synthetic DNAs" as used herein means DNAs assembled from oligonucleotide building blocks that are chemically synthesized using procedures known to those skilled in the art. These building blocks are ligated and annealed to form DNA segments, which are then enzymatically assembled to construct the entire DNA. "Chemically synthesized", as related to a sequence of DNA, means that the component nucleotides were assembled in vitro. Manual chemical synthesis of DNA may be accomplished using well-established procedures, or automated chemical synthesis can be performed using one of a number of commercially available machines.

Both terms "polypeptide" and "protein", as used herein, mean a polymer composed of amino acids connected by peptide bonds. An amino acid unit in a polypeptide (or protein) is called a residue. The terms "polypeptide" and "protein" also applies to any amino acid polymers in which one or more amino acid residue is an artificial chemical analogue of a corresponding naturally occurring amino acid, as well as to any naturally occurring amino acid polymers. The essential nature of such analogues of naturally occurring amino acids is that, when incorporated into a polypeptide, that polypeptide is specifically reactive to antibodies elicited to the same polypeptide but consisting entirely of naturally occurring amino acids. It is well known in the art that proteins or polypeptides may undergo modification, including but not limited to, disulfide bond formation, gamma-carboxylation of glutamic acid residues, glycosylation, lipid attachment, phosphorylation, oligomerization, hydroxylation and ADP-ribosylation. Exemplary modifications are described in most basic texts, such as, for example, *Proteins - Structure and Molecular Properties*, 2nd ed., T. E. Creighton, W. H. Freeman and Company, New York (1993), herein incorporated by reference in its entirety. Many detailed reviews are available on this subject, such as, for example, those provided by Wold, F., Post-translational Protein Modifications. Perspectives and Prospects, pp. 1-12 in *Post-translational Covalent Modification of Proteins*, B. C. Johnson, Ed., Academic Press, New York (1983); Seifter *et al.*, *Meth. Enzymol.* 182:626-M (1990) and Rattan *et al.*, *Protein Synthesis: Post-translational Modifications and Aging*, *Ann. N.Y. Acad. Sci.* 663:48-62 (1992), herein incorporated by reference in their entirety. Modifications can occur anywhere in a polypeptide, including the

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peptide backbone, the amino acid side-chains and the amino or carboxyl termini. In fact, blockage of the amino or carboxyl group in a polypeptide, or both, by a covalent modification, is common in naturally occurring and synthetic polypeptides and such modifications may be present in polypeptides of the present invention, as well. For instance, the amino terminal residue of polypeptides made in *E. coli* or other cells, prior to proteolytic processing, almost invariably will be N-formylmethionine. During post-translational modification of the polypeptide, a methionine residue at the NH₂ terminus may be deleted. Accordingly, this invention contemplates the use of both the methionine containing and the methionine-less amino terminal variants of the polypeptide of the invention. Thus, as used herein, the terms "protein" and "polypeptide" include any protein or polypeptide that is modified by any biological or non-biological process.

The terms "amino acid" and "amino acids" refer to all naturally occurring amino acids and, unless otherwise limited, known analogs of natural amino acids that can function in a similar manner as naturally occurring amino acids. This definition is meant to include norleucine, ornithine, homocysteine, and homoserine.

The term "amino acid sequence" means the sequence of amino acids in a polypeptide (or protein) that is written starting with the amino-terminal (N-terminal) residue and ending with the carboxyl-terminal (C-terminal) residue.

The term "an amino acid subsequence" means a portion of the amino acid sequence of a polypeptide. An amino acid subsequence generally has a length of 3 to 50 amino acid residues.

Both terms "substantially purified polypeptide" and "substantially purified protein", as used herein, means a polypeptide or protein that is separated substantially from all other molecules normally associated with it in its native state and is the predominant species present in a preparation. A substantially purified molecule may be greater than 60% free, preferably 75% free, more preferably 90% free, and most preferably 95% free from the other molecules (exclusive of solvent) present in the natural mixture.

Percentage of sequence identity is determined by comparing two optimally aligned sequences over a comparison window, wherein the portion of the polynucleotide or amino acid sequence in the comparison window may comprise additions or deletions (i.e., gaps) as compared to the reference sequence (which does not comprise additions or deletions) for optimal alignment of the two sequences. The percentage is calculated by determining the number of

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positions at which the identical nucleic acid base or amino acid residue occurs in both sequences to yield the number of matched positions, dividing the number of matched positions by the total number of positions in the window of comparison and multiplying the result by 100 to yield the percentage of sequence identity.

5 The term "identity" refers to amino acid or nucleic acid sequences that when compared using the *local homology* algorithm of Smith and Waterman (T. F. Smith and M. S. Waterman, J. Mol. Biol. (147) 195-197 (1981). in the SSEARCH3 3.0t75 (W.R. Pearson, Genomics (11)635-650 (1991) or BLAST 2.2.1 (Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. Lipman , Nucleic Acids Res. (25)3389-10 3402 (1997) programs which are exactly alike.

 The term "similarity" refers to two amino acids, which are similar as defined by similarity matrix BLOSSUM62 (S. Henikoff and J. G. Henikoff, , Proc. Natl. Acad. Sci. U.S.A. (89) 10915-10919 (1992) which is used in BLAST 2.2.1 (Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. 15 Lipman , Nucleic Acids Res. (25)3389-3402 (1997). The BLAST uses alias "positive" for similarity. These two terms, similarity and positive-ness, are interchangeable.

 The term "percent identity" for a pair of protein sequences refers to the number of identical amino acid residues in a two-sequence alignment reported by BLAST, divided by the total number of amino acid residues in the same alignment, expressed in percentage.

20 The term "percent similarity" for a pair of protein sequences refers to the number of similar ("Positive" in BLAST output) amino acid residues in a two-sequence alignment reported by BLAST, divided by the total number of amino acid residues in the same alignment, expressed in percentage.

 Both terms "substantially identical" and "substantial identity", as reference to two amino 25 acid sequences or two nucleotide sequences, means that one amino acid sequence or one nucleotide sequence has at least 50% sequence identity compared to the other amino acid sequence or nucleotide sequence as a reference sequence using the Gap program in the WISCONSIN PACKAGE version 10.0-UNIX from Genetics Computer Group, Inc. based on the method of Needleman and Wunsch (J. Mol. Biol. 48:443-453 (1970), herein incorporated by 30 reference in its entirety) using the set of default parameters for pair-wise comparison (for amino

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acid sequence comparison: Gap Creation Penalty = 8, Gap Extension Penalty = 2; for nucleotide sequence comparison: Gap Creation Penalty = 50; Gap Extension Penalty = 3).

Polypeptides, which are "substantially similar" share sequences as noted above except that residue positions which are not identical may differ by conservative amino acid changes. Conservative amino acid substitutions refer to the interchangeability of residues having similar side chains. "Conservative amino acid substitutions" mean substitutions of one or more amino acids in a native amino acid sequence with another amino acid(s) having similar side chains, resulting in a silent change. Conserved substitutes for an amino acid within a native amino acid sequence can be selected from other members of the group to which the naturally occurring amino acid belongs. For example, a group of amino acids having aliphatic side chains is glycine, alanine, valine, leucine, and isoleucine; a group of amino acids having aliphatic-hydroxyl side chains is serine and threonine; a group of amino acids having amide-containing side chains is asparagine and glutamine; a group of amino acids having aromatic side chains is phenylalanine, tyrosine, and tryptophan; a group of amino acids having basic side chains is lysine, arginine, and histidine; and a group of amino acids having sulfur-containing side chains is cysteine and methionine. Preferred conservative amino acids substitution groups are: valine-leucine, valine-isoleucine, phenylalanine-tyrosine, lysine-arginine, alanine-valine, aspartic acid-glutamic acid, and asparagine-glutamine.

One skilled in the art will recognize that the values of the above substantial identity of nucleotide sequences can be appropriately adjusted to determine corresponding sequence identity of two nucleotide sequences encoding the polypeptides of the present invention by taking into account codon degeneracy, conservative amino acid substitutions, reading frame positioning and the like. Substantial identity of nucleotide sequences for these purposes normally means sequence identity of at least 35%.

The term "codon degeneracy" means divergence in the genetic code permitting variation of the nucleotide sequence without affecting the amino acid sequence of an encoded polypeptide. The skilled artisan is well aware of the "codon-bias" exhibited by a specific host cell in usage of nucleotide codons to specify a given amino acid. Therefore, when synthesizing a gene for ectopic expression in a host cell, it is desirable to design the gene such that its frequency of codon usage approaches the frequency of preferred codon usage of the host cell.

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In another aspect, the present invention provides an isolated nucleic acid molecule in a DNA construct comprising a promoter operably linked to nucleotide sequence or complement thereof, wherein the nucleotide sequence hybridizes under stringent conditions to the complement of a second nucleotide sequence encoding a polypeptide having an amino acid sequence selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71.

The present invention also provides a method for obtaining an isolated nucleic acid molecule of the present invention, the method comprising the steps of: (a) probing a cDNA or genomic library with a hybridization probe comprising a nucleotide sequence encoding all or a portion of the amino acid sequence of a polypeptide, wherein the amino acid sequence of the polypeptide is selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, 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: 52, SEQ ID NO: 54, SEQ ID NO: 67, SEQ ID NO: 69, and SEQ ID NO: 71 (b) identifying a DNA clone that hybridizes under stringent conditions to hybridization probe; (c) isolating the DNA clone identified in step (b); and (d) sequencing the cDNA insert or genomic fragment contained in the DNA clone isolated in step (c) wherein the sequenced nucleic acid molecule encodes all or a substantial portion of the amino acid sequence of the present invention.

Hybridization conditions are sequence dependent and will be different in different circumstances. As used herein "stringent conditions" are selected to be about 5°C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength and pH. The "thermal melting point" is the temperature (under defined ionic strength and pH) at which 50% of a target molecule hybridizes to a completely complementary molecule. Appropriate stringent conditions which promote DNA hybridization, for example, 6.0 X sodium chloride/sodium citrate (SSC) at about 45°C, followed by a wash of 2.0 X SSC at 50°C, are known to those skilled in the art or can be found in *Current Protocols in Molecular Biology*, John Wiley & Sons, N.Y. (1989), 6.3.1-6.3.6, incorporated herein by reference in its entirety. For example, the salt

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concentration in the wash step can be selected from a low stringent condition of about 2.0 X SSC at 50°C to a high stringency of about 0.2 X SSC at 50°C. In addition, the temperature in the wash step can be increased from low stringent conditions at room temperature, about 22°C, to high stringent conditions at about 65°C. Both temperature and salt concentration may be varied, or either the temperature or the salt concentration may be held constant while the other variable is changed. For the purposes of this disclosure, stringent conditions include at least one wash in 2.0 X SSC at a temperature of at least about 50°C for 20 minutes, or equivalent conditions.

The nucleic acid molecules of the present invention may be combined with other non-native, or "heterologous" sequences in a variety of ways. By "heterologous" sequences it is meant any sequence that is not naturally found joined to the nucleotide sequence encoding polypeptide of the present invention, including, for example, combinations of nucleotide sequences from the same plant which are not naturally found joined together, or the two sequences originate from two different species.

The term "homologous" refers to two or more genes that are derived from a single gene in a common ancestor. The term "homology is attributed to descent from a common ancestor. (J. J. Doyle and B. S. Gaut, *Plant Molecular Biology* (42)1-23, 2000.

The term "orthologous" refers to different homologous sequences in different species that arose from a common ancestral gene during speciation; may or may not be responsible for a similar function.

The term paralogous refers to homologous sequences within a single species that arose by gene duplication.

The term "domain" refers to a discrete portion of a protein assumed to fold independently of the rest of the protein and possessing its own function. Domain is also used to refer a discrete portion of nucleotide when translated provides a portion of protein which is assumed to fold independently of the rest of the translated nucleotide sequence and possessing its own function.

The term "Motif" refers to a short conserved region in a protein or nucleotide sequence. Motifs are frequently highly conserved parts of domains.

The term "genome" as it applies to plant cells encompasses not only chromosomal DNA found within the nucleus, but organelle DNA found within subcellular components of the cell. DNAs of the present invention introduced into plant cells can therefore be either chromosomally integrated or organelle-localized.

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The term "operably linked", as used in reference to a regulatory sequence and a structural nucleotide sequence, means that the regulatory sequence causes regulated expression of the operably linked structural nucleotide sequence. "Expression" means the transcription and stable accumulation of sense or antisense RNA derived from the nucleic acid molecule of the present invention. Expression may also refer to translation of mRNA into a polypeptide. "Sense" RNA means RNA transcript that includes the mRNA and so can be translated into polypeptide or protein by the cell. "Antisense RNA" means a RNA transcript that is complementary to all or part of a target primary transcript or mRNA and that blocks the expression of a target gene (U.S. Pat. No. 5,107,065, incorporated herein by reference). The complementarity of an antisense RNA may be with any part of the specific gene transcript, i.e., at the 5' non-coding sequence, 3' non-translated sequence, introns, or the coding sequence. "RNA transcript" means the product resulting from RNA polymerase-catalyzed transcription of a DNA sequence. When the RNA transcript is a perfect complementary copy of the DNA sequence, it is referred to as the primary transcript or it may be a RNA sequence derived from post-transcriptional processing of the primary transcript and is referred to as the mature RNA.

The term "overexpression" means the expression of a polypeptide encoded by an exogenous nucleic acid molecule introduced into a host cell, wherein said polypeptide is either not normally present in the host cell, or wherein said polypeptide is present in said host cell at a higher level than that normally expressed from the endogenous gene encoding said polypeptide.

By "ectopic expression" it is meant that expression of a nucleic acid molecule encoding a polypeptide in a cell type other than a cell type in which the nucleic acid molecule is normally expressed, at a time other than a time at which the nucleic acid molecule is normally expressed or at an expression level other than the level at which the nucleic acid molecule normally is expressed.

"Antisense inhibition" means the production of antisense RNA transcripts capable of suppressing the expression of the target polypeptide. "Co-suppression" means the production of sense RNA transcripts capable of suppressing the expression of identical or substantially similar foreign or endogenous genes (U.S. Patent No. 5,231,020, incorporated herein by reference).

The term "a gene" means the segment of DNA that is involved in producing a polypeptide. Such segment of DNA includes regulatory sequences preceding (5' non-coding sequences) and following (3' non-coding sequences) the coding region as well as intervening

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sequences (introns) between individual coding segments (exons). A "Native gene" means a gene as found in nature with its own regulatory sequences. "Chimeric gene" means any gene that is not a native gene, comprising regulatory and coding sequences that are not found together in nature. Accordingly, a chimeric gene may comprise regulatory sequences and coding sequences that are derived from different sources, or regulatory sequences and coding sequences derived from the same source, but arranged in a manner different than that found in nature. "Endogenous gene" or "endogenous DNA molecule" means a native gene in its natural location in the genome of an organism. A "foreign gene" means a gene not normally found in the host organism, but that is introduced into the host organism by gene transfer. Foreign genes can comprise native genes inserted into a non-native organism, or chimeric genes. A "transgene" is a gene that has been introduced into the genome by a transformation procedure.

"Regulatory sequences" mean nucleotide sequences located upstream (5' non-coding sequences), within, or downstream (3' non-translated sequences) of a structural nucleotide sequence, and which influence the transcription, RNA processing or stability, or translation of the associated structural nucleotide sequence. Regulatory sequences may include promoters, translation leader sequences, introns, and polyadenylation recognition sequences.

The term "promoter sequence" means a nucleotide sequence that is capable of, when located in *cis* to a structural nucleotide sequence encoding a polypeptide, functioning in a way that directs expression of one or more mRNA molecules that encodes the polypeptide. Such promoter regions are typically found upstream of the trinucleotide ATG sequence at the start site of a polypeptide-coding region. Promoter sequences can also include sequences from which transcription of transfer RNA (tRNA) or ribosomal RNA (rRNA) sequences are initiated. Transcription involves the synthesis of a RNA chain representing one strand of a DNA duplex. By "representing" it is meant that the RNA is identical in sequence with one strand of the DNA; it is complementary to the other DNA strand, which provides the template for its synthesis. Transcription takes place by the usual process of complementary base pairing, catalyzed and scrutinized by the enzyme RNA polymerase. The reaction can be divided into three stages described as initiation, elongation and termination. Initiation begins with the binding of RNA polymerase to the double stranded (DS or ds) DNA. The sequence of DNA required for the initiation reaction defines the promoter. The site at which the first nucleotide is incorporated is called the start-site or start-point of transcription. Elongation describes the phase during which

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the enzyme moves along the DNA and extends the growing RNA chain. Elongation involves the disruption of the DNA double stranded structure in which a transiently unwound region exists as a hybrid RNA-DNA duplex and a displaced single strand of DNA. Termination involves recognition of the point at which no further bases should be added to the chain. To terminate
5 transcription, the formation of phosphodiester bonds must cease and the transcription complex must come apart. When the last base is added to the RNA chain, the RNA-DNA hybrid is disrupted, the DNA reforms into a duplex state, and the RNA polymerase enzyme and RNA molecule are both released from the DNA. The sequence of DNA required for the termination reaction is called the terminator.

10 The promoter sequence consists of proximal and more distal upstream elements, the latter elements often referred to as enhancers. Accordingly, an "enhancer" is a DNA sequence that can stimulate promoter activity and may be an innate element of the promoter or a heterologous element inserted to enhance the level or tissue-specificity of a promoter. Promoters may be derived in their entirety from a native gene, or be composed of different elements derived from
15 different promoters found in nature, or even comprise synthetic DNA segments. It is understood by those skilled in the art that different promoters may direct the expression of a gene in different tissues or cell types, or at different stages of development, or in response to different environmental conditions.

Promoters, which are known or are found to cause transcription of DNA in plant cells,
20 can be used in the present invention. Such promoters may be obtained from a variety of sources such as plants and plant viruses. A number of promoters, including constitutive promoters, inducible promoters and tissue-specific promoters, that are active in plant cells have been described in the literature. In addition to promoters that are known to cause transcription of DNA in plant cells, other promoters may be identified for use in the current invention by screening a
25 plant cDNA library for genes that are selectively or preferably expressed in the target tissues and then determine the promoter regions.

The term "constitutive promoter" means a regulatory sequence which causes expression of a structural nucleotide sequence in most cells or tissues at most times. Constitutive promoters are active under most environmental conditions and states of development or cell differentiation.
30 A variety of constitutive promoters are well known in the art. Examples of constitutive promoters that are active in plant cells include but are not limited to the nopaline synthase (NOS)

promoters; DNA plant virus promoters including, but not limited to the caulimovirus promoters for example, cauliflower mosaic virus (CaMV) 19S and 35S, and figwort mosaic virus promoters; the bacilliform virus promoter for example sugar cane bacilliform virus, rice tungro bacilliform virus, among others; plant actin promoters, such as the *Arabidopsis* and rice actin gene promoter (see, e.g., Huang *et al.*, *Plant Mol. Biol.* 33:125-139 (1997), US Patent No. 5,641,876), herein incorporated by reference in its entirety). These promoters when used in a DNA construct are heterologous to the linked gene sequence when they are derived from a different organism, plant species, or a different gene.

The term "inducible promoter" means a regulatory sequence which causes conditional expression of a structural nucleotide sequence under the influence of changing environmental conditions or developmental conditions. Examples of inducible promoters include but are not limited to the senescence-induced promoter for the senescence-associated gene, SAG12, (Gan and Amasino, *Science* 270: 1986-1988 (1995), herein incorporated by reference in its entirety); the light-inducible promoter from the small subunit of ribulose-1,5-bis-phosphate carboxylase (ssRUBISCO); the drought-inducible promoter of maize (Busk *et al.*, *Plant J.* 11:1285-1295 (1997), herein incorporated by reference in its entirety); the cold, drought, and high salt inducible promoter from potato (Kirch, *Plant Mol. Biol.* 33:897-909 (1997), herein incorporated by reference in its entirety); a nitrate-inducible promoter derived from the spinach nitrite reductase gene (Back *et al.*, *Plant Mol. Biol.* 17:9 (1991), herein incorporated by its entirety); salicylic acid inducible promoter (Uknes *et al.*, *Plant Cell* 5:159-169 (1993); Bi *et al.*, *Plant J.* 8:235-245 (1995) herein incorporated by reference in their entireties); the auxin-response elements E1 promoter fragment (AuxREs) in the soybean (*Glycine max* L.) (Liu *et al.*, *Plant Physiol.* 115:397-407 (1997), herein incorporated by reference in its entirety); the auxin-responsive *Arabidopsis* GST6 promoter (also responsive to salicylic acid and hydrogen peroxide) (Chen *et al.*, *Plant J.* 10: 955-966 (1996), herein incorporated by reference in its entirety); the auxin-inducible parC promoter from tobacco (Sakai *et al.*, *Plant Cell Physiol.* 37:906-913 (1996), herein incorporated by reference in its entirety); a plant biotin response element (Streit *et al.*, *Mol. Plant Microbe Interact.* 10:933-937 (1997), herein incorporated by reference in its entirety); the promoter responsive to the stress hormone abscisic acid (Sheen *et al.*, *Science* 274:1900-1902 (1996), herein incorporated by reference in its entirety); the maize In2-2 promoter activated by benzenesulfonamide herbicide safeners (De Veylder *et al.*, *Plant Cell*

Physiol. 38:568-577 (1997), herein incorporated by reference in its entirety); a tetracycline-inducible promoter, such as the promoter for the *Avena sativa* L. (oat) arginine decarboxylase gene (Masgrau *et al.*, *Plant J.* 11:465-473 (1997), herein incorporated by reference in its entirety); and a salicylic acid-responsive element (Stange *et al.*, *Plant J.* 11:1315-1324 (1997),
5 herein incorporated by reference in its entirety).

The term "tissue-specific promoter" or "organ enhanced promoter" means a regulatory sequence that causes transcriptions or enhanced transcriptions of DNA in specific cells or tissues at specific times during plant development, such as in vegetative tissues or reproductive tissues. Examples of tissue-specific promoters under developmental control include promoters that
10 initiate transcription only (or primarily only) in certain tissues, such as vegetative tissues, e.g., roots, leaves or stems, or reproductive tissues, such as fruit, ovules, seeds, pollen, pistils, flowers, or any embryonic tissue. Reproductive tissue specific promoters may be, e.g., ovule-specific, embryo-specific, endosperm-specific, integument-specific, seed coat-specific, pollen-specific, petal-specific, sepal-specific, or some combination thereof. One skilled in the art will
15 recognize that a tissue-specific promoter may drive expression of operably linked sequences in tissues other than the target tissue. Thus, as used herein a tissue-specific promoter is one that drives expression preferentially in the target tissue, but may also lead to some expression in other tissues as well.

A variety of promoters specifically active in vegetative tissues, such as leaves, stems,
20 roots and tubers, can be used to express the nucleic acid molecules of the present invention. Examples of tuber-specific promoters include but are not limited to the class I and II patatin promoters (Bevan *et al.*, *EMBO J.* 8: 1899-1906 (1986); Koster-Topfer *et al.*, *Mol Gen Genet.* 219: 390-396 (1989); Mignery *et al.*, *Gene.* 62: 27-44 (1988); Jefferson *et al.*, *Plant Mol. Biol.* 14: 995-1006 (1990), herein incorporated by reference in their entireties), the promoter for the
25 potato tuber ADPGPP genes, both the large and small subunits; the sucrose synthase promoter (Salanoubat and Belliard, *Gene.* 60: 47-56 (1987), Salanoubat and Belliard, *Gene.* 84: 181-185 (1989), herein incorporated by reference in their entirety); and the promoter for the major tuber proteins including the 22 kd protein complexes and proteinase inhibitors (Hannapel, *Plant Physiol.* 101: 703-704 (1993), herein incorporated by reference in its entirety). Examples of leaf-
30 specific promoters include but are not limited to the ribulose biphosphate carboxylase (RBCS or RuBISCO) promoters (see, *e.g.*, Matsuoka *et al.*, *Plant J.* 6:311-319 (1994), herein incorporated

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by reference in its entirety); the light harvesting chlorophyll a/b binding protein gene promoter (see, e.g., Shiina *et al.*, *Plant Physiol.* 115:477-483 (1997); Casal *et al.*, *Plant Physiol.* 116:1533-1538 (1998), herein incorporated by reference in their entireties); and the *Arabidopsis thaliana* myb-related gene promoter (Atmyb5) (Li *et al.*, *FEBS Lett.* 379:117-121 (1996), herein incorporated by reference in its entirety). Examples of root-specific promoter include but are not limited to the promoter for the acid chitinase gene (Samac *et al.*, *Plant Mol. Biol.* 25: 587-596 (1994), herein incorporated by reference in its entirety); the root specific subdomains of the CaMV35S promoter that have been identified (Lam *et al.*, *Proc. Natl. Acad. Sci. (U.S.A.)* 86:7890-7894 (1989), herein incorporated by reference in its entirety); the ORF13 promoter from *Agrobacterium rhizogenes* which exhibits high activity in roots (Hansen *et al.*, *Mol. Gen. Genet.* 254:337-343 (1997), herein incorporated by reference in its entirety); the promoter for the tobacco root-specific gene TobRB7 (Yamamoto *et al.*, *Plant Cell* 3:371-382 (1991), herein incorporated by reference in its entirety); and the root cell specific promoters reported by Conkling *et al.* (Conkling *et al.*, *Plant Physiol.* 93:1203-1211 (1990), herein incorporated by reference in its entirety). Another class of useful vegetative tissue-specific promoters is meristematic (root tip and shoot apex) promoters. For example, the "SHOOTMERISTEMLESS" and "SCARECROW" promoters, which are active in the developing shoot or root apical meristems (Di Laurenzio *et al.*, *Cell* 86:423- 433 (1996); Long, *Nature* 379:66-69 (1996); herein incorporated by reference in their entireties), can be used. Another example of a useful promoter is that which controls the expression of 3-hydroxy-3- methylglutaryl coenzyme A reductase HMG2 gene, whose expression is restricted to meristematic and floral (secretory zone of the stigma, mature pollen grains, gynoeceium vascular tissue, and fertilized ovules) tissues (see, e.g., Enjuto *et al.*, *Plant Cell.* 7:517-527 (1995), herein incorporated by reference in its entirety). Also another example of a useful promoter is that which controls the expression of knl-related genes from maize and other species which show meristem-specific expression (see, e.g., Granger *et al.*, *Plant Mol. Biol.* 31:373-378 (1996); Kerstetter *et al.*, *Plant Cell* 6:1877-1887 (1994); Hake *et al.*, *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* 350:45-51 (1995), herein incorporated by reference in their entireties). Another example of a meristematic promoter is the *Arabidopsis thaliana* KNAT1 promoter. In the shoot apex, KNAT1 transcript is localized primarily to the shoot apical meristem; the expression of KNATI in the shoot meristem decreases during the floral transition

and is restricted to the cortex of the inflorescence stem (see, e.g., Lincoln *et al.*, *Plant Cell* 6:1859-1876 (1994), herein incorporated by reference in its entirety).

Suitable seed-specific promoters can be derived from the following genes: MAC1 from maize (Sheridan *et al.*, *Genetics* 142:1009-1020 (1996), herein incorporated by reference in its entirety); Cat3 from maize (GenBank No. L05934, Abler *et al.*, *Plant Mol. Biol.* 22:10131-1038 (1993), herein incorporated by reference in its entirety); viviparous-1 from *Arabidopsis* (Genbank No. U93215); Atimyc1 from *Arabidopsis* (Urao *et al.*, *Plant Mol. Biol.* 32:571-57 (1996); Conceicao *et al.*, *Plant* 5:493-505 (1994), herein incorporated by reference in their entireties); napA from *Brassica napus* (GenBank No. J02798); the napin gene family from *Brassica napus* (Sjodahl *et al.*, *Planta* 197:264-271 (1995), herein incorporated by reference in its entirety).

The ovule-specific promoter for BEL1 gene (Reiser *et al.* *Cell* 83:735-742 (1995), GenBank No. U39944; Ray *et al.*, *Proc. Natl. Acad. Sci. USA* 91:5761-5765 (1994), all of which are herein incorporated by reference in their entireties) can also be used. The egg and central cell specific MEA (FIS1) and FIS2 promoters are also useful reproductive tissue-specific promoters (Luo *et al.*, *Proc. Natl. Acad. Sci. USA*, 97:10637-10642 (2000); Vielle-Calzada, *et al.*, *Genes Dev.* 13:2971-2982 (1999); herein incorporated by reference in their entireties).

A maize pollen-specific promoter has been identified in maize (Guerrero *et al.*, *Mol. Gen. Genet.* 224:161-168 (1990), herein incorporated by reference in its entirety). Other genes specifically expressed in pollen have been described (see, e.g., Wakeley *et al.*, *Plant Mol. Biol.* 37:187-192 (1998); Ficker *et al.*, *Mol. Gen. Genet.* 257:132-142 (1998); Kulikauskas *et al.*, *Plant Mol. Biol.* 34:809-814 (1997); Treacy *et al.*, *Plant Mol. Biol.* 34:603-611 (1997); all of which are herein incorporated by reference in their entireties).

Promoters derived from genes encoding embryonic storage proteins, which includes the gene encoding the 2S storage protein from *Brassica napus* (Dasgupta *et al.*, *Gene* 133:301-302 (1993), herein incorporated by reference in its entirety); the 2S seed storage protein gene family from *Arabidopsis*; the gene encoding oleosin 20kD from *Brassica napus* (GenBank No. M63985); the genes encoding oleosin A (GenBank No. U09118) and oleosin B (GenBank No. U09119) from soybean; the gene encoding oleosin from *Arabidopsis* (GenBank No. Z17657); the gene encoding oleosin 18kD from maize (GenBank No. J05212, Lee, *Plant Mol. Biol.* 26:1981-1987 (1994), herein incorporated by reference in its entirety); and the gene encoding

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low molecular weight sulphur rich protein from soybean (Choi et al., Mol. Gen. Genet. 246:266-268 (1995), herein incorporated by reference in its entirety), can also be used.

Promoters derived from zein encoding genes (including the 15 kD, 16 kD, 19 kD, 22 kD, 27 kD, and gamma genes) (Pedersen *et al.*, Cell 29: 1015-1026 (1982), herein incorporated by reference in its entirety) can be also used. The zeins are a group of storage proteins found in maize endosperm.

Other promoters known to function, for example, in maize, include the promoters for the following genes: *waxy*, *Brittle*, *Shrunken 2*, Branching enzymes I and II, starch synthases, debranching enzymes, oleosins, glutelins, and sucrose synthases. A particularly preferred promoter for maize endosperm expression is the promoter for the glutelin gene from rice, more particularly the Osgt-1 promoter (Zheng *et al.*, Mol. Cell Biol. 13: 5829-5842 (1993), herein incorporated by reference in its entirety). Examples of promoters suitable for expression in wheat include those promoters for the ADPglucose pyrophosphorylase (ADPGPP) subunits, the granule bound and other starch synthases, the branching and debranching enzymes, the embryogenesis-abundant proteins, the gliadins, and the glutenins. Examples of such promoters in rice include those promoters for the ADPGPP subunits, the granule bound and other starch synthases, the branching enzymes, the debranching enzymes, sucrose synthases, and the glutelins. A particularly preferred promoter is the promoter for rice glutelin, Osgt-1. Examples of such promoters for barley include those for the ADPGPP subunits, the granule bound and other starch synthases, the branching enzymes, the debranching enzymes, sucrose synthases, the hordeins, the embryo globulins, and the aleurone specific proteins.

A tomato promoter active during fruit ripening, senescence and abscission of leaves and, to a lesser extent, of flowers can be used (Blume *et al.*, Plant J. 12:731-746 (1997), herein incorporated by reference in its entirety). Other exemplary promoters include the pistil specific promoter in the potato (*Solanum tuberosum* L.) SK2 gene, encoding a pistil-specific basic endochitinase (Ficker *et al.*, Plant Mol. Biol. 35:425-431 (1997), herein incorporated by reference in its entirety); the Blec4 gene from pea (*Pisum sativum* cv. Alaska), active in epidermal tissue of vegetative and floral shoot apices of transgenic alfalfa. This makes it a useful tool to target the expression of foreign genes to the epidermal layer of actively growing shoots. The tissue specific E8 promoter from tomato is also useful for directing gene expression in fruits.

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It is recognized that additional promoters that may be utilized are described, for example, in U.S. Patent Nos. 5,378,619, 5,391,725, 5,428,147, 5,447,858, 5,608,144, 5,608,144, 5,614,399, 5,633,441, 5,633,435, and 4,633,436, all of which are herein incorporated in their entirety. In addition, a tissue specific enhancer may be used (Fromm *et al.*, *The Plant Cell* 5 1:977-984 (1989), herein incorporated by reference in its entirety). It is further recognized that since in most cases the exact boundaries of regulatory sequences have not been completely defined, DNA fragments of different lengths may have identical promoter activity.

The "translation leader sequence" means a DNA sequence located between the promoter sequence of a gene and the coding sequence. The translation leader sequence is present in the 10 fully processed mRNA upstream of the translation start sequence. The translation leader sequence may affect processing of the primary transcript to mRNA, mRNA stability or translation efficiency. Examples of translation leader sequences have been described (US Patent No. 5,659,122 and Turner and Foster, *Molecular Biotechnology* 3:225 (1995) herein incorporated by reference in its entirety).

15 The "3' non-translated sequences" means DNA sequences located downstream of a structural nucleotide sequence and include sequences encoding polyadenylation and other regulatory signals capable of affecting mRNA processing or gene expression. The polyadenylation signal functions in plants to cause the addition of polyadenylate nucleotides to the 3' end of the mRNA precursor. The polyadenylation sequence can be derived from the natural 20 gene, from a variety of plant genes, or from T-DNA. An example of the polyadenylation sequence is the nopaline synthase 3' sequence (NOS 3'; Fraley *et al.*, *Proc. Natl. Acad. Sci. USA* 80: 4803-4807 (1983), herein incorporated by reference in its entirety). The use of different 3' non-translated sequences is exemplified by Ingelbrecht *et al.*, *Plant Cell* 1:671-680 (1989), herein incorporated by reference in its entirety.

25 The isolated nucleic acid molecules of the present invention may also include introns. Generally, optimal expression in monocotyledonous and some dicotyledonous plants is obtained when an intron sequence is inserted between the promoter sequence and the structural gene sequence or, optionally, may be inserted in the structural coding sequence to provide an interrupted coding sequence. An example of such an intron sequence is the HSP 70 intron 30 described in WO 93/19189, herein incorporated by reference in its entirety.

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The laboratory procedures in recombinant DNA technology used herein are those well known and commonly employed in the art. Standard techniques are used for cloning, DNA and RNA isolation, amplification and purification. Generally enzymatic reactions involving DNA ligase, DNA polymerase, restriction endonucleases and the like are performed according to the manufacturer's specifications. These techniques and various other techniques are generally performed according to Sambrook et al., *Molecular Cloning - A Laboratory Manual*, 2nd. ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, New York (1989).

Isolation and identification of nucleic acid molecules encoding polypeptides of the present invention from soybean, corn, rice and other species are described in detail in Examples. All or a substantial portion of the nucleic acid molecules of the present invention may be used to isolate cDNAs and nucleic acid molecules encoding homologous polypeptides from the same or other plant species.

A "substantial portion" of a nucleotide sequence comprises enough of the sequence to afford specific identification and/or isolation of a nucleic acid molecule comprising the sequence. Nucleotide sequences can be evaluated either manually by one skilled in the art, or by using computer-based sequence comparison and identification tools that employ algorithms such as BLAST (Basic Local Alignment Search Tool; Altschul *et al. J Mol. Biol.* 215:403-410 (1993)). In general, a sequence of thirty or more contiguous nucleotides is necessary in order to putatively identify a nucleotide sequence as homologous to a gene. Moreover, with respect to nucleotide sequences, gene-specific oligonucleotide probes comprising 30 or more contiguous nucleotides may be used in sequence-dependent methods of gene identification (e.g., Southern hybridization) and isolation (e.g., in situ hybridization of bacterial colonies or bacteriophage plaques). In addition, short oligonucleotides of 12 or more nucleotides may be used as amplification primers in PCR in order to obtain a particular nucleic acid molecule comprising the primers. The skilled artisan, having the benefit of the sequences as reported herein, may now use all or a substantial portion of the disclosed sequences for purposes known to those skilled in this art. Accordingly, the instant invention comprises the complete sequences as reported in the accompanying Sequence Listing, as well as substantial portions of those sequences as defined above.

Isolation of nucleic acid molecules encoding homologous polypeptides using sequence-dependent protocols is well known in the art. Examples of sequence-dependent protocols include, but are not limited to, methods of nucleic acid molecule hybridization, and methods of

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DNA and RNA amplification as exemplified by various uses of nucleic acid molecule amplification technologies (e.g., polymerase chain reaction, ligase chain reaction).

For example, structural nucleic acid molecules encoding other polypeptide of the present invention, either as cDNAs or genomic DNAs, could be isolated directly by using all or a substantial portion of the nucleic acid molecules of the present invention as DNA hybridization probes to screen cDNA or genomic libraries from any desired plant employing methodology well known to those skilled in the art. Methods for forming such libraries are well known in the art. Specific oligonucleotide probes based upon the nucleic acid molecules of the present invention can be designed and synthesized by methods known in the art. Moreover, the entire sequences of the nucleic acid molecules can be used directly to synthesize DNA probes by methods known to the skilled artisan such as random primer DNA labeling, nick translation, or end-labeling techniques, or RNA probes using available in vitro transcription systems. In addition, specific primers can be designed and used to amplify a part or all of the sequences. The resulting amplification products can be labeled directly during amplification reactions or labeled after amplification reactions, and used as probes to isolate full-length cDNA or genomic DNAs under conditions of appropriate stringency.

Alternatively, the nucleic acid molecules of interest can be amplified from nucleic acid samples using amplification techniques. For instance, the disclosed nucleic acid molecules may be used to define a pair of primers that can be used with the polymerase chain reaction (Mullis, *et al.*, *Cold Spring Harbor Symp. Quant. Biol.* 51:263-273 (1986); Erlich *et al.*, EP 50,424; EP 84,796, EP 258,017, EP 237,362; Mullis, EP 201,184; Mullis *et al.*, US 4,683,202; Erlich, US 4,582,788; and Saiki, R. *et al.*, US 4,683,194, all of which are herein incorporated by reference in their entireties) to amplify and obtain any desired nucleic acid molecule directly from mRNA, from cDNA, from genomic libraries or cDNA libraries. PCR and other in vitro amplification methods may also be useful, for example, to clone nucleotide sequences that encode for polypeptides to be expressed, to make nucleic acid molecules to use as probes for detecting the presence of the desired mRNA in samples, for nucleic acid sequencing, or for other purposes. In addition, two short segments of the nucleic acid molecules of the present invention may be used in polymerase chain reaction protocols to amplify longer nucleic acid molecules encoding homologues of a polypeptide of the present invention from DNA or RNA. For example, the skilled artisan can follow the RACE protocol (Frohman *et al.*, *Proc. Natl. Acad. Sci. USA*

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85:8998 (1988), herein incorporated by reference in its entirety) to generate cDNAs by using PCR to amplify copies of the region between a single point in the transcript and the 3' or 5' end. Primers oriented in the 3' and 5' directions can be designed from the nucleic acid molecules of the present invention. Using commercially available 3'RACE or 5'RACE systems (Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A.), specific 3' or 5' cDNA fragments can be isolated (Ohara et al., Proc. Natl. Acad. Sci. USA 86:5673 (1989); Loh et al., Science 243:217 (1989), both of which are herein incorporated by reference in their entireties). Products generated by the 3' and 5' RACE procedures can be combined to generate full-length cDNAs (Frohman and Martin, Techniques.1: 165 (1989), herein incorporated by reference in its entirety).

Nucleic acid molecules of interest may also be synthesized, either completely or in part, especially where it is desirable to provide plant-preferred sequences, by well-known techniques as described in the technical literature. See, e.g., Carruthers et al., Cold Spring Harbor Symp. Quant. Biol. 47:411-418 (1982), and Adams et al., J. Am. Chem. Soc. 105:661 (1983), both of which are herein incorporated by reference in their entireties. Thus, all or a portion of the nucleic acid molecules of the present invention may be synthesized using codons preferred by a selected plant host. Plant-preferred codons may be determined, for example, from the codons used most frequently in the proteins expressed in a particular plant host species. Other modifications of the gene sequences may result in mutants having slightly altered activity

Availability of the nucleotide sequences encoding polypeptide of the present invention facilitates immunological screening of cDNA expression libraries. Synthetic polypeptides representing portions of the amino acid sequences of polypeptides of the present invention may be synthesized. These polypeptides can be used to immunize animals to produce polyclonal or monoclonal antibodies with specificity for polypeptides comprising the amino acid sequences. These antibodies can be then be used to screen cDNA expression libraries to isolate full-length cDNA clones of interest (Lemer, Adv. Immunol. 36: 1 (1984); Sambrook et al., Molecular Cloning: A Laboratory Manual; Cold Spring Harbor Laboratory Press: Cold Spring Harbor, (1989)). It is understood that people skilled in the art are familiar with the standard resource materials which describe specific conditions and procedures for the construction, manipulation and isolation of antibodies (see, for example, Harlow and Lane, In *Antibodies: A Laboratory Manual*, Cold Spring Harbor Press, Cold Spring Harbor, New York (1988)).

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Another aspect of the present invention relates to methods of making a DNA construct by obtaining a nucleic acid molecule comprising a nucleotide sequence encoding a polypeptide of the present invention, the amino acid sequence of which has at least 50% sequence identity to a polypeptide selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71.

One aspect of the present invention provides an isolated nucleic acid molecule comprising a nucleotide sequence or complement thereof, wherein the nucleotide sequence encodes a polypeptide having an amino acid sequence that has at least 50% sequence identity, preferably at least 60%, more preferably at least 70% sequence identity, even more preferably at least 80% or 90% sequence identity, and most preferably at least 95% to 98% sequence identity to a polypeptide selected from group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71.

One method of the present invention is for obtaining a nucleic acid molecule encoding all or a substantial portion of the amino acid sequence of a polypeptide of the present invention comprising: (a) probing a cDNA or genomic library with a hybridization probe comprising a nucleotide sequence encoding all or a substantial portion of a polypeptide having an amino acid sequence set forth in any of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, SEQ ID NO: 71 or an amino acid sequence set forth in any of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 10,

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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: 67, SEQ ID NO: 69, SEQ ID NO: 71 with conservative amino acid substitutions; (b) identifying a DNA clone that hybridizes under stringent conditions to the hybridization probe; (c) isolating the DNA clone identified in step (b); and (d) sequencing the cDNA or genomic fragment that comprises the clone isolated in step (c) wherein the sequenced nucleic acid molecule encodes all or a substantial portion of the amino acid sequence of polypeptide of the present invention.

Another method of the present invention for obtaining a nucleic acid molecule encoding all or a substantial portion of the amino acid sequence of a polypeptide of the present invention comprising: (a) synthesizing a first and a second oligonucleotide primers, wherein the sequences of the first and second oligonucleotide primers encode two different portions of a polypeptide having an amino acid sequence selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, SEQ ID NO: 71; and (b) amplifying and obtaining the nucleic acid molecule directly from mRNA samples, from genomic libraries or from cDNA libraries using the first and second oligonucleotide primers of step (a) wherein the nucleic acid molecule encodes all or a substantial portion of the amino acid sequence polypeptide of the present invention.

All or a substantial portion of the nucleic acid molecules of the present invention may also be used as probes for genetically and physically mapping the genes that they are a part of, and as markers for traits linked to those genes. Such information may be useful in plant breeding in order to develop lines with desired phenotypes. For example, the nucleic acid molecules of the present invention may be used as restriction fragment length polymorphism (RFLP) markers. Southern blots (Maniatis et al., *Molecular Cloning: A Laboratory Manual*, Second Edition (1989) Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y., herein incorporated by

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reference in its entirety) of restriction-digested plant genomic DNA may be probed with the nucleic acid fragments of the present invention. The resulting banding patterns may then be subjected to genetic analyses using computer programs such as MapMaker (Lander et al., *Genomics 1*:174-181 (1987), herein incorporated by reference in its entirety) in order to construct
5 a genetic map. In addition, the nucleic acid fragments of the present invention may be used to probe Southern blots containing restriction endonuclease-treated genomic DNAs of a set of individuals representing parent and progeny of a defined genetic cross. Segregation of the DNA polymorphisms is noted and used to calculate the position of the nucleotide sequence of the present invention in the genetic map previously obtained using this population (Botstein *et al.*,
10 *Am. J. Hum. Genet.* 32:314-331 (1980), herein incorporated by reference in its entirety).

The production and use of plant gene-derived probes for use in genetic mapping is described in Bernatzky and Tanksley, *Plant Mol. Biol. Reporter 4*:37-41 (1986), herein incorporated by reference in its entirety. Numerous publications describe genetic mapping of specific cDNA clones using the methodology outlined above or variations thereof. For example,
15 F2 intercross populations, backcross populations, randomly mated populations, near isogenic lines, exotic germplasms, and other sets of individuals may be used for mapping. Such methodologies are well known to those skilled in the art.

Nucleic acid probes derived from the nucleic acid molecules of the present invention may also be used for physical mapping (i.e., placement of sequences on physical maps; see Hoheisel
20 *et al.*, In: *Nonmammalian Genomic Analysis: A Practical Guide*, Academic press 1996, pp. 319-346, and references cited therein).

In another embodiment, nucleic acid probes derived from the nucleic acid molecules of the present invention may be used in direct fluorescence *in situ* hybridization (FISH) mapping (Trask, *Trends Genet.* 7:149-154 (1991), herein incorporated by reference in its entirety).
25 Although current methods of FISH mapping favor use of large clones (several to several hundred KB; see Laan *et al.*, *Genome Res.* 5:13-20 (1995), herein incorporated by reference in its entirety), improvements in sensitivity may allow performance of FISH mapping using shorter probes.

A variety of nucleic acid amplification-based methods of genetic and physical mapping
30 may be carried out using the nucleotide molecules of the present invention. Examples include allele-specific amplification (Kazazian et al., *J. Lab. Clin. Med.* 11:95-96 (1989), herein

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incorporated by reference in its entirety), polymorphism of PCR-amplified fragments (CAPS; Sheffield *et al.*, *Genomics* 16:325-332 (1993), herein incorporated by reference in its entirety), allele-specific ligation (Landegren *et al.*, *Science* 241:1077-1080 (1988) herein incorporated by reference in its entirety), nucleotide extension reactions (Sokolov *et al.*, *Nucleic Acid Res.* 18:3671 (1990) herein incorporated by reference in its entirety), Radiation Hybrid Mapping (Walter *et al.*, *Nat. Genet.* 7:22-28 (1997) herein incorporated by reference in its entirety) and Happy Mapping (Dear and Cook, *Nucleic Acid Res.* 17:6795-6807 (1989) herein incorporated by reference in its entirety). For these methods, the sequence of a nucleic acid fragment is used to design and produce primer pairs for use in the amplification reaction or in primer extension reactions. The design of such primers is well known to those skilled in the art. In methods employing PCR-based genetic mapping, it may be necessary to identify DNA sequence differences between the parents of the mapping cross in the region corresponding to the nucleotide sequence. This, however, is generally not necessary for mapping methods.

Isolated nucleic acid molecules of the present invention may find use in the identification of loss of function mutant phenotypes of a plant, due to a mutation in one or more endogenous genes encoding polypeptides of the present invention. This can be accomplished either by using targeted gene disruption protocols or by identifying specific mutants for these genes contained in a population of plants carrying mutations in all possible genes (Ballinger and Benzer, *Proc. Natl. Acad. Sci. USA* 86:9402-9406 (1989); Koes *et al.*, *Proc. Natl. Acad. Sci. USA* 92:8149-8153 (1995); Bensen *et al.*, *Plant Cell* 7:75-84 (1995) all of which are incorporated herein by reference in their entirety). The latter approach may be accomplished in two ways. First, short segments of the nucleic acid molecules of the present invention may be used in polymerase chain reaction protocols in conjunction with a mutation tag sequence primer on DNAs prepared from a population of plants in which mutator transposons or some other mutation-causing DNA element has been introduced. The amplification of a specific DNA fragment with these primers indicates the insertion of the mutation tag element in or near the plant gene encoding polypeptide(s) of the present invention. Alternatively, the nucleic acid molecules of the present invention may be used as a hybridization probe against PCR amplification products generated from the mutation population using the mutation tag sequence primer in conjunction with an arbitrary genomic site primer, such as that for a restriction enzyme site-anchored synthetic adapter. Isolated nucleic acid molecules of the present invention can be mutated by a variety of methods well known in

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the art (Shortle D. et al., *Annu. Rev. Genet.* 15: 265, 1981; Itakura K. et al., *Ann Rev. Biochem* 53: 323, 1984; Botstein D. & Shortle D., *Science* 229: 1193, 1985; Smith M., *Ann. Rev. Genet.* 19: 423 1985; and Sambrook, *et al.*, In: *Molecular Cloning, A Laboratory Manual, 2nd Edition, Cold Spring Harbor Press, Cold Spring Harbor, New York 1989*, herein incorporated by
5 reference in its entirety). These mutations can also be introduced by causing point mutation or site directed mutagenesis by using commercially available kits such as QuickChange™ from Stratagene (11011 North Torrey Pines Road, La Jolla, CA 92037). For example site directed mutations can include but are not limited to, truncations, substitutions, additions, terminations, fusions of polypeptides or nucleic acid. With either method, a plant containing a mutation in the
10 endogenous gene encoding polypeptides of the present invention can be identified and obtained. Such plant can also be obtained by *in situ* site directed mutagenesis. This mutant plant can then be used to determine or confirm the natural function of the polypeptides of the present invention disclosed herein.

Methods for introducing genetic mutations into plant genes are well known. For instance,
15 seeds or other plant material can be treated with a mutagenic chemical substance, according to standard techniques. Such chemical substances include, but are not limited to, the following: diethyl sulfate, ethylene imine, ethyl methanesulfonate and N-nitroso-N-ethylurea. Alternatively, ionizing radiation from sources such as, for example, X-rays or gamma rays can be used. Desired mutants are selected by assaying for increased seed mass, oil content and other properties.

20 “C-terminal region” refers to the region of a peptide, polypeptide, or protein chain from the middle thereof to the end that carries the amino acid having a free carboxyl group.

“N-terminal region” refers to the region of a peptide, polypeptide, or protein chain from the amino acid having a free amino group to the middle of the chain.

Antisense Oligonucleotides or Polynucleotides of the Present Invention

25 The present invention, in another aspect, provides an antisense oligonucleotide or polynucleotide encoding an RNA molecule which hybridizes to at least a portion of an RNA transcript of an endogenous gene encoding a polypeptide of the present invention, wherein the RNA molecule hybridizes with the RNA transcript such that expression of the endogenous gene is altered.

30 The present invention, in another aspect, provides DNA construct wherein a promoter that functions in plant is operably linked to an antisense oligonucleotide or polynucleotide

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encoding an RNA molecule which hybridizes under stringent hybridization conditions to at least a portion of an RNA transcript of an endogenous gene encoding a polypeptide the amino acid sequence of which has at least 50% sequence identity to a member selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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: 67, SEQ ID NO: 69, and SEQ ID NO: 71. The antisense oligonucleotide or polynucleotide can be full length or preferably has about six to about 100 nucleotides. In one embodiment of this invention, the antisense oligonucleotide or polynucleotide hybridizes under stringent conditions to either at least a corresponding portion of one strand of a nucleotide sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, SEQ ID NO: 70:, or the RNA transcript transcribed from the a nucleotide sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, and SEQ ID NO: 70. In another embodiment, the antisense oligonucleotide or polynucleotide hybridizes under stringent conditions to a corresponding portion of the 5' non-coding region or 3' non-translated region of the RNA transcript. In another embodiment, the antisense oligonucleotide or polynucleotide further comprises a sequence encoding a catalytic RNA or ribozyme.

The antisense oligonucleotides or polynucleotides of the present invention may find particular use in antisense technology to suppress endogenous gene expression to control sizes of

organs in transgenic plants. To accomplish this, a nucleic acid molecule derived from a nucleotide sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, and SEQ ID NO: 70 is cloned and operably linked to a promoter such that the antisense strand of RNA will be transcribed. The construct is then transformed into plants and the antisense strand of RNA is produced. In plant cells, it has been suggested that antisense RNA inhibits gene expression by preventing the accumulation of mRNA that encodes the enzyme of interest (see, *e.g.*, Sheehy *et al.*, *Proc. Nat. Acad. Sci. USA* 85:8805-8809 (1988), and U.S. Patent No. 4,801,340; both of which are herein incorporated by reference in their entireties).

The antisense oligonucleotide or polynucleotide to be introduced generally will be substantially identical to at least a portion of the endogenous gene or genes of the present invention to be repressed. The sequence, however, needs not to be perfectly identical to inhibit or suppress expression of the endogenous gene or genes of the present invention.

For antisense suppression, the introduced sequence also needs not be full length relative to either the primary transcription product or fully processed mRNA. Generally, higher homology can be used to compensate for the use of a shorter sequence. Furthermore, the introduced sequence need not have the same intron or exon pattern, and homology of non-coding segments may be equally effective. Normally, a sequence of between about 6 nucleotides and about full length nucleotides should be used, though a sequence of at least about 500 to about 1700 nucleotides is preferred, a sequence of at least about 200 nucleotides is more preferred, and a sequence of about 6 to about 100 nucleotides is especially preferred.

In another embodiment of this invention, the antisense oligonucleotide or polynucleotide is substantially complementary to at least a corresponding portion of one strand of a nucleotide sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, SEQ ID NO: 70, or the RNA transcript transcribed from the a nucleotide sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, SEQ ID NO: 70. In another embodiment, the antisense oligonucleotide or polynucleotide is substantially complementary to a corresponding portion of the 5' non-coding portion or 3' non-coding portion of one strand of a DNA molecule which has substantial sequence identity to a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, or SEQ ID NO: 70.

The antisense oligonucleotides or polynucleotides of the present invention may further comprise a nucleotide sequence encoding a catalytic RNA molecule or a ribozyme. It is known to a skilled person in the art that catalytic RNA molecules or ribozymes can also be used to inhibit expression of genes of the present invention. It is possible to design ribozymes that specifically pair with virtually any target RNA and cleave the phosphodiester backbone at a specific location, thereby functionally inactivating the target RNA. In carrying out this cleavage, the ribozyme is not itself altered, and is thus capable of recycling and cleaving other molecules, making it a true enzyme. The inclusion of ribozyme sequences within antisense RNAs confers RNA-cleaving activity upon them, thereby increasing the activity of the recombinant DNA constructs.

A number of classes of ribozymes have been identified. One class of ribozymes is derived from a number of small circular RNAs, which are capable of self-cleavage and replication in plants. The RNAs replicate either alone (viroid RNAs) or with a helper virus

(satellite RNAs). Examples include RNAs from avocado sunblotch viroid and the satellite RNAs from tobacco ringspot virus, lucerne transient streak virus, velvet tobacco mottle virus, *Solanum nodiflorum* mottle virus and subterranean clover mottle virus. The design and use of target RNA-specific ribozymes is described in Haseloff *et al. Nature* 334:585-591 (1988), herein
5 incorporated by reference in its entirety.

The present invention also provides antibodies that specifically bind to the polypeptides of the present invention and recombinant DNA constructs that comprise nucleic acid molecules of the present invention.

Plant Recombinant DNA Constructs and Transformed Plants

10 The term "transgenic plant" means a plant that contains an exogenous nucleic acid, which can be derived from the same plant species or from a different species. By "exogenous" it is meant that a nucleic acid molecule originates from outside the plant to which the nucleic acid molecule is introduced. An exogenous nucleic acid molecule can have a naturally occurring or non-naturally occurring nucleotide sequence. One skilled in the art understands that an
15 exogenous nucleic acid molecule can be a heterologous nucleic acid molecule derived from a different plant species than the plant into which the nucleic acid molecule is introduced or can be a nucleic acid molecule derived from the same plant species as the plant into which it is introduced.

The term "genome" as it applies to plant cells encompasses not only chromosomal DNA
20 found within the nucleus, but organelle DNA found within subcellular components of the cell. DNAs of the present invention introduced into plant cells can therefore be either chromosomally integrated or organelle-localized. The term "genome" as it applies to bacteria encompasses both the chromosome and plasmids within a bacterial host cell. Encoding DNAs of the present invention introduced into bacterial host cells can therefore be either chromosomally integrated or
25 plasmid-localized.

Exogenous nucleic acid molecules may be transferred into a plant cell by the use of a recombinant DNA construct (or vector) designed for such a purpose. The present invention also provides a plant recombinant DNA construct (or vector) for producing transgenic plants, wherein the plant recombinant DNA construct (or vector) comprises a sense oligonucleotide or
30 polynucleotide or an antisense oligonucleotide or polynucleotide. Methods that are well known to those skilled in the art may be used to prepare the plant recombinant DNA construct (or

vector) of the present invention. These methods include in vitro recombinant DNA techniques, synthetic techniques, and in vivo genetic recombination. Such techniques are described in Sambrook et al., *Molecular Cloning, A Laboratory Manual*, Cold Spring Harbor Press, Plainview, N.Y. (1989); and Ausubel et al., *Current Protocols in Molecular Biology*, John Wiley & Sons, New York, N.Y. (1989).

A plant recombinant DNA construct (or vector) of the present invention contains a sense oligonucleotide or polynucleotide or an antisense oligonucleotide or polynucleotide and operably linked regulatory sequences or control elements. Exemplary regulatory sequences include but are not limited to promoters, translation leader sequences, introns, 3' non-translated sequences. The promoters can be constitutive, inducible, or preferably tissue-specific promoters.

A plant recombinant DNA construct (vector) of the present invention will typically comprise a selectable marker that confers a selectable phenotype on plant cells. Selectable markers may also be used to select for plants or plant cells that contain the exogenous nucleic acid molecules of the present invention. The marker may encode biocide resistance, antibiotic resistance (e.g., kanamycin, G418 bleomycin, hygromycin, etc.), or herbicide resistance (e.g., glyphosate, etc.). Examples of selectable markers include, but are not limited to, a neo gene (Potrykus *et al.*, *Mol. Gen. Genet.* 199:183-188 (1985)) which codes for kanamycin resistance and can be selected for using kanamycin, G418, etc.; a bar gene which codes for bialaphos resistance; a mutant EPSP synthase gene (Hinchee *et al.*, *Bio/Technology* 6:915-922 (1988)) which encodes glyphosate resistance; a nitrilase gene which confers resistance to bromoxynil (Stalker *et al.*, *J. Biol. Chem.* 263:6310-6314 (1988)); a mutant acetolactate synthase gene (ALS) which confers imidazolinone or sulphonylurea resistance (European Patent Application 154,204 (Sept. 11, 1985)); and a methotrexate resistant DHFR gene (Thillet *et al.*, *J. Biol. Chem.* 263:12500-12508 (1988)).

A plant recombinant DNA construct (vector) of the present invention may also include a screenable marker. Screenable markers may be used to monitor expression. Exemplary screenable markers include a β -glucuronidase or uidA gene (GUS) which encodes an enzyme for which various chromogenic substrates are known (Jefferson, *Plant Mol. Biol. Rep.* 5:387-405 (1987); Jefferson *et al.*, *EMBO J.* 6:3901-3907 (1987)); an R-locus gene, which encodes a product that regulates the production of anthocyanin pigments (red color) in plant tissues (Dellaporta *et al.*, *Stadler Symposium* 11:263-282 (1988)); a β -lactamase gene (Sutcliffe *et al.*,

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Proc. Natl. Acad. Sci. (U.S.A.) 75:3737-3741 (1978)), a gene which encodes an enzyme for which various chromogenic substrates are known (e.g., PADAC, a chromogenic cephalosporin); a luciferase gene (Ow *et al.*, *Science* 234:856-859 (1986)) a xylE gene (Zukowsky *et al.*, *Proc. Natl. Acad. Sci. (U.S.A.)* 80:1101-1105 (1983)) which encodes a catechol dioxygenase that can
5 convert chromogenic catechols; an α -amylase gene (Ikatsu *et al.*, *Bio/Technol.* 8:241-242 (1990)); a tyrosinase gene (Katz *et al.*, *J. Gen. Microbiol.* 129:2703-2714 (1983)) which encodes an enzyme capable of oxidizing tyrosine to DOPA and dopaquinone which in turn condenses to melanin; an α -galactosidase, which will turn a chromogenic α -galactose substrate.

Included within the terms "selectable or screenable marker genes" are also genes which
10 encode a secretable marker whose secretion can be detected as a means of identifying or selecting for transformed cells. Examples include markers that encode a secretable antigen that can be identified by antibody interaction, or even secretable enzymes which can be detected catalytically. Secretable proteins fall into a number of classes, including small, diffusible proteins detectable, e.g., by ELISA, small active enzymes detectable in extracellular solution
15 (e.g., α -amylase, β -lactamase, phosphinothricin transferase), or proteins which are inserted or trapped in the cell wall (such as proteins which include a leader sequence such as that found in the expression unit of extension or tobacco PR-S). Other possible selectable and/or screenable marker genes will be apparent to those of skill in the art.

In addition to a selectable marker, it may be desirable to use a reporter gene. In some
20 instances a reporter gene may be used with or without a selectable marker. Reporter genes are genes which are typically not present in the recipient organism or tissue and typically encode for proteins resulting in some phenotypic change or enzymatic property. Examples of such genes are provided in K. Wising *et al.* *Ann. Rev. Genetics*, 22, 421 (1988), which is incorporated herein by reference. Preferred reporter genes include the beta-glucuronidase (GUS) of the uidA locus of *E.*
25 *coli*, the chloramphenicol acetyl transferase gene from Tn9 of *E. coli*, the green fluorescent protein from the bioluminescent jellyfish *Aequorea victoria*, and the luciferase genes from firefly *Photinus pyralis*. An assay for detecting reporter gene expression may then be performed at a suitable time after said gene has been introduced into recipient cells. A preferred such assay entails the use of the gene encoding beta-glucuronidase (GUS) of the uidA locus of *E. coli* as
30 described by Jefferson *et al.*, (*Biochem. Soc. Trans.* 15, 17-19 (1987)) to identify transformed cells.

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In preparing the recombinant DNA constructs (vectors) of the present invention, the various components of the construct or fragments thereof will normally be inserted into a convenient cloning vector, e.g., a plasmid that is capable of replication in a bacterial host, e.g., *E. coli*. Numerous cloning vectors exist that have been described in the literature, many of which are commercially available. After each cloning, the cloning vector with the desired insert may be isolated and subjected to further manipulation, such as restriction digestion, insertion of new fragments or nucleotides, ligation, deletion, mutation, resection, etc. so as to tailor the components of the desired sequence. Once the construct has been completed, it may then be transferred to an appropriate vector for further manipulation in accordance with the manner of transformation of the host cell.

A plant recombinant DNA construct (vector) of the present invention may also include a chloroplast transit peptide, in order to target the polypeptide of the present invention to the plastid. The term "plastid" means the class of plant cell organelles that includes amyloplasts, chloroplasts, chromoplasts, elaioplasts, eoplasts, etioplasts, leucoplasts, and proplastids. These organelles are self-replicating, and contain what is commonly referred to as the "chloroplast genome," a circular DNA molecule that ranges in size from about 120 to about 217 kb, depending upon the plant species, and which usually contains an inverted repeat region. Many plastid-localized polypeptides are expressed from nuclear genes as precursors and are targeted to the plastid by a chloroplast transit peptide (CTP), which is removed during the import steps. Examples of such chloroplast polypeptides include the small subunit of ribulose-1,5-biphosphate carboxylase (ssRUBISCO, SSU), 5-enolpyruvateshikimate-3-phosphate synthase (EPSPS), ferredoxin, ferredoxin oxidoreductase, the light-harvesting-complex protein I and protein II, and thioredoxin F. It has been demonstrated that non-plastid polypeptides may be targeted to the chloroplast by use of polypeptide fusions with a CTP and that a CTP sequence is sufficient to target a polypeptide to the plastid. Those skilled in the art will also recognize that various other recombinant DNA constructs can be made that utilize the functionality of a particular plastid transit peptide to import the enzyme into the plant cell plastid depending on the promoter tissue specificity.

The present invention also provides a transgenic plant comprising in its genome an exogenous nucleic acid molecule which comprises: (A) a 5' non-coding sequence which functions in the cell to cause the production of an RNA molecule; which is operably linked to

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(B) an antisense oligonucleotide or polynucleotide or a sense oligonucleotide or polynucleotide of this invention; which is operably linked to (C) a 3' non-translated sequence that functions in said cell to cause termination of transcription.

Transgenic plants of the present invention preferably have incorporated into their genome or transformed into their chloroplast or plastid genomes an exogenous nucleic acid molecule (or "transgene") that comprises a sense oligonucleotide or polynucleotide or an antisense oligonucleotide or polynucleotide. Transgenic plants are also meant to comprise progeny (descendant, offspring, *etc.*) of any generation of such a transgenic plant. A seed of any generation of all such transgenic plants wherein said seed comprises a sense oligonucleotide or polynucleotide or an antisense oligonucleotide or polynucleotide of the present invention is also an important aspect of the invention.

The DNA constructs of the present invention may be introduced into the genome of a desired plant host by a variety of conventional transformation techniques, which are well known to those skilled in the art. Preferred methods of transformation of plant cells or tissues are the *Agrobacterium* mediated transformation method and the biolistics or particle-gun mediated transformation method. Suitable plant transformation vectors for the purpose of *Agrobacterium* mediated transformation include those derived from a Ti plasmid of *Agrobacterium tumefaciens*, as well as those disclosed, e.g., by Herrera-Estrella et al., *Nature* 303:209 (1983); Bevan, *Nucleic Acids Res.* 12: 8711-8721 (1984); Klee et al., *Bio-Technology* 3(7): 637-642 (1985); and EPO publication 120,516. In addition to plant transformation vectors derived from the Ti or root-inducing (Ri) plasmids of *Agrobacterium*, alternative methods can be used to insert the DNA constructs of this invention into plant cells. Such methods may involve, but are not limited to, for example, the use of liposomes, electroporation, chemicals that increase free DNA uptake, free DNA delivery via microprojectile bombardment, and transformation using viruses or pollen.

A plasmid expression vector suitable for the introduction of an antisense oligonucleotide or polynucleotide or a sense oligonucleotide or polynucleotide in monocots using electroporation or particle-gun mediated transformation is composed of the following: a promoter that is inducible or constitutive or tissue-specific; an intron that provides a splice site to facilitate expression of the gene, such as the Hsp70 intron (U.S. Patent Number 5,859,347); and a 3' polyadenylation sequence such as the nopaline synthase 3' sequence (NOS 3'; Fraley et al., Proc.

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Natl. Acad. Sci. USA 80: 4803-4807(1983)). This expression cassette may be assembled on high copy replicons suitable for the production of large quantities of DNA.

An example of a useful Ti plasmid cassette vector for plant transformation is described in US Patent No. 6,147,278 herein incorporated by reference in its entirety, and contains a gene
5 encoding an EPSPS enzyme conferring glyphosate resistance (denominated *aroA*; CP4), which is an excellent selection marker gene for many plants. The gene is fused to the *Arabidopsis* EPSPS chloroplast transit peptide (CTP2) and expressed from the FMV promoter as described therein.

When adequate numbers of cells (or protoplasts) containing an antisense oligonucleotide or polynucleotide or a sense oligonucleotide or polynucleotide of the present invention are
10 obtained, the cells (or protoplasts) can be cultured to regenerate into whole plants. Such regeneration techniques rely on manipulation of certain phytohormones in a tissue culture growth medium, typically relying on a biocide and/or herbicide marker which has been introduced together with the desired nucleotide sequences. Choice of methodology for the regeneration step is not critical, with suitable protocols being available for hosts from Leguminosae (alfalfa,
15 soybean, clover, etc.), Umbelliferae (carrot, celery, parsnip), Cruciferae (cabbage, radish, canola/rapeseed, etc.), Cucurbitaceae (melons and cucumber), Gramineae (wheat, barley, rice, maize, etc.), Solanaceae (potato, tobacco, tomato, peppers), various floral crops, such as sunflower, and nut-bearing trees, such as almonds, cashews, walnuts, and pecans. See, for example, Ammirato et al., Handbook of Plant Cell Culture - Crop Species. Macmillan Publ. Co.
20 (1984); Shimamoto et al., Nature 338:274-276 (1989); Fromm, UCLA Symposium on Molecular Strategies for Crop Improvement, April 16-22, 1990. Keystone, CO (1990); Vasil et al., Bio/Technology 8:429-434 (1990); Vasil et al., Bio/Technology 10:667-674 (1992); Hayashimoto, Plant Physiol. 93:857-863 (1990); and Datta et al., Bio-technology 8:736-740 (1990). Plant regeneration from cultured protoplasts is described in Evans et al., Protoplasts
25 Isolation and Culture, Handbook of Plant Cell Culture, pp. 124-176, MacMillilan 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 et al., Ann. Rev. Plant Phys. 38:467-486 (1987).

30 A transgenic plant formed using *Agrobacterium* transformation methods typically contains a single exogenous gene on one chromosome. Such transgenic plants can be referred to

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as being heterozygous for the added exogenous gene. More preferred is a transgenic plant that is homozygous for the added exogenous gene; *i.e.*, a transgenic plant that contains two added exogenous genes, one gene at the same locus on each chromosome of a chromosome pair. A homozygous transgenic plant can be obtained by sexually mating (selfing) an independent segregant transgenic plant that contains a single exogenous gene, germinating some of the seed produced and analyzing the resulting plants produced for the exogenous gene of interest.

The development or regeneration of transgenic plants containing the exogenous nucleic acid molecule that encodes a polypeptide of interest is well known in the art. Preferably, the regenerated plants are self-pollinated to provide homozygous transgenic plants, as discussed above. Otherwise, pollen obtained from the regenerated plants is crossed to seed-grown plants of agronomically important lines. Conversely, pollen from plants of these important lines is used to pollinate regenerated plants. A transgenic plant of the present invention containing desired polypeptides of the present invention are cultivated using methods well known to one skilled in the art.

Plants that can be made to have larger organ size by practice of the present invention include, but are not limited to, *Acacia*, alfalfa, aneth, apple, apricot, artichoke, arugula, asparagus, avocado, banana, barley, beans, beet, blackberry, blueberry, broccoli, brussels sprouts, cabbage, canola, cantaloupe, carrot, cassava, cauliflower, celery, cherry, cilantro, citrus, clementines, coffee, corn, cotton, cucumber, Douglas fir, eggplant, endive, escarole, eucalyptus, fennel, figs, gourd, grape, grapefruit, honey dew, jicama, kiwifruit, lettuce, leeks, lemon, lime, loblolly pine, mango, melon, mushroom, nut, oat, okra, onion, orange, an ornamental plant, papaya, parsley, pea, peach, peanut, pear, pepper, persimmon, pine, pineapple, plantain, plum, pomegranate, poplar, potato, pumpkin, quince, radiata pine, radicchio, radish, raspberry, rice, rye, sorghum, southern pine, soybean, spinach, squash, strawberry, sugarbeet, sugarcane, sunflower, sweet potato, sweetgum, tangerine, tea, tobacco, tomato, turf, a vine, watermelon, wheat, yams, and zucchini.

The present invention also provides cells of the transgenic plants of the present invention which could be used for regenerating a plant or any organ of plant with present invention.

The present invention also further provides a method for altering a specific organ or organs in a plant to generate smaller plant or plant organ, the method comprising the steps of: a) introducing into the genome of the plant an exogenous nucleic acid molecule comprising in the

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5' to 3' direction i) a promoter that functions in the cells of said plant, said promoter operably linked to; ii) an antisense oligonucleotide or polynucleotide or a sense oligonucleotide or polynucleotide of the present invention, said antisense oligonucleotide or polynucleotide operably linked to; iii) a 3' non-translated nucleotide sequence that functions in said cells of said plant to cause transcriptional termination; b) obtaining transformed plant cells containing the exogenous nucleic acid molecule of step (a); and c) regenerating from said transformed plant cells a transformed plant in which the expression of an endogenous gene of the present invention is suppressed or inhibited.

The following examples are provided to better elucidate the practice of the present invention and should not be interpreted in any way to limit the scope of the present invention. Those skilled in the art will recognize that various modifications, truncations, etc., can be made to the methods and genes described herein while not departing from the spirit and scope of the present invention. In the following examples references are made to proprietary databases and proprietary libraries, e.g., of DNA clones, available to the inventors from Monsanto Company.

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EXAMPLES

EXAMPLE 1

Stock Plant Material and Growth Conditions

Arabidopsis thaliana var Columbia seeds were obtained from *Lehle seeds* (LEHLE SEEDS 1102 South Industrial Blvd., Suite D, Round Rock TX 78681 USA). For growing seeds into plants, 2.5 inch pots are prepared with soil covered with bridal veil or a mesh screen, making sure that the soil is not packed too tightly and the mesh is in contact with the soil surface (this ensures that the germinating seedlings will be able to get through the mesh. Seeds are sown and covered with a germination dome. Seeds are vernalized for 3-4 days. Plants are grown under conditions of 16 hours light / 8 hours dark at 20-22° C, 70% humidity. They are watered twice weekly, and fertilize from below with 1/2 X (half of the strength recommended by the manufacturer) Peters 20-20-20 fertilizer (from Hummert International, Earth City, MO). Micronutrients are added (Hummert's Dyna-grain Soluble Trace Elements) (in full strength recommended by the manufacturer) every other week. After about 1-2 weeks, dome is removed and pots are thinned to

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one or two plants per pot. Clip the primary bolt, when it develops, to encourage more secondary bolt formation. In 5-7 days the plants will be ready for infiltration.

Isolation of Gene and its modification: Senescing leaves were removed from plants grown as indicated in above Example 1. Senescing leaves were flash frozen in liquid nitrogen until ready for isolation of RNA. RNA was prepared from senescent *Arabidopsis* leaves by the Trizol method (Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A.) essentially as recommended by the manufacturer. SAG13 cDNA was isolated by reverse transcription from the above senescent leaf RNA using Superscript II kit (Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A.) according to the manufacturer's directions. To isolate DNA molecules of the present invention, two gene specific primers, MF16 5'SAG13 and MF17 3'SAG13, were designed based on the *SAG 13* sequence information (AF192276) from Genbank and custom synthesized by Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A. The sequence of MF16 5'SAG13 is ATA TTT AAC AAG CCA TGG CAA AGG A, and MF17 3'SAG13 is ATA TGT GTT TGA ATT CAT AGT CTT GAA identified as SEQ ID NO: 55 and SEQ ID NO: 56 respectively in the sequence list, which annealed on SAG 13 gene to introduce Nco 1 site at 5'end and Eco R1 site at 3'end of the gene. PCR was then performed to amplify the *SAG-13* cDNA using the above prepared cDNA as the template, and MF16 5'SAG13 and MF17 3'SAG13 as the primers. The thermal cycling conditions were as follows: 94°C, 40 second, followed 30 cycles of 94°C, 25 seconds; 55°C, 30 seconds and 68°C, 2 minutes 30 seconds (All reagents and equipment for PCR can be procured from Applied Biosystems, 850 Lincoln Center Drive, Foster City, CA 94404, USA). The amplified cDNA was purified by gel-electrophoresis to obtain the gene identified as SEQ ID NO: 1.

All other sequences shown in table from Example 8 are isolated from different plant species by designing appropriate PCR primers based on the sequence information provided in table of Example 8. For isolating these sequences, total RNA is isolated from appropriate crop and other desired plant species by pooling tissues of different developmental stages of all vegetative and reproductive organs. Sequences can be cloned out from total RNA by methodology shown in above paragraph. In order to isolate genes of the invention from microorganisms DNA is isolated from the desired microorganism. Isolation of DNA from microorganism is well known in the art (Sambrook, *et al.*, In: *Molecular Cloning, A Laboratory Manual, 2nd Edition*, Cold Spring Harbor Press, Cold Spring Harbor, New York 1989, herein

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incorporated by reference in its entirety). This DNA along with oligonucleotide PCR primers can be used in a polymerase chain reaction by any one skilled in the art to isolate genes of the invention.

When the amplified product SEQ ID NO: 1 was sequenced, it was discovered to have an
5 extra "T" between position 622-628, as compared to SEQ ID NO: 3. Addition of an extra "T" in
the amplified product caused the generation of termination codon at position 211 of SEQ ID NO:
4 and a change of amino acid at position 210 from aspartic acid (D) to arginine (R). It is known
in the art that polymerase chain reaction can cause point mutations (Cline J, Braman JC, Hogrefe
HH; *Nucleic Acids Res.*; 24(18):3546-51, 1996) as shown between SEQ ID No: 1 and SEQ ID
10 NO: 3. However such a mutation can be created within an isolated nucleic acid molecule at a
desired position by a number of methods know in the art (Shortle D. et al., *Annu. Rev. Genet.* 15:
265, 1981; Itakura K. et al., *Ann Rev. Biochem* 53: 323, 1984; Botstein D. & Shortle D., *Science*
229: 1193, 1985; Smith M., *Ann. Rev. Genet.* 19: 423 1985; and Sambrook, *et al.*, In: *Molecular*
15 *Cloning, A Laboratory Manual, 2nd Edition, Cold Spring Harbor Press, Cold Spring Harbor,*
New York 1989, herein incorporated by reference in its entirety). These mutations can also be
introduced by causing a point mutation or site directed mutagenesis in nucleotide SEQ ID NO: 3
by using commercially available kits such as QuickChangeTM Site Directed Mutagenesis Kit
from Stratagene (11011 North Torrey Pines Road, La Jolla, CA 92037) to result in a peptide with
20 about 90% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a
stop codon at amino acid position 241 ± 3 , or to result in a peptide with about 80% of the N-
terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino
acid position 214 ± 3 , or to result in about 70% of the N-terminal of SEQ ID NO: 4 by altering a
nucleotide sequence to encode a stop codon at amino acid position 188 ± 3 , or to result in about
60% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop
25 codon at amino acid position 161 ± 3 , or to result in about 50% of the N-terminal of SEQ ID
NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 134 ± 3 of
the reading frame. These mutations can also be introduced *in situ* by causing point mutation or
site directed mutagenesis in nucleotide SEQ ID NO: 3 by using *in situ* mutagenesis technology
provided by Valigen Inc. (Newtown PA. USA) (US Patent Number 6,211,351; US Patent
30 Number 6,271,360, WO 01/24615 A1, and WO 01/25460 A2, herein incorporated by reference

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in its entirety) to result in a peptide with about 90% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 241 ± 3 , or to result in a peptide with about 80% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 214 ± 3 , or to result in about 70% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 188 ± 3 , or to result in about 60% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 161 ± 3 , or to result in about 50% of the N-terminal of SEQ ID NO: 4 by altering a nucleotide sequence to encode a stop codon at amino acid position 134 ± 3 of the reading frame. Additional mutations can be introduced to provide a translational product that is larger or smaller than SEQ ID NO: 1. It can be determined that these modified polynucleotide sequences will provide a desired phenotype without undue experimentation

EXAMPLE 2

Genetic elements of cloning vector for expressing protein in plants (Figure 1 and 4)

The DNA constructs used are double border plant transformation constructs that contain DNA segments that provide replication function and antibiotic selection in bacterial cells, for example, an *E. coli* origin of replication such as *ori322*, a broad host range origin of replication such as *oriV* or *oriRi*, and a coding region for a selectable marker such as Spc/Str that encodes for Tn7 aminoglycoside adenytransferase (*aadA*) conferring resistance to spectinomycin or streptomycin, or a gentamicin (Gm, Gent) selectable marker. For plant transformation, the host bacterial strain is *Agrobacterium tumefaciens* ABI or LBA4404.

The genetic elements of the DNA constructs are assembled to have in operable linkage a promoter that functions in plants. Additionally an antibiotic or herbicide marker cassette, an epitope tag (For example Flag® peptide catalog number F-3290, SIGMA, P.O. Box 14508 St. Louis, MO 63178 USA) at the 3' termination region of gene of interest is included in the DNA construct. The multiple cloning site in this DNA construct encodes BglII, NcoI, EcoRI, Sall, and XhoI. The epitope tag region was encoded with Sall and XhoI restriction sites for optional removal of the epitope tag. The NcoI site encodes a Kozak sequence for efficient translation of the protein products. DNA constructs used in the method of the current invention comprise any promoter known to function to cause the transcription in plant cells and any antibiotic tolerance

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encoding polynucleotide sequence known to confer antibiotic tolerance to plant cells. The antibiotic tolerance polynucleotide sequences include, but are not limited to polynucleotide sequences encoding for proteins involved in antibiotic tolerance to kanamycin, neomycin, hygromycin, and other antibiotics known in the art. Antibiotic tolerance gene in such a vector can be replaced by herbicide tolerance encoding for 5-enolpyruvylshikimate-3-phosphate synthase (EPSPS, described in U.S. Patent Numbers 5,627,061, and 5,633,435, herein incorporated by reference in its entirety; Padgett *et al.* (1996) *Herbicide Resistant Crops*, Lewis Publishers, 53-85, and in Penaloza-Vazquez, *et al.* (1995) *Plant Cell Reports* 14:482-487) and *aroA* (U.S. Patent Number 5,094,945) for glyphosate tolerance, bromoxynil nitrilase (Bxn) for Bromoxynil tolerance (U.S. Patent Number 4,810,648), phytoene desaturase (*crtI* (Misawa *et al.*, (1993) *Plant Journal* 4:833-840, and (1994) *Plant Jour* 6:481-489) for tolerance to norflurazon, acetohydroxy acid synthase (AHAS, Sathasiivan *et al.* (1990) *Nucl. Acids Res.* 18:2188-2193) and the *bar* gene for tolerance to glufosinate (DeBlock, *et al.* (1987) *EMBO J.* 6:2513-2519. Herbicides for which transgenic plant tolerance has been demonstrated and the method of the present invention can be applied include, but are not limited to: glyphosate, glufosinate, sulfonylureas, imidazolinones, bromoxynil, delapon, cyclohezanedione, protoporphyrionogen oxidase inhibitors, and isoxaslutole herbicides.

Genetic elements of transgene DNA constructs used for plant transformation and expression of transgenes in plants include, but are not limited to: the P-E35S promoter (U.S. Patent No.5,539,142, 5,196,525, 5,322,938 and 5,164,316 herein incorporated by reference in its entirety). P-E35S promoter can be replaced by P-CaMV.35S promoter (U.S. Patent No. 5,858,742, herein incorporated by reference in its entirety), or by enhanced P-CaMV.35S from Cauliflower mosaic virus containing a duplication of the -90-300 region as described in U. S. Patent No. 5,424,200, herein incorporated by reference in its entirety; or the Figwort mosaic virus promoter, P-FMV, as described in U. S. Patent No. 5,378,619, herein incorporated by reference in its entirety; or the P-AtEF1a (P-AtEF1 or EF1a) a promoter region from the *Arabidopsis thaliana* elongation factor gene 1a; the Gbox10 and Gbox11 motif (Fumiharuru *et al.*, (1999) *Plant J.* 18:443-448); or the DC3 promoter region from carrot (Seffens *et al.*, *Develop. Genet.* 11:65-76); or the TP12 promoter (GenBank accession no. U68483); DNA molecules encoding plastid transit peptides, for example, the *Arabidopsis* EPSPS chloroplast transit peptide, At.CTP2 as described in U. S. Patent No. 5,633,435, herein incorporated by reference in its

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entirety. The method of the present invention enables one of skill in the art of plant molecular biology to design and assemble plant expression cassettes that contain promoters of known and unknown function. The genetic elements of the DNA construct further comprise 5' leader polynucleotides for example, the Hsp70 non-translated leader sequence from *Petunia hybrida* as described in U. S. Patent No. 5,362,865, herein incorporated by reference in its entirety. The genetic elements further comprise herbicide tolerance genes that include, but are not limited to, for example, the *aroA*:CP4 coding region for EPSPS glyphosate resistant enzyme isolated from *Agrobacterium tumefaciens* (AGRTU) strain CP4 as described in U. S. Patent No. 5,633,435, herein incorporated by reference in its entirety. The genetic elements of the DNA construct further comprise 3' termination regions that include, but are not limited to, the E9 3' termination region of the pea RbcS gene that functions as a polyadenylation signal; the *nos* is the 3' end of the nopaline synthase gene that functions as a polyadenylation signal. The genetic elements of the DNA construct further comprise the right border (RB) and left borders (LB) of the Ti plasmid of *Agrobacterium tumefaciens* octopine and nopaline strains.

15 EXAMPLE 3

Cloning of isolated DNA Molecule: Amplified and purified product SEQ ID NO: 1 from Example 1 was digested by Nco1 and EcoR1 restriction enzymes (BRL/Life Technologies, Inc., Gaithersburg, MD). The digested product was purified again by gel electrophoresis before ligating to binary vector pMON23435 that had been linearized by Nco 1 and Eco R1 and T4 DNA ligase (BRL/Life Technologies, Inc., Gaithersburg, MD). The ligation reaction was performed according to the manufacturer's instructions. The resulting plasmid was confirmed by restriction mapping and sequencing. After ligation of the Nco-EcoR1 fragment of SEQ ID NO: 1 into vector pMON23435, the new construct was referred to as construct pMON57521. Construct pMON57521 was transformed in *Arabidopsis* plants by *Agrobacterium* mediated transformation procedure.

Amplified and purified product SEQ ID NO: 1 can also be cloned in antisense orientation in appropriate cloning sites of vector pMON23435 to express the opposite strand of SEQ ID NO:1 by one skilled in the art.

EXAMPLE 4

30 Cloning of DNA molecules of the present invention: This example illustrates how all other DNA molecules as shown in table from example 9 are isolated from different plant species

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by designing appropriate PCR primers based on the DNA sequence information provided in the table of Example 8. For isolating these DNA molecules one skilled in the art will isolate total RNA from a crop or other desired plant species by pooling tissues of different developmental stages of all vegetative and reproductive organs. DNA molecules are cloned out from total RNA
5 by methodology shown in Example 2. In order to isolate genes of the invention from microorganisms one will have to isolate DNA from desired microorganism. Isolation of DNA from microorganism is well known in the art (Sambrook, *et al.*, In: *Molecular Cloning, A Laboratory Manual, 2nd Edition, Cold Spring Harbor Press, Cold Spring Harbor, New York* 1989, herein incorporated by reference in its entirety). This DNA along with oligonucleotide
10 PCR primers can be used in a polymerase chain reaction by any one skilled in the art to isolate genes of the invention.

Design of primers and reaction conditions are determined as described in the art. (PCR Strategies, Edited by Michael A. Innis; David H. Gelfand; John J. Sninsky; Academic Press 1995 and PCR Protocols, A Guide to Method and Applications, Edited by Michael A. Innis; David H.
15 Gelfand; John Sninsky; and Thomas J. White, Academic Press 1990, herein incorporated by reference in its entirety). All reagents for isolating sequences of the invention can be procured from Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A. Isolated DNA molecule sequences can be cloned at appropriate cloning sites in sense or antisense orientation of a plant expression vector shown in Example 3 or a similar vector capable of ectopically expressing the
20 gene of interest of the present invention in sense or antisense orientation as known in the art.

EXAMPLE 5

This example illustrates how *Agrobacterium* cells are transformed and how transformed cells are cultured.

Transformation:

- 25 1. Electroporate 2 μ l of DNA construct into 20 μ l of ABI competent cells;
2. Pipette transformed cells directly onto LB plates containing Spectinomycin (75 μ g/ml), Kanamycin (50 μ g/ml), Chloramphenicol (25 μ g/ml). Add 50 μ l of SOC media to plate and spread;
3. Incubate plated transformation at 28°C for 2 days (or can grow over weekend).

ABI Cell Culture:

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1. Pick 3 colonies per ABI plate and grow each in 4ml LB media containing Spectinomycin (75µg/ml), Kanamycin (50µg/ml), Chloramphenicol (25µg/ml);
2. Incubate 4 ml cultures of at 28°C, shaking, for 2 days. (Culture tubes should be at an angle).

Glycerol Stocks, & DNA Preps:

- 5 1. Make three 1 ml ABI glycerol stocks per 4 ml culture, using 500µl of culture and 500µl of 40% glycerol. Freeze and store at -80°C.
2. Miniprep remaining culture (about 2.5ml), using a Qiagen miniprep kit and protocol (Qiagen Genomics, Inc., Seattle, WA), ensuring add PB buffer wash step and EB buffer (10mM Tris-Cl, pH 8.5) to 70°C before eluting DNA from column. The resulting volume per miniprep
10 sample should be 50µl.

Digest Confirmation:

1. Using the Pollux and construct maps, choose two digests to perform: one to verify insert integrity, one to verify vector integrity (will need to refer to plasmid maps);
2. Digest 17µl miniprep DNA per digest, resulting in a final digest volume of 20µl;
- 15 3. Run 20µl of each digest on 1% agarose gel vs. 1Kb DNA ladder; and
4. For 2 of 3 confirmed clones, streak LB plates containing Spectinomycin (75µg/ml), Kanamycin (50µg/ml), Chloramphenicol (25µg/ml) from ABI glycerol stocks and allow to grow at 28°C for 2 days (or can grow over weekend).

All other reagents used in example 4 can be procured from Sigma Chemical Company.
20 Saint Louis, MO, USA

EXAMPLE 6

This example demonstrates how to transform *Arabidopsis* plants with gene constructs of present invention *Arabidopsis* plants may be transformed by any one of many available methods. For example, *Arabidopsis* plants may be transformed using *in planta* transformation method by
25 vacuum infiltration (see, Bechtold *et al.*, *In planta Agrobacterium* mediated gene transfer by infiltration of adult *Arabidopsis thaliana* plants. CR Acad. Sci. Paris Sciences de la vie/life sciences 316: 1194-1199 (1993), herein incorporated by reference in its entirety). Plants can be grown as described in Example 1.

Agrobacterium Preparation (Small scale and Large scale cultures):

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Agrobacterium strain ABI is streaked onto an LB plate containing Spectinomycin 100 mg/L, Streptomycin 100 mg/L, Chloramphenicol 25mg/L, and Kanamycin 50mg/L (denoted SSCK). Two days prior to infiltration, a loop of *Agrobacterium* is placed into a tube containing 10 mls LB/SSCK and put on a shaker in the dark at 28°C to grow overnight. The following day, the *Agrobacterium* is diluted 1:50 in 400 mls of bacterial grown medium such as SSCK and put on a shaker at 28°C to grow for 16-20 hours.

Infiltration

Harvest the *Agrobacterium* cells by pouring into a 500 ml centrifuge bottle and spinning at 3500 rpm for 20-25 minutes. Pour off the supernatant. Dry the pellet and then resuspend in 25 ml Infiltration Medium (MS Basal Salts 0.5%, Gamborg's B-5 Vitamins 1%, Sucrose 5%, MES 0.5 g/L, pH 5.7) with 0.44 nM benzylaminopurine (BAP) (10 µl of a 1.0 mg/L stock in DMSO per liter) and 0.02% Vac-In-Stuff (Silwet L-77) from Lehle Seeds (Round Rock, TX). The BAP and Silwet L-77 are added fresh the day of infiltration. Add 200 µl of Silwet L-77, and 20 µl of BAP (0.5 mg/L stock). Using Infiltration Medium as your blank, take the OD₆₀₀ of a 1:10 dilution of the *Agrobacterium* suspensions. Calculate the volume needed for 400 ml of *Agrobacterium* suspension/infiltration medium, OD₆₀₀ = 0.6, for the vacuum infiltration.

$$\text{Equation: } \frac{(\text{final volume}) * (\text{final OD600})}{\text{OD600}} = \text{Volume needed for final OD600 of 0.6}$$

Place resuspended culture in a Rubbermaid container inside a vacuum dessicator. Invert pots containing plants to be infiltrated into the solution so that the entire plant is covered, including the rosette, but not too much of the soil is submerged. Soak the plants with water for at least 30 min. prior to infiltration. (This keeps the soil from soaking up the *Agrobacterium* suspension).

Draw a vacuum of ~23-27 in. Hg for 10 min. Quickly release the vacuum. Briefly drain the pots, place them on their sides in a diaper-lined tray, cover the tray with a dome to maintain humidity, and return to growth chamber. The following day, uncover the pots, set them upright, and remove the diaper. Do not water plants for ~5 days. After the 5 days are up, allow the plants to be watered and to continue to grow under the same conditions as before. (The leaves that were infiltrated may degenerate but the plant should survive until it is finished flowering).

Harvesting and Sterilizing Seed

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Cone the plants, individually, by using the Lehle Aracons (Lehle Seeds, Round Rock, TX) approximately 2 weeks after infiltration. After all of the seed is matured and has set (~ 4 weeks post-infiltration), remove the plants from water to dry down the seeds. Approximately 2 weeks later harvest the seeds by cutting the branches below the cone. Clean the seed by using a sieve to catch the silique and branch material and allow the seed to go through. Place the seed in an envelope or in 15ml conical tubes.

Transfer desired amount of seeds to 15ml conical tubes prior to sterilization. Loosen the lid to the conicals and place them on their side in a vacuum dessicator with a beaker containing 400 ml of bleach Clorox (Clorox Company, Oakland, CA) and 4 ml of Hydrochloric Acid. (Add the HCl to the Clorox in a fume hood). Pull a vacuum just to seal the dessicator, and close the suction (i.e. so that the dessicator is still under a vacuum but the vacuum is not still being directly pulled) for ~ 16 hrs. After sterilization, release the vacuum and place tubes containing seed in a sterile hood (keep caps loose so gas can still be released).

Plate ("sprinkle") the seed on selection plates containing MS Basal Salts 4.3 g/L, Gamborg's B-5 (500 X) 2.0 g/L, Sucrose 10 g/L, MES 0.5 g/L, and 8 g/L Phytagar (Life Technologies, Inc., Rockville, MD) with Carbenicillin 250mg/L, Cefotaxime 100 mg/L. Selection levels will either be kanamycin 60 mg/L, Glyphosate 60 μ M, or Bialaphos 10mg/L.

A very small amount of seed can be first plated out to check for contamination. If there is contamination, re-sterilized seeds for ~ 4 more hours and check for contamination again. The second sterilization is usually not necessary, but sometimes the seed harbors a fungal contaminant and repeat sterilizations are needed. (The sterilization duration generally is shorter than 16 hours because of significantly decreased germination rates starting at 24 hr. sterilization duration). Seal plates with parafilm and place in a cold room to vernalize for ~ 2-4 days. After seeds are vernalized, place in percival with cool white bulbs.

25 **Transfer to Soil**

After 5-10 days at ~26°C and a 16/8 light-dark cycle, the transformants will be visible as green plants. After another 1-2 weeks, plants will have at least one set of true leaves. Transfer plants to soil, cover with a germination dome, and move to a growth chamber with normal *Arabidopsis* growth conditions. Keep covered until new growth is apparent (usually 5-7 days).

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EXAMPLE 7

Following subsections of Example 7 are used for describing phenotypic changes after transformation and growth of *Arabidopsis* plants as described in Example 6. Three different events obtained from transformation of pMON57521(Figure 2) into *Arabidopsis thaliana* ecotype Columbia were selected and grown side by side with wild type plants. Growth conditions were 16 hr light, 8 hr night, 21 degrees Centigrade and 70% relative humidity. Plants were observed at all growth stages and photographed using an Olympus camera C-2500 L (Olympus America Inc., 2 Corporate Center Drive, Melville, NY 11747) as described by the manufacturer. Microscopic images of plant organ were acquired after proper dissection on a stereoscopic microscope Nikon SMZ 1500 (NIKON, 1300 Walt Whitman Road, Melville, NY 11747) equipped with MagnaFire Digital imaging system. Image quantitation was done as per manufacturer's instructions by using MagnaFire Digital imaging system's software s Pro™ or Lucis™ softwares (Optronics, 175 Cremona Drive, Goleta, CA 93117). The three events showed high phenotypic consistency in preliminary experiments and hence only two events were chosen for further downstream work.

Plants transformed with Construct pMON57521 were shown to exhibit 2 to 3 times larger flower and floral organs such as stamen and pistils when compared to the wild type, non transformed plants of the same species. Transgenic plants were also observed to have at least 2 times more lateral roots and 2 to 3 times thicker stem size when compared to the lateral roots and stems of wild type, non transformed plants of the same species. Approximately 2 fold increased trichome numbers and distribution were also observed. Over expression of the polypeptide encoded by the sequence in the pMON57521 construct of the present invention in the transformed plants, yielded plants that exhibited a 100 % increase in individual seed size and weight. Approximately 20 % increase in seed yield/plant was also observed for plants transformed with pMON57521 when compared to seed yield/plant of wild type, non transformed plants of the same species. The lines were followed up to 6 generations after transformation and appeared to be highly consistent in maintaining the observed phenotypic changes.

(A) Seed Size: Seeds from transformed plants exhibit increased seed size that was approximately twice as large as the seeds from wild type plants as shown in following table. Image sizes were determined by the Pro™ or Lucis™ software (Optronics, 175 Cremona Drive,

Goleta, CA 93117) based on the image pixel value of wild type and transformed plants' seed under the same resolution. Plant lines 8752-1, 8752-2, 8752-6, and 8752-7 in Table 1 correspond to different transgenic plant lines produced by transforming wild *Arabidopsis* plants with pMON57521.

5 Table 1: Seed Size

	Seed size	Average	Std deviation
WT	1820	1873	89
	1847		
	1904		
	1784		
	2013		
8752-1	3762	3627	191
	3441		
	3812		
	3400		
	3720		
8752-6	3700	3685	172
	3802		
	3872		
	3623		
	3428		
8748-2	3675	3664	96
	3784		
	3724		
	3543		
	3597		
8748-7	3712	3754	88
	3625		
	3771		
	3821		
	3845		

(B) Seed Weight: The seed from plants containing pMON57521 lines was also found to be heavier than WT seed. Seed weight was measured from 2 lines produced from event 8752 (8752-1 and 8752-6) and 2 lines from event 8748 (8748-2 and 8748-7) and compared with wild

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type. Shown below is a representative seed weight analysis. For each line, 3 replications were measured (i.e. 50 seeds were counted three times and measured each time). This analysis was repeated with different seed counts (eg. 100 seed or 150 seed) and was found to be highly reproducible. For example, weight per 100 seed is similarly high ~ 0.0025 gm. for wild type and 0.0046 gm. for seeds from plants transformed with sequence of the present invention. Data was also highly reproducible within a given line and across lines. The average weight of a single seed from a line transformed with sequence of the present invention was extrapolated to be ~0.048 mg as opposed to 0.026 mg for a WT seed. Plant lines 8752-1, 8752-6, 8748-2, and 8748-7 in Table 2 correspond to different transgenic plant lines produced by transforming wild *Arabidopsis* plants with pMON57521.

Table 2: Seed Weight

	Weight (gm./50 seed)	Average	Std deviation
WT	0.0013	0.001333	5.7735E-05
	0.0014		
	0.0013		
8752-1	0.0024	0.0024	1E-04
	0.0025		
	0.0023		
8752-6	0.0024	0.002433	5.7735E-05
	0.0025		
	0.0024		
8748-2	0.0025	0.0024	1E-04
	0.0023		
	0.0024		
8748-7	0.0025	0.0024	1E-04
	0.0023		
	0.0024		

(C) Seed Number: Lines over expressing sequence of the present invention were found to have fewer seed per silique as compared to wild type plants. The average number of seed /silique of equivalent maturity was 34 for transgenic plants of the present invention as compared

to 52 for wild type plants. The analysis was repeated three times with highly reproducible results within and between events.

Table 3: Seed Number

	Number /silique	Average	Std deviation
WT	46	52	7.402702
	62		
	50		
	57		
	55		
	42		
8752-1	34	34	4.195235
	32		
	30		
	30		
	38		
	40		

5 **(D) Seed Yield:** There was an increase of > 20% seed yield in transgenic plants of the present invention. Seed yield when measured in terms of weight of total seeds per plant.

Table 4: Seed Yield

	Yield(gm./plant)	Average	Std deviation
WT	0.31	0.274	0.024129
	0.25		
	0.28		
	0.27		
	0.25		
	0.30		
	0.26		
	0.28		
	0.3		
	0.24		
	0.25		
	0.29		
8752-1	0.33	0.336667	0.021034
	0.31		

	0.37		
	0.35		
	0.37		
	0.33		
	0.33		
	0.36		
	0.33		
	0.33		
	0.31		
	0.32		

EXAMPLE 8

This example describes changes in the branching pattern after transformation and growth of Arabidopsis plants as described in Example 6.

5 Arabidopsis plants over-expressing polypeptide molecules corresponding to SEQ ID NO: 2 appear to have more branches than wild type plants. This results in a net increase in silique number. Although each transgenic silique appears to have fewer seed than an equivalent wild type silique, the increase in branching appears to compensate resulting in a net increase in overall seed yield. The branching pattern of Arabidopsis over-expressing lines was compared to wild
 10 type plants and the transgenic plants were shown to have more branches than WT plants. All measurements were taken at growth stage 6.5 which corresponds to an Arabidopsis growth stage where the plant is still actively growing (mid-flowering). By growth stage 6.9 the plants have started senescing and flower production has stopped with < 5 open flowers. Thus the growth stage 6.5 gives a fairly accurate estimate of the growth rate of the plant prior to senescence. An
 15 average of the silique number at growth stage 6.5 was estimated for both WT and transgenics and total silique number is shown in the following table. This indicates again that silique number was at least 28% greater in lines over-expressing the polypeptide molecules corresponding SEQ ID NO: 2 (transformed lines) as compared to wild type lines at the same growth stage. As the plant proceeds to senescence, the relative increase appears to stay
 20 consistent and together with the increase in seed weight in the transgenic plant, contributes to the overall increase in seed yield per plant. Growth stage estimation was based on "Growth stage-

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based phenotypic analysis of Arabidopsis: A model for high throughput functional genomics in plants. Plant Cell. 13(7):1499, 2001”

Table 5 Number of Silique. WT is wild type plant and Transformed line is transgenic plant line expressing polypeptide molecules corresponding to SEQ ID. NO: 2.

5

WT siliques	Average WT siliques	Transformed line siliques	Average Transformed line siliques
88	88	124	113
87		123	
81		126	
78		112	
84		104	
80		102	
96		112	
98		113	
100		105	

EXAMPLE 9

This example describes increase of seeds in transgenic soybean plants expressing polypeptide molecule corresponding to SEQ ID NO: 2.

10 Soybean plants were transformed with pMON73955 (Figure 7) to constitutively express SEQ ID NO: 2. Soybean transformation is performed essentially as described in WO 00/42207, herein incorporated by reference in its entirety. R1 seed from 10 events out of 44 events were advanced in Puerto Rico (PR) based on the gene copy number in plants. Soybean transformation was performed essentially as described in WO 00/42207, herein incorporated by reference in its
 15 entirety. Preliminary data indicates that 1 of the 10 events in PR showed a phenotype similar to that seen for the Arabidopsis over-expression of SEQ ID NO: 2 (pMON 57521 Figure 2) i.e. more pods and more branches as well as short, fat pods with fewer seed. More two-seeded pods were observed and the seed was larger than seed from wild type and negative plants (but seed size was not doubled as seen in Arabidopsis). In addition the positives (transgenic line
 20 expressing genes of interest) from this event “A” produced more seed than the negatives (transgenic line NOT expressing genes of interest) as shown in Table 7. Table 6 shows the seed weight of individual plants (events) from the R1 generation and seed weight in the R2

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generation. R2 seed weight in this table is an average of seed weight of all lines from the original R1 event. All numbers are expressed as a percentage of wild-type (WT). Detailed R2 seed weight and seed number data from single lines (positives versus negatives) of the best performing event "A" is shown in Table 7. In general a good correlation is observed between positives and negatives for seed size and seed yield.

Table 6: R1 and R2 seed weights of 10 independent soybean transgenic events expressing SEQ ID NO: 2. Values are expressed as a percentage of wild-type where WT = 100%

Pedigree	% R1 Seed weight	% R2 Seed weight
Line A-1	154%	163%
Line A-2	108%	80%
Line A-3	97%	109%
Line A-4	105%	110%
Line A-5	91%	83%
Line A-6	105%	78%
Line A-7	102%	95%
Line A-8	108%	183%
Line A-9	105%	115%
Line A-10	105%	116%

Table 7: R2 seed weights of individual lines from the large-seeded transgenic event "A" expressing SEQ ID NO: 2.

Line	Total number of seed produced	Corresponding seed weight (gms)	Seed weight of 100 seed (gms)
A-1 POS	144	28	19.44
A-1 POS	76	15	19.74
A-1 POS	126	30	23.81
A-1 POS	124	21	16.94
A-1 POS	166	34	20.48
A-1 POS	118	23	19.49
A-1 NEG	98	17	17.35
A-1 NEG	50	11	22
A-1 NEG	97	16	16.49
A-1 NEG	66	12	18.18
A-1 NEG	58	10	17.24
A-1 NEG	75	15	20
A-1 NEG	92	16	17.39

A-1 NEG	75	14	18.67
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EXAMPLE 10

This example describes over expression of the protein in bacterial cells for purification of the protein so as to screen for the activity of polypeptide molecules.

5 **(A) Cloning of the nucleotide molecules of the present invention for expression of corresponding peptides.**

pMON57521 (Figure 2), pMON73963 (Figure 3), or *Nostoc punctiforme* (*Nostoc*) genomic DNA is used as a DNA template source for polymerase chain reaction amplification (PCR) of DNA for cloning into bacterial cells so as to express the *Arabidopsis thaliana* protein corresponding to
10 SEQ ID NO: 2, SEQ ID NO: 4, or *Nostoc* protein corresponding to SEQ ID NO: 50. Design of primers and PCR reaction conditions are determined as described in the article PCR Strategies, Edited by Michael A. Innis; David H. Gelfand; & John J. Sninsky; Academic Press 1995 and PCR Protocols, A Guide to Method and Applications, Edited by Michael A. Innis; David H. Gelfand; John J. Sninsky; & Thomas J. White Academic Press 1990, herein incorporated by
15 reference in its entirety. All reagents for performing the PCR reaction can be procured from Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A. The thermocycler needed for performing PCR reaction was procured from Applied Biosystems (Perkin-Elmer Corp., Applied Biosystems Div., Foster City, CA). The polynucleotide sequence of the amplicon produced from *Arabidopsis* is shown in SEQ ID NO:1 and SEQ ID NO: 3. PCR primer pairs SEQ ID NO: 57 and SEQ ID NO: 58 were used for performing the reaction to obtain the *Arabidopsis* amplicon.
20 *Arabidopsis* amplicons were subcloned into a pET-28b vector (*E coli* expression vector, Novagen, Madison, WI, USA) and were sequenced by using sequencing primers SEQ ID NO: 59 and SEQ ID NO: 60 to confirm the sequence of the cloned polynucleotide molecule. Similarly *Nostoc* amplicons were generated by using *Nostoc* genomic DNA as template with primers pairs
25 SEQ ID NO: 61 and SEQ ID NO: 63 or SEQ ID NO: 62 and SEQ ID NO: 63. *Nostoc* amplicons were also subcloned in to the pET-28b vector and were sequenced by using sequencing primers SEQ ID NO: 64 and SEQ ID NO: 65 to confirm the polynucleotide sequence of the cloned polynucleotide molecule. Primer Sequence for amplification and sequencing of molecules of invention from *Arabidopsis* and *Nostoc* are shown in Figure 10. These constructs contained a
30 candidate from *Arabidopsis* SEQ ID NO: 3 and the closest *Nostoc* homologue SEQ ID NO: 49,

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both with and without an N-terminal His-tag. An N-terminal His-tag was chosen based on the crystal structures (Nakajima et al. *PNAS* 95, 4876, 1998) of the related proteins (tropinone reductases) which suggested that the N terminus would not interfere with the dimerization domain. The resultant vectors pMON 63132 (Figure 4, Nostoc Histag), pMON 63133 (Figure 5, *A. thaliana*), pMON 63134 (Figure 6, *A. thaliana* Histag), and pMON 63135 (Figure 7 Nostoc) were used in the following examples for expression of the protein molecules. As described herein any molecules of present invention can be cloned for expression of its corresponding peptide or protein molecule.

(B) Over expression/Purification of Protein molecules of the present invention:

Appropriate clones (as shown in Figures 4-7) expressing all four versions of above-mentioned proteins in *E. coli* were identified by appropriate antibiotic selection as described by the manufacturer (Novagen 441 Charmany Drive Madison, WI, 53719, USA). Expressed protein bands of the predicted sizes were identified by analytical SDS poly acrylamide gel electrophoresis (Laemmli, U.K., *Nature*, 227, 680, 1970) to confirm the expression of desired protein. A gel with the His tagged and Non tagged proteins for all four proteins clearly shows the size differences. Protein obtained from the plasmid pMON63134 (Figure 6) was chosen for assay work described below. Protein extracts were prepared by freeze thawing and sonication as described Cull and McHenry (Cull M and McHenry C.S. *Methods in Enzymology* 182, 147-153, 1990). Pellets were resuspended in an appropriate volume of either 50 mM Tris-HCl pH 7.4 250 mM NaCl and 500 μ M EDTA with 0.05% CHAPS and 1mM ABESF or 50 mM KH_2PO_4 pH-7.2 250 mM NaCl and 250 μ M EDTA with 0.05% CHAPS and 1mM ABESF, buffer. Crude extracts containing His-tagged protein were applied to equilibrated His-Trap (Pharmacia Piscataway, NJ) columns and purified by standard procedures as described by the manufacturer of the columns. Steps included a 10 mM imidazole elution /wash and elution in 250 mM imidazole. As a result the Arabidopsis tagged protein was partially purified (approximately 85-90 % by SDS-PAGE stained with Coomassie Blue). Purified samples were gel filtered into 50 mM KH_2PO_4 pH-7.2 250 mM NaCl and 250 μ M EDTA, 10% glycerol and stored at -80°C until use. Protein concentrations were 1.5 to 1.7 mg /mL, based on the method of Bradford (Bradford M, *Anal Biochem*, 72, 248, 1976 and Bio-Rad Laboratories Procedure bulletin 1123) with BSA standard (Bradford reagent and BSA was from Bio-Rad Laboratories Headquarters 1000 Alfred

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Nobel Drive Hercules, CA 94547). The assays described below used 10 or 15 uL of this solution (15 to 22 µg per reaction).

(C) Catalytic or Enzymatic activity determination of peptide/protein molecules of present invention.

5

Spectrophotometric assays were used for determination of enzyme activities of purified proteins. Assays were based on earlier described methods (Portsteffen et al. *Phytochemistry*, 37, 391, 1994). Consumption or production of NADPH was observed at 340 nm. Assays were performed in a Varian Cary 50 Bio spectrophotometer (Varian Instruments Inc. AA, ICP, UV, Fluorescence Products 679 Springvale RoadMulgrave, Victoria 3170Australia). All assays were run at 30°C for 10 to 30 minutes in 1mL volumes using 100 mM KH₂PO₄ pH 6.5, 250 µM EDTA or 50 mM KH₂PO₄ pH-7.2, 250 mM NaCl, and 250 µM EDTA as reaction buffer. If metals were added they were supplemented to reactions in cuvettes from 100 mM stocks of either ZnCl₂ and or MgCl₂. Substrates included those found in table 8. All assays contained 3 to 15 30µg of protein 200 to 400 µM NADPH or NADP and 100 µM to 10 mM substrate (acetone 60mM). Controls contained no substrate to determine the background rate of consumption and degradation of NADPH. NADPH solutions were freshly made and checked by UV-vis for degradation. Enzyme solutions were supplemented with 1mM MgCl₂ and 1mM Zn Cl₂ which were at 1.5 to 1.7 mg /mL concentrations. The assays used 10 or 15 µL of this solution (ca. 15 to 20 22 µg per reaction). Stock solutions of the substrates (20 to 500mM) were made up in a reaction buffer or as noted. Steroids solutions were made in 100% EtOH. From these stock solutions additions to the assays were made directly. Precipitation of steroids was observed. Thus, the soluble concentration of steroids in reaction mixture is not accurately known in each case. However, good estimate of substrate (sterol) concentration would be 100 to 200 µM for a 25 saturated solution under these conditions. There was a clear difference in rate observed by changing concentration from 26 µM to 260 µM indicating that saturation may be between the two points both for enzyme activity and substrate solubility. At these concentrations precipitation did not have an effect on the absorbance changes at 340 nm as evidenced by controls containing substrate and NADPH. In addition to cell extract control, a substrate control 30 was also employed because solutions of some substrates tested can have a lower pH at

described assay conditions which may affect rates of NADPH degradation and enzyme activity under assay conditions. Results of the substrates tested can be found in Table 10 below.

These results (Table 8 to 10) show that the *A. thaliana* protein molecule SEQ ID NO: 4 has significant steroid reductase activity. Combined with other biological data these results suggest that SEQ ID NO: 4 is likely a steroid reductase with true substrates that are most likely brassinosteroids. These results do not preclude the possibility that another compound class might act as substrate but results suggest that the substrate is a ketone moiety potentially in a steroid ring system.

Table 8: Zero Order Result Table using acetone as substrate (60mM) shows evidence of reductase activity. Turnover (consumption of NADPH) is observed in the reductive direction. Slopes (for acetone, lines 1 and 2) indicate the difference from control is 10 fold. Controls (lines 3 and 4) were identical except for the omission of acetone.

Sample	Start (min)	Stop (min)	Slope (Abs/min)	A Start	A Stop	S.D.
Acetone	0.000	10.00	-0.0118	0.8869	0.7640	0.0016
Acetone	0.000	10.00	-0.0111	0.8655	0.7502	0.0025
Control	0.000	10.00	-0.0006	0.8609	0.8547	0.0006
Control	0.000	10.00	-0.0010	0.8773	0.8683	0.0011

Start=start time, Stop=stop time, Slope is change in absorbance over time, A start is starting absorbance, A stop is ending absorbance.


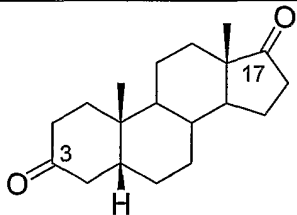
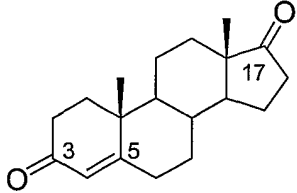
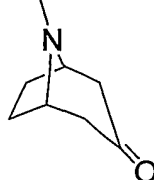
Table 9: Zero Order Result Table. This table shows elected data for 5 α -androstane-3,17-dione in a reductase assay. This example shows the results for three reactions with acetone, three reactions with 5 α -androstane-3,17-dione and two controls. Table 8 shows the initial rates as estimated by linear fit between 0 and 5 minutes. The results for 5 α -androstane-3,17-dione shown here are the best of any substrate at the same concentration (approx 200 μ M). 4-androstene-3,17-dione is the next best substrate with a rate on the order of 2.5 to 4 times slower than this under the same conditions (not shown). Acetone results at similar concentrations (Table 7 shows results at high acetone concentration) are on the order of 10 to 20 times slower and nearly at background levels. Similar results have been obtained with multiple batches of enzyme and substrate.

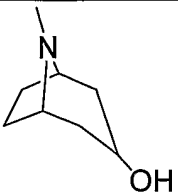
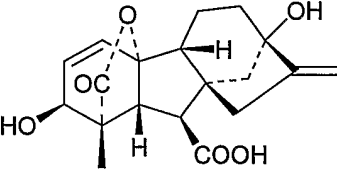
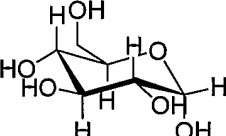

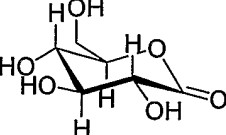
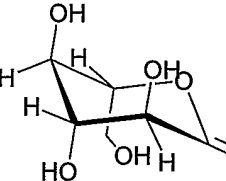
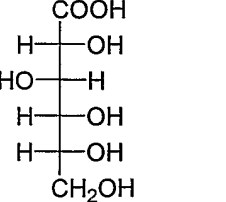
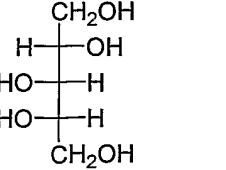
- 63 -

Sample	Start (min)	Stop (min)	Slope (Abs/min)	A Start	A Stop	S.D.
5-alpha	0.000	5.000	-0.0632	1.4399	1.1322	0.0018
5-alpha	0.000	5.000	-0.0611	1.4105	1.1092	0.0022
5-alpha	0.000	5.000	-0.0652	1.4710	1.1652	0.0030
acetone	0.000	5.000	-0.0049	1.5050	1.4793	0.0018
acetone	0.000	5.000	-0.0047	1.3902	1.3961	0.0012
acetone	0.000	5.000	-0.0041	1.4584	1.4352	0.0014
Control	0.000	5.000	-0.0030	1.5256	1.5118	0.0013
Control	0.000	5.000	-0.0035	1.4898	1.4743	0.0018

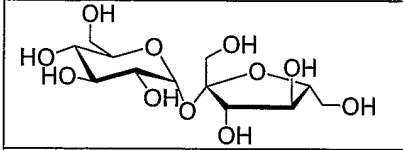
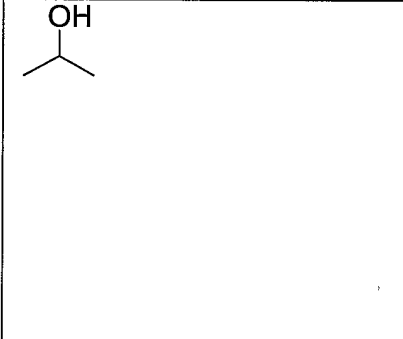
Start=start time, Stop=stop time, Slope is change in absorbance over time, A start is starting absorbance, A stop is ending absorbance.

Table 10: Substrates tested.

Substrate	Name	Activity	Comments /references
	acetone	Yes 65mM Much lower at lower concentrations	
	5 α -androstane- 3,17-dione	YES 100 μ M	17 β activity Chemico- biological interactions 130-132 (2001)783- 803
	4-androstene- 3,17-dione	YES 100 μ M	3 α activity Chemico- biological interactions 130-132 (2001)783- 803 5 α activity (PNAS , 94, 3354- 3359, 1997)
	Tropinone	No retested with new enzyme 1 and 10 mM	(Photochemistry, 37, 391-400, 1994)

	Tropine	No retested with new enzyme 1 and 10 mM	(Photochemistry, 37, 391-400, 1994)
	Gibberellic Acid (GA ₃)	No 1 and 10 mM	
Additional substrates tested	Name	Activity	Chosen based on informatics results 2nd tier possibilities
	D-Glucose	No 25mM	
	Glucose phosphate	6 No 25mM	
	Glucono 1,5 lactone	1,5 No 25mM	Stereoisomer of δ-gluconolactone. Easily hydrolyzed in H ₂ O.
	δ-gluconolactone	No 25mM	Stereoisomer of Glucono 1,5 lactone. pH of solution is very low i.e. below 3.
	Potassium gluconate	d- No 25mM	Potassium salt used
	Arabinitol	No 25mM	

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	Sucrose	No 25mM	
	isopropanol	Yes 68mM	At high concentrations it is a substrate for the reverse reaction but at same concentrations is a poorer substrate than acetone indicating potential preference for the reductase activity under these conditions.

EXAMPLE 11

In the following example a large number of DNA sequences were searched using BlastA (Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang,
5 Webb Miller, and David J. Lipman, *Nucleic Acids Res.* (25)3389-3402 (1997)) to find various homologs, paralogs or orthologs of the gene of the present invention. These sequences are determined from cDNA libraries prepared from a variety plant species and tissues.

For construction of cDNA libraries, tissue is harvested and immediately frozen in liquid nitrogen and stored at -80°C until total RNA extraction. Total RNA is purified from tissue using
10 Trizol reagent from Life Technologies (Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A.), as recommended by the manufacturer. Poly A⁺ RNA (mRNA) is purified using magnetic oligo dT beads essentially as recommended by the manufacturer (Dynabeads, Dynal Corporation, Lake Success, New York U.S.A.).

Construction of plant cDNA libraries is well known in the art and a number of cloning
15 strategies exist. A number of cDNA library construction kits are commercially available. The SuperscriptTM Plasmid System for cDNA synthesis and Plasmid Cloning (Gibco BRL, Life Technologies, Gaithersburg, Maryland U.S.A.) is used, following the conditions suggested by the manufacturer.

The cDNA libraries are plated on LB agar containing the appropriate antibiotics for
20 selection and incubated at 37° for sufficient time to allow the growth of individual colonies. Single selective media colonies are individually placed in each well of a 96-well microtiter plates

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containing LB liquid including the selective antibiotics. The plates are incubated overnight at approximately 37°C with gentle shaking to promote growth of the cultures.

The plasmid DNA is isolated from each clone using Qiaprep plasmid isolation kits, using the conditions recommended by the manufacturer (Qiagen Inc., Santa Clara, California U.S.A.).

5 The template plasmid DNA clones are used for subsequent sequencing. For sequencing the cDNA libraries, a commercially available sequencing kit, such as the ABI PRISM dRhodamine Terminator Cycle Sequencing Ready Reaction Kit with AmpliTaq® DNA Polymerase, FS, is used under the conditions recommended by the manufacturer (PE Applied Biosystems, Foster City, CA). The cDNAs of the present invention are generated by sequencing
10 initiated from the 5' end or 3' end of each cDNA clone. Entire inserts or only part of the inserts (ESTs or expressed sequenced tags) are sequenced.

A number of DNA sequencing techniques are known in the art, including fluorescence-based sequencing methodologies. These methods have the detection, automation and instrumentation capability necessary for the analysis of large volumes of sequence data.
15 Currently, the 377 and 3700 DNA Sequencer (Perkin-Elmer Corp., Applied Biosystems Div., Foster City, CA) allows the most rapid electrophoresis and data collection. With these types of automated systems, fluorescent dye-labeled sequence reaction products are detected and data entered directly into the computer, producing a chromatogram that is subsequently viewed, stored, and analyzed using the corresponding software programs. These methods are known to
20 those of skill in the art and have been described and reviewed (Birren *et al.*, *Genome Analysis: Analyzing DNA*, 1, Cold Spring Harbor, New York, the entirety of which is herein incorporated by reference).

The generated ESTs (including any full-length cDNA inserts or complete coding sequences) are combined with ESTs and full-length cDNA sequences in public databases such as
25 GenBank. Duplicate sequences are removed, and duplicate sequence identification numbers are replaced. The combined dataset is then clustered and assembled using Pangea Systems tool identified as CAT v.3.2. First, the EST sequences are screened and filtered, *e.g.* high frequency words are masked to prevent spurious clustering; sequence common to known contaminants such as cloning bacteria are masked; high frequency repeated sequences and simple sequences are
30 masked; unmasked sequences of less than 100 bp are eliminated. The screened and filtered ESTs are combined and subjected to a word-based clustering algorithm which calculates sequence pair

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distances based on word frequencies and uses a single linkage method to group like sequences into clusters of more than one sequence, as appropriate. Clustered sequences are assembled individually using an iterative method based on PHRAP/CRAW/MAP providing one or more self-consistent consensus sequences and inconsistent singleton sequences. The assembled clustered sequence files are checked for completeness and parsed to create data representing each consensus contiguous sequence (contig), the initial EST sequences, and the relative position of each EST in a respective contig. The sequence of the 5' most clone is identified from each contig. The initial sequences that are not included in a contig are separated out.

Above described databases with nucleotide and peptide sequences were queried with sequences of the present invention to get following homologues, orthologs or paralogs as shown in following table. The BLAST 2.2.1 software (Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. Lipman, Nucleic Acids Res. (25)3389-3402 (1997), with BLOSUM62 matrix and "no Filter" options, was used in the queries. As when necessary, frame-shifts in the DNA sequences of the homologues were detected by aligning the DNA sequence of the homologue in question to the protein sequence of present invention, using the "frame+_n2p" program with default parameters in the GenCore software package (Compugen Inc., 1998). Such frame-shifts were conceptually corrected to yield open reading frames. The "translate" program with default parameters in the same package was used to translate open reading frames to corresponding peptide sequences based on standard genetic codes.

SEQ ID NO: 1 exhibits 52.857% percent identity with its closest known functional gene, identified using the BLAST 2.2.1 software with BLOSUM62 matrix and "no Filter" options (Genbank accession number gi|1717752|sp|P50162|TRN1_DATST). SEQ ID NO: 1 exhibits identity of 111 amino acid residues over its entire length of 210 amino acids with tropionone reductase of *Datura stramonium* (Genbank accession number gi|1717752|sp|P50162|TRN1_DATST). Due to this relationship, it is possible that enzymes with similar activities can also function in present invention, also DNA molecules encoding proteins that are closely related to tropionone reductase are an aspect of the present invention.

Table 11.

Seq num	Gene Bank Identity	Species
1	none	<i>Arabidopsis thaliana</i>

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2	none	<i>Arabidopsis thaliana</i>
3	none	<i>Arabidopsis thaliana</i>
4	none	<i>Arabidopsis thaliana</i>
5	gi 3980405 gb AAC95208.1 _cds	<i>Arabidopsis thaliana</i>
6	gi 3980405 gb AAC95208.1	<i>Arabidopsis thaliana</i>
7	gi 3980415 gb AAC95218.1 _cds	<i>Arabidopsis thaliana</i>
8	gi 3980415 gb AAC95218.1	<i>Arabidopsis thaliana</i>
9	gi 3980416 gb AAC95219.1 _cds	<i>Arabidopsis thaliana</i>
10	gi 3980416 gb AAC95219.1	<i>Arabidopsis thaliana</i>
11	gi 3980398 gb AAC95201.1 _cds	<i>Arabidopsis thaliana</i>
12	gi 3980398 gb AAC95201.1	<i>Arabidopsis thaliana</i>
13	gi 13605590 gb AF361621.1 AF361621	<i>Arabidopsis thaliana</i>
14	gi 13605591 gb AAK32789.1 AF361621_1	<i>Arabidopsis thaliana</i>
15	gi 8978342 dbj BAA98195.1 _cds	<i>Arabidopsis thaliana</i>
16	gi 8978342 dbj BAA98195.1	<i>Arabidopsis thaliana</i>
17	gi 2880044 gb AAC02738.1 _cds	<i>Arabidopsis thaliana</i>
18	gi 2880044 gb AAC02738.1	<i>Arabidopsis thaliana</i>
19	gi 3980401 gb AAC95204.1 _cds	<i>Arabidopsis thaliana</i>
20	gi 3980401 gb AAC95204.1	<i>Arabidopsis thaliana</i>
21	gi 3980403 gb AAC95206.1 _cds	<i>Arabidopsis thaliana</i>
22	gi 3980403 gb AAC95206.1	<i>Arabidopsis thaliana</i>
23	gi 3980406 gb AAC95209.1 _cds	<i>Arabidopsis thaliana</i>
24	gi 3980406 gb AAC95209.1	<i>Arabidopsis thaliana</i>
25	ARATH-14MAR01-CLUSTER10347_2	<i>Arabidopsis thaliana</i>
26	ARATH-14MAR01-CLUSTER10347_2_prot	<i>Arabidopsis thaliana</i>
27	ARATH-14MAR01-CLUSTER10347_3	<i>Arabidopsis thaliana</i>
28	ARATH-14MAR01-CLUSTER10347_3_prot	<i>Arabidopsis thaliana</i>
29	ARATH-14MAR01-CLUSTER10347_5	<i>Arabidopsis thaliana</i>
30	ARATH-14MAR01-CLUSTER10347_5_prot	<i>Arabidopsis thaliana</i>
31	ARATH-14MAR01-CLUSTER136303_1	<i>Arabidopsis thaliana</i>
32	ARATH-14MAR01-CLUSTER136303_1_prot	<i>Arabidopsis thaliana</i>
33	None	<i>Glycine max</i>
34	None	<i>Glycine max</i>
35	None	<i>Glycine max</i>
36	None	<i>Glycine max</i>
37	None	<i>Glycine max</i>
38	None	<i>Glycine max</i>
39	None	<i>Glycine max</i>
40	None	<i>Glycine max</i>

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41	None	<i>Glycine max</i>
42	None	<i>Glycine max</i>
43	None	<i>Zea mays</i>
44	None	<i>Zea mays</i>
45	None	<i>Zea mays</i>
46	None	<i>Zea mays</i>
47	None	<i>Oryza sativa</i>
48	None	<i>Oryza sativa</i>
49	gi 17231948 ref NP_488496.1 _cds	<i>Nostoc sp. PCC7120</i>
50	gi 17231948 ref NP_488496.1	<i>Nostoc sp. PCC7120</i>
51	None	<i>Xanthomonas campestris</i>
52	None	<i>Xanthomonas campestris</i>
53	XYL200910	<i>Xylella fastidiosa</i>
54	XYL200910_prot	<i>Xylella fastidiosa</i>

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CLAIMS:

1. A method of increasing seed and organ size of a plant comprising the steps of:
 - a) transforming said plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, wherein said DNA molecule is selected from the group consisting of: SEQ ID NO: 1, SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7, 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: 66, SEQ ID NO: 68, and SEQ ID NO: 70 which is operably linked to a 3' termination region; and
 - b) selecting a desired plant from a population of transformed plants containing said DNA construct; wherein said desired plant exhibits increased seed and organ size compared to a plant of a same plant species not transformed to contain said DNA construct.
2. The method of claim 1, wherein said promoter is a caulimovirus promoter.
3. The method of claim 1, wherein said promoter comprises a heterologous plant constitutive promoter.
4. The method of claim 1, wherein said promoter is a tissue specific promoter or an organ enhanced promoter.
5. A method of increasing seed and organ size of a plant comprising the steps of:
 - a) transforming said plant with a DNA construct comprising a promoter that functions in plants, operably linked to a DNA molecule that encodes a protein, wherein said protein comprises at least an N-terminal 50% portion of a polypeptide selected from the group consisting of: SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8,

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5 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: 67, SEQ ID NO: 69,
and SEQ ID NO: 71, wherein said DNA construct is operably linked to a 3' termination
region; and

b) selecting a plant from a population of transformed plants containing said DNA
construct; wherein said plant exhibits increased seed and organ size compared to a plant
10 of a same plant species not transformed to contain said DNA construct.

6. The method of claim 5, wherein said plant species is selected from the group consisting
of: *Acacia*, alfalfa, aneth, apple, apricot, artichoke, arugula, asparagus, avocado, banana, barley,
beans, beet, blackberry, blueberry, broccoli, brussels sprouts, cabbage, canola, cantaloupe, carrot,
15 cassava, cauliflower, celery, cherry, cilantro, citrus, clementines, coffee, corn, cotton, cucumber,
Douglas fir, eggplant, endive, escarole, eucalyptus, fennel, figs, gourd, grape, grapefruit, honey
dew, jicama, kiwifruit, lettuce, leeks, lemon, lime, loblolly pine, mango, melon, millet,
mushroom, nut, oat, okra, onion, orange, an ornamental plant, papaya, parsley, pea, peach,
peanut, pear, pepper, persimmon, pine, pineapple, plantain, plum, pomegranate, poplar, potato,
20 pumpkin, quince, radiata pine, radicchio, radish, raspberry, rice, rye, sorghum, southern pine,
soybean, spinach, squash, strawberry, sugarbeet, sugarcane, sunflower, sweet potato, sweetgum,
tangerine, tea, tobacco, tomato, turf, a vine, watermelon, wheat, yams, and zucchini.

7. A transgenic plant with increased seed and organ size compared to a non transformed
25 plant of a same plant species, said transgenic plant comprising a DNA construct, wherein said
DNA construct encodes a protein comprising at least an N-terminal 50% portion selected from
the group consisting of: SEQ ID NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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
30 NO: 32, SEQ ID NO: 34, SEQ ID NO: 36, SEQ ID NO: 38, SEQ ID NO: 40, SEQ ID NO: 42,

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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: 67, SEQ ID NO: 69, and SEQ ID NO: 71.

8. The transgenic plant of claim 7, wherein said DNA construct encodes a protein
5 comprising at least an N-terminal 60% portion selected from the group consisting of: SEQ ID
NO: 2, SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, 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:
10 46, SEQ ID NO: 48, SEQ ID NO: 50, SEQ ID NO: 52, SEQ ID NO: 54, SEQ ID NO: 67, SEQ
ID NO: 69, and SEQ ID NO: 71.

9. The transgenic plant of claim 7, wherein said plant species is selected from the group
consisting of: *Acacia*, alfalfa, aneth, apple, apricot, artichoke, arugula, asparagus, avocado,
15 banana, barley, beans, beet, blackberry, blueberry, broccoli, brussels sprouts, cabbage, canola,
cantaloupe, carrot, cassava, cauliflower, celery, cherry, cilantro, citrus, clementines, coffee, corn,
cotton, cucumber, Douglas fir, eggplant, endive, escarole, eucalyptus, fennel, figs, gourd, grape,
grapefruit, honey dew, jicama, kiwifruit, lettuce, leeks, lemon, lime, loblolly pine, mango,
melon, millet, mushroom, nut, oat, okra, onion, orange, an ornamental plant, papaya, parsley,
20 pea, peach, peanut, pear, pepper, persimmon, pine, pineapple, plantain, plum, pomegranate,
poplar, potato, pumpkin, quince, radiata pine, radicchio, radish, raspberry, rice, rye, sorghum,
southern pine, soybean, spinach, squash, strawberry, sugarbeet, sugarcane, sunflower, sweet
potato, sweetgum, tangerine, tea, tobacco, tomato, turf, a vine, watermelon, wheat, yams, and
zucchini.

25

10. A progeny plant produced from said transgenic plant of claim 7.

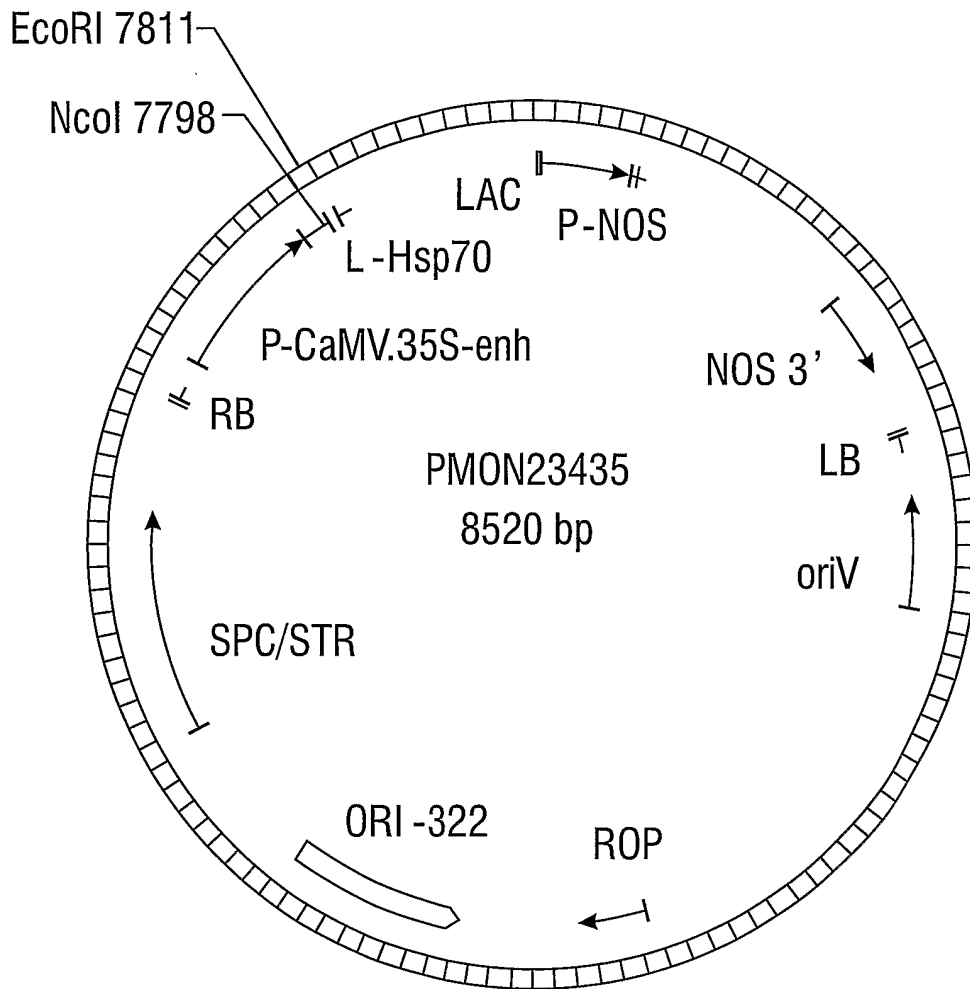


FIG. 1

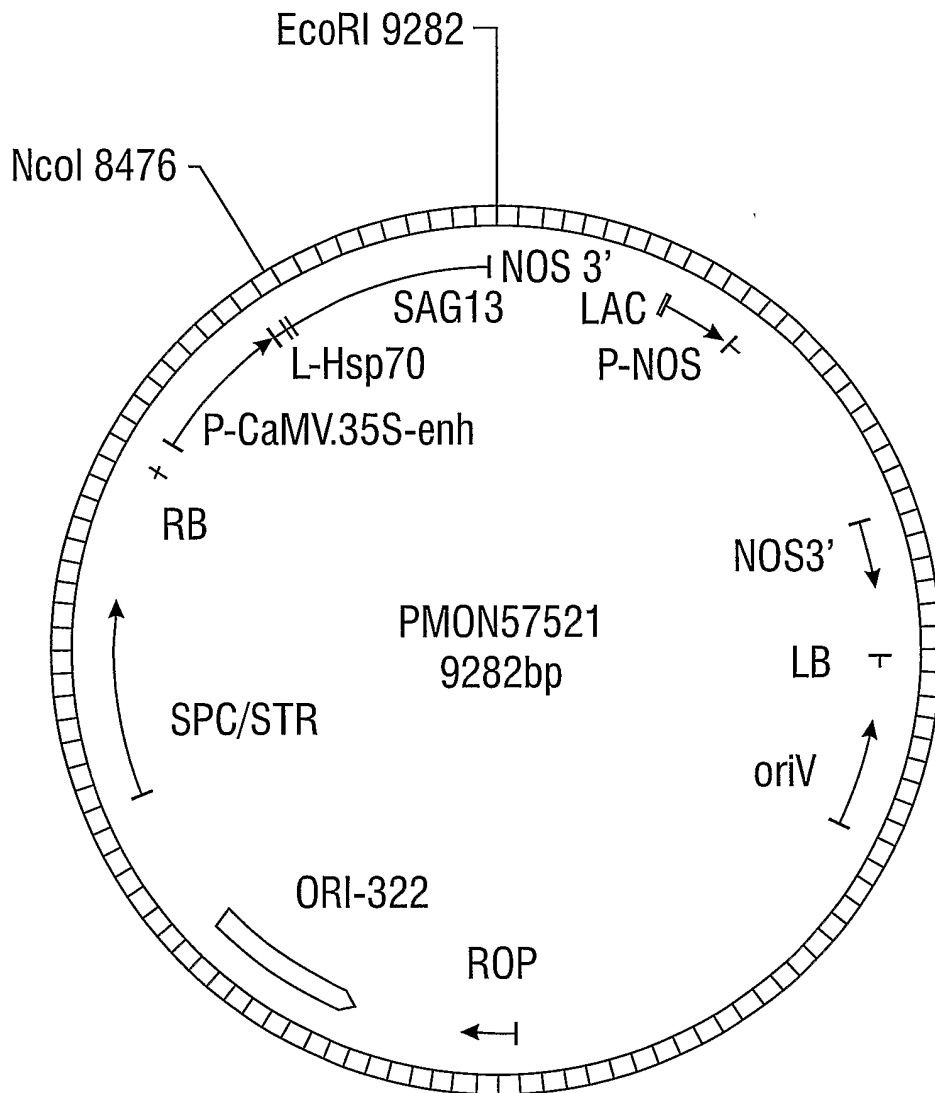


FIG. 2

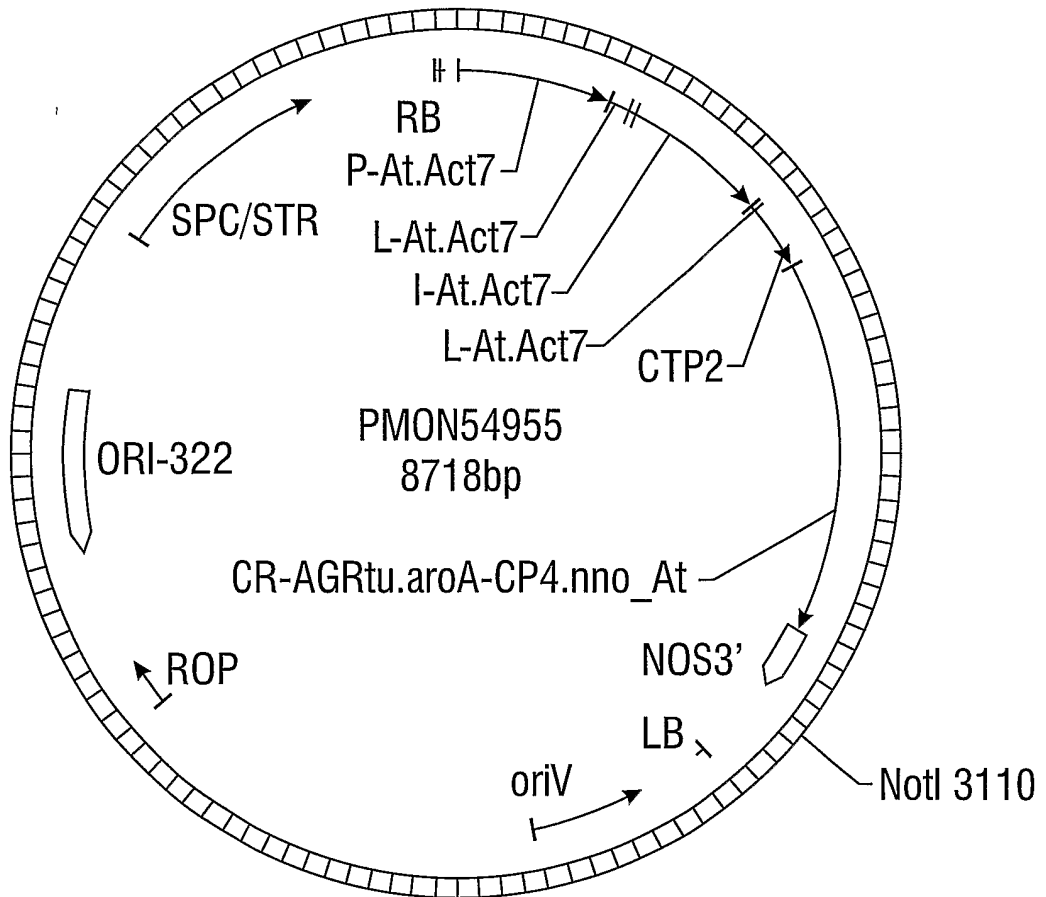


FIG. 3

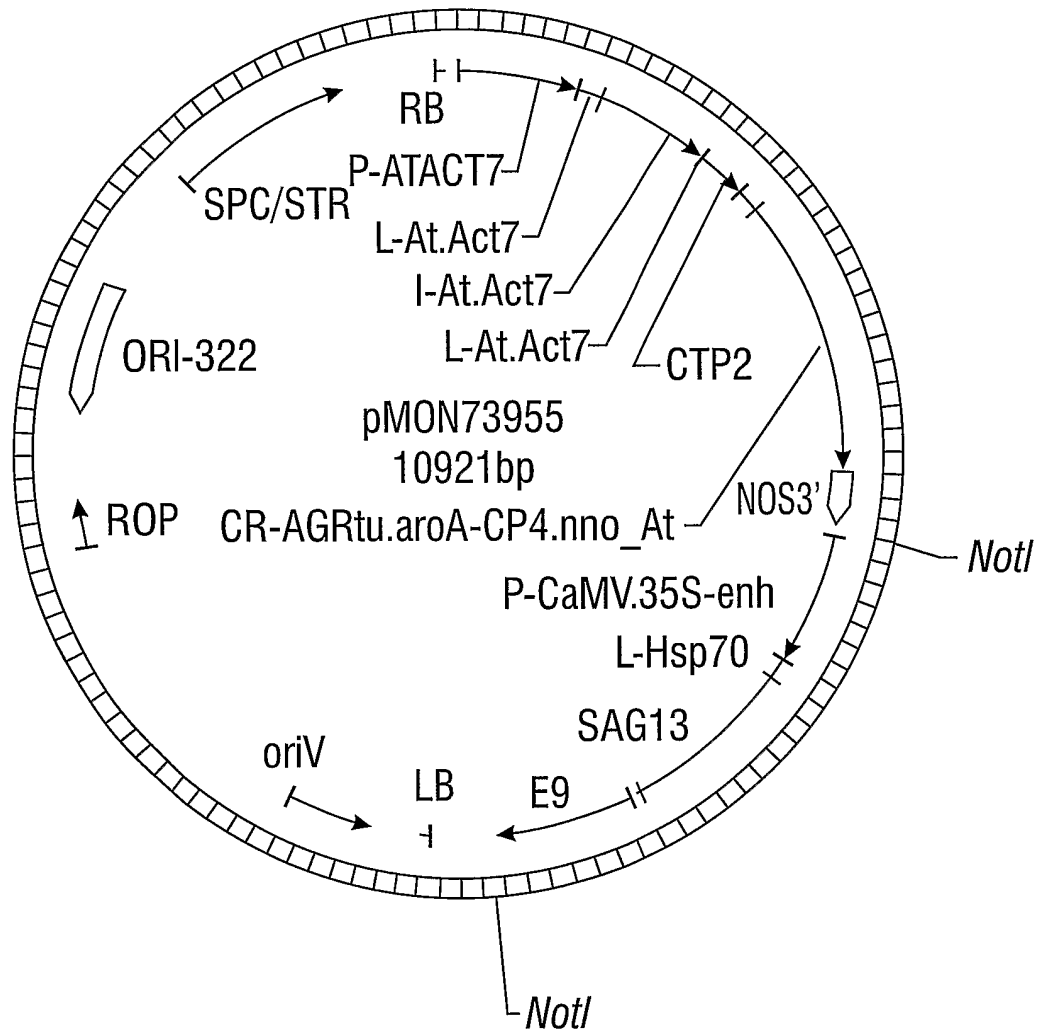


FIG. 4

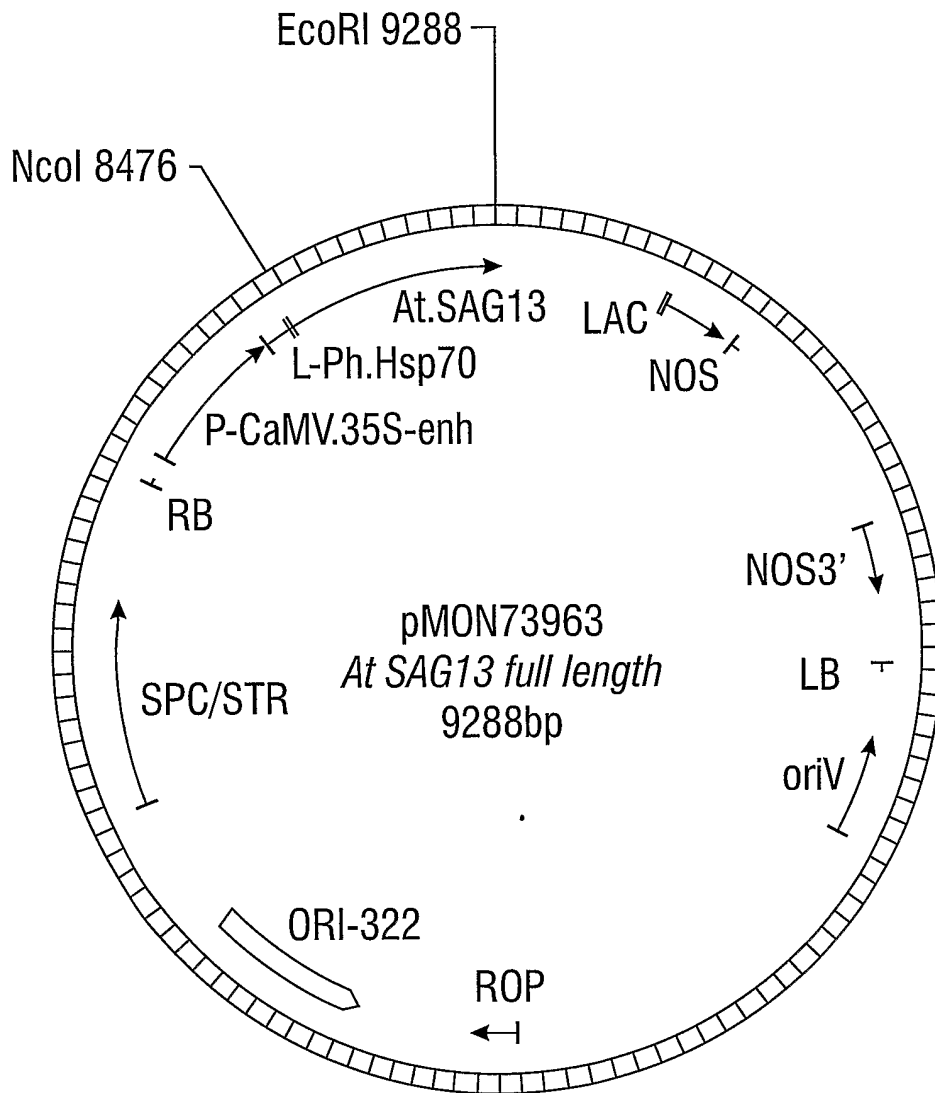


FIG. 5

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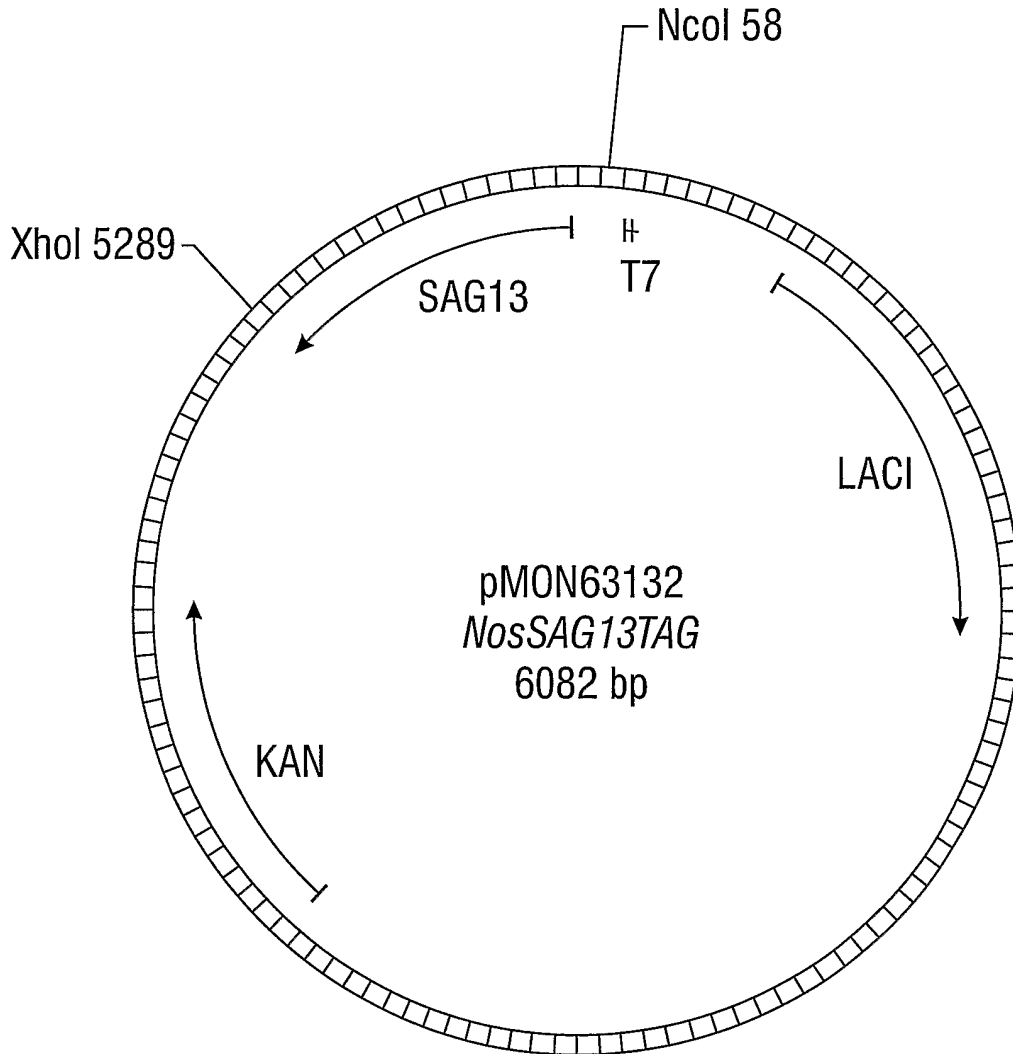


FIG. 6

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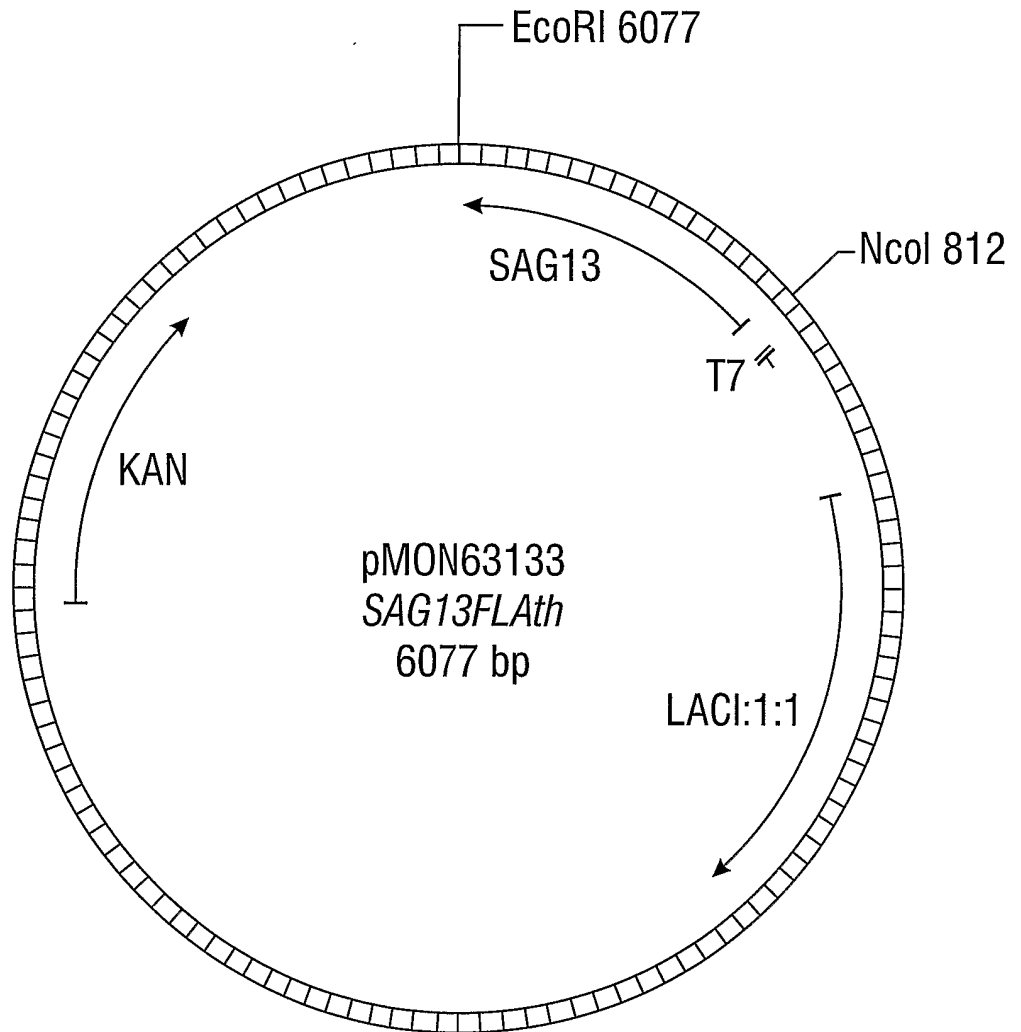


FIG. 7

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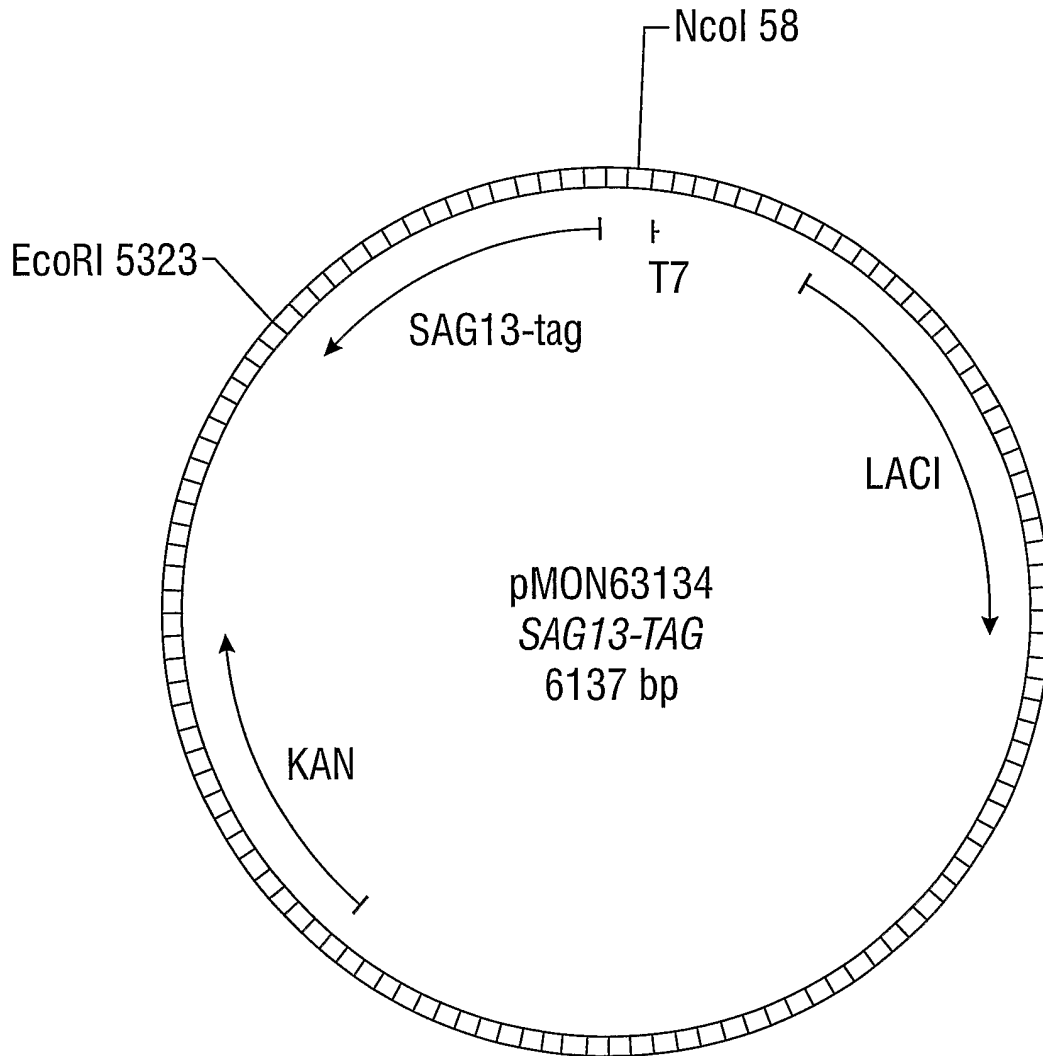


FIG. 8

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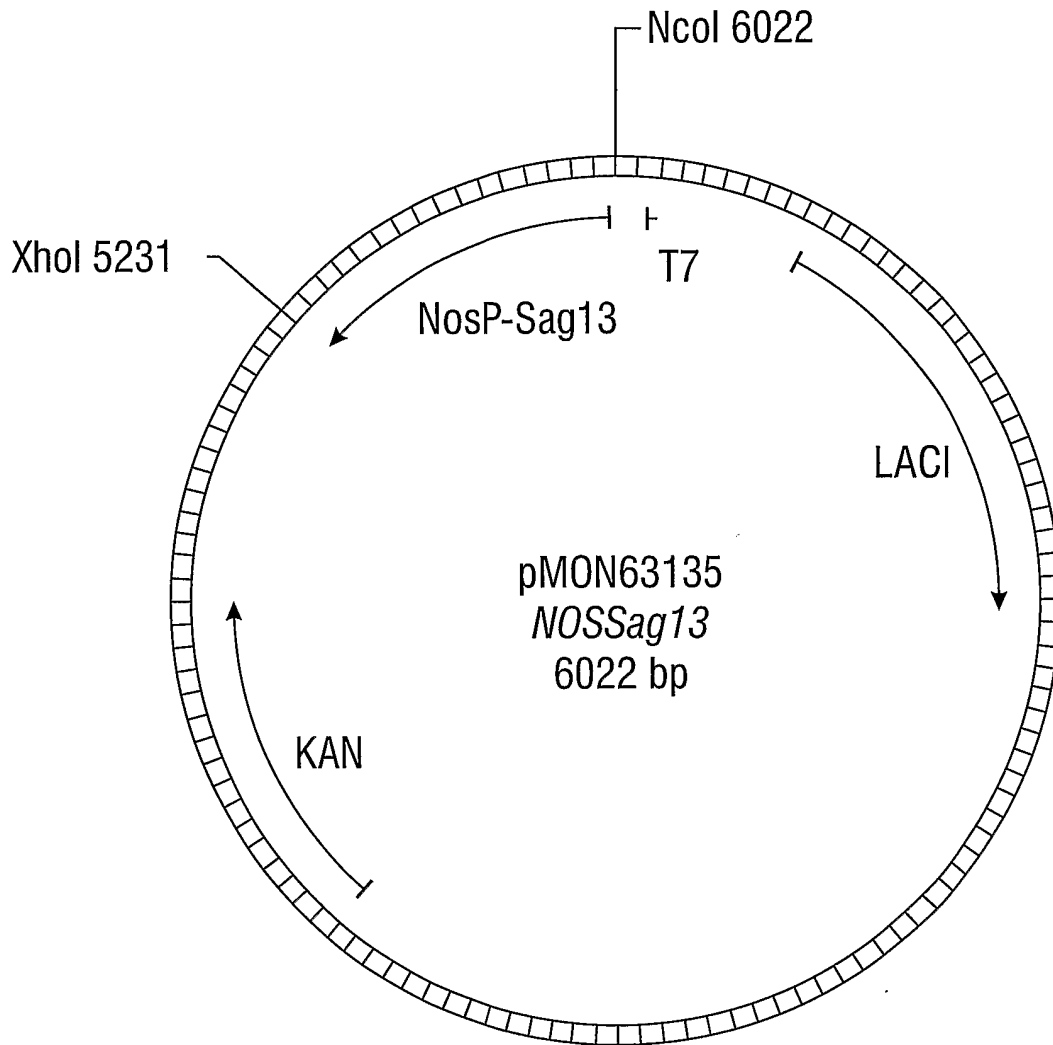


FIG. 9

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Primers	Sequence ID No.	Structural Comments
Arabidopsis Primers		
GTAAGATATCATATGGCAAAGG AAGGG	SEQ ID No: 57	BgIII and NdeI site N-terminal histag
GCGCGAATTCTCGAGTCATAGTC TTGAAGGAAAAAC	SEQ ID No: 58	Has EcoRI and XhoI
GGCTACAAATCTCGAGTCAGC	SEQ ID No: 59	
GCTGACTCGAGATTGTAGCC	SEQ ID No: 60	
Nostoc Primers		
GTAAGATATCATATGAGTGATGC CTACGGTG	SEQ ID No: 61	NdeI site N-terminal histag
GTAAGATCTGCCACCATGGGTGA TGCCACGGTG	SEQ ID No: 62	Modified 2nd codon for NcoI site Ser to Gly)
GAGAAAGCTTCTCGAGTTAAAAA CCGAAAGCC	SEQ ID No: 63	(HindIII/XhoI)
CCTCTACTGCAAAATAGGG	SEQ ID No: 64	
CCCTATTTTCAGTAGAGG	SEQ ID No: 65	

FIG. 10

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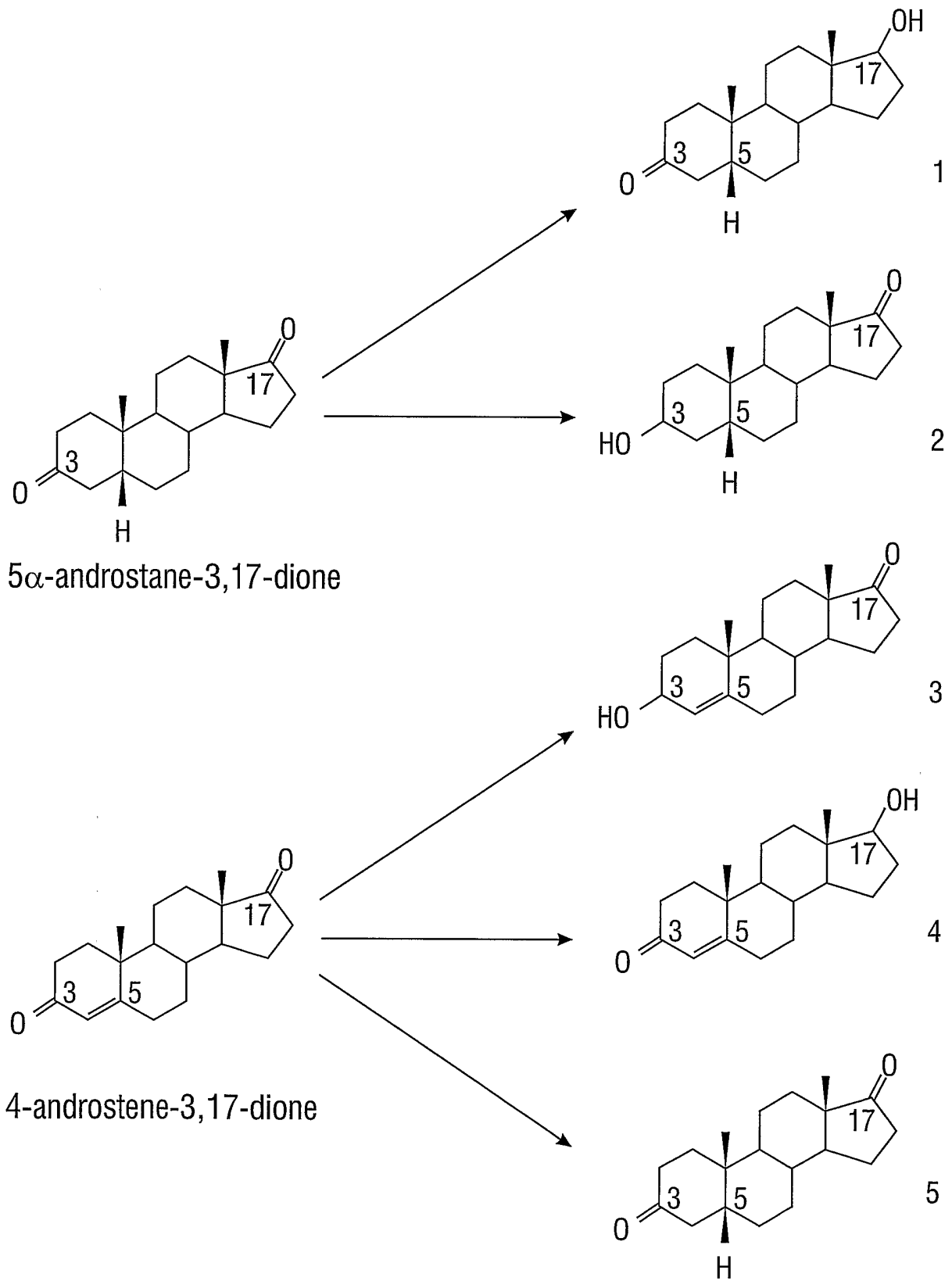


FIG. 11

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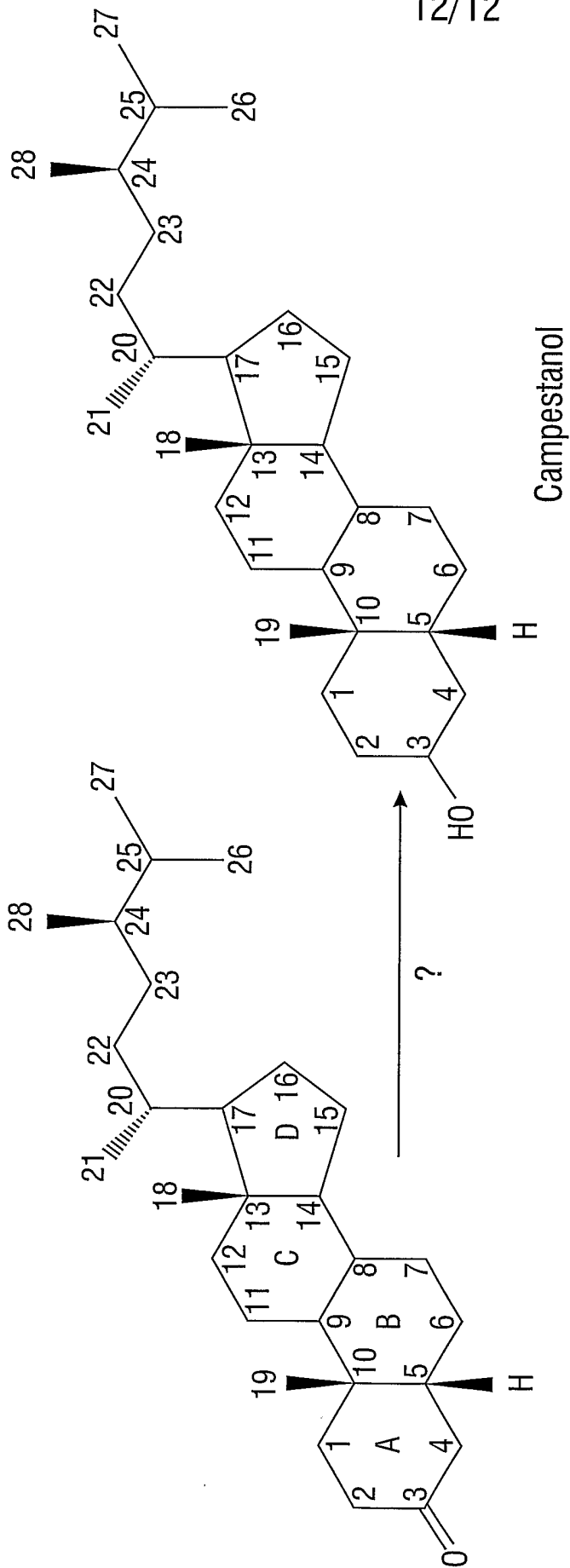


FIG. 12

		20						25								30
Val	Val	Glu	Glu	Leu	Ala	Met	Leu	Gly	Ala	Lys	Val	His	Thr	Cys	Ala	
		35					40					45				
Arg	Asp	Glu	Thr	Gln	Leu	Gln	Glu	Arg	Leu	Arg	Glu	Trp	Gln	Ala	Lys	
	50					55					60					
Gly	Phe	Gln	Val	Thr	Thr	Ser	Val	Cys	Asp	Val	Ser	Ser	Arg	Asp	Gln	
65					70					75					80	
Arg	Val	Lys	Leu	Met	Glu	Thr	Val	Ser	Ser	Leu	Tyr	Gln	Gly	Lys	Leu	
				85					90					95		
Asn	Ile	Leu	Val	Asn	Asn	Val	Gly	Thr	Ser	Ile	Phe	Lys	Pro	Thr	Thr	
			100					105						110		
Glu	Tyr	Thr	Ala	Glu	Asp	Phe	Ser	Phe	Val	Met	Ala	Thr	Asn	Leu	Glu	
		115					120						125			
Ser	Ala	Phe	His	Leu	Ser	Gln	Leu	Ala	His	Pro	Leu	Leu	Lys	Ala	Ser	
	130					135					140					
Gly	Ser	Gly	Ser	Ile	Val	Leu	Ile	Ser	Ser	Ala	Ala	Gly	Val	Val	His	
145					150					155					160	
Val	Asn	Val	Gly	Ser	Ile	Tyr	Gly	Ala	Thr	Lys	Gly	Ala	Met	Asn	Gln	
				165					170						175	
Leu	Ala	Arg	Asn	Leu	Ala	Cys	Glu	Trp	Ala	Ser	Asp	Asn	Ile	Arg	Thr	
			180					185					190			
Asn	Ser	Val	Cys	Pro	Trp	Tyr	Ile	Thr	Thr	Pro	Leu	Ser	Asn	Asp	Phe	
		195					200					205				
Phe	Arg															
	210															

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 <213> Arabidopsis thaliana

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 tgggagcaaaa agtccacacg tgtgccagag acgaaactca gcttcaagaa cgcttacgtg 180

aatggcaagc aaaaggggtt caggtcacca cttctgtctg cgacgtctct tctcgtgacc 240
 aacgagtgaa actcatggaa actgtttcct ctctctacca aggaaaactc aacatcctcg 300
 tcaacaatgt gggaacgtca atattcaagc cgaccacaga gtatacagca gaagatttct 360
 cttttgtaat ggctacaaat ctcgagtcag ctttccatct ctcacagctt gccacccat 420
 tgttaaaagc ctctggctca gggagcatcg tgctcatatc ctctgctgct ggagtcgtgc 480
 atgtcaatgt tggatccatc tatggagcaa ccaaaggagc catgaatcag ctggctagaa 540
 acttagcttg cgaatgggag agcgacaaca taaggacgaa ctctgtttgt ccttgggtata 600
 tcacaactcc ttttaagtaac gattttttcg atgaagagtt taagaaagaa gcggtgcgta 660
 cgacaccaat ggggcgtgtg gagaagcaaa tgaagtctca ccgcttgtgg catttctttg 720
 tcttccttca gcttcttata ttactggta gaccatttgc gttgacggcg gtgccactgt 780
 taacgggtttt tccttcaaga ctatgaattc 810

<210> 4
 <211> 268
 <212> PRT
 <213> Arabidopsis thaliana

<400> 4

Met Ala Lys Glu Gly Gly Leu Gly Glu Asn Ser Arg Trp Ser Leu Gly
 1 5 10 15

Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Glu Ala
 20 25 30

Val Val Glu Glu Leu Ala Met Leu Gly Ala Lys Val His Thr Cys Ala
 35 40 45

Arg Asp Glu Thr Gln Leu Gln Glu Arg Leu Arg Glu Trp Gln Ala Lys
 50 55 60

Gly Phe Gln Val Thr Thr Ser Val Cys Asp Val Ser Ser Arg Asp Gln
 65 70 75 80

Arg Val Lys Leu Met Glu Thr Val Ser Ser Leu Tyr Gln Gly Lys Leu
 85 90 95

Asn Ile Leu Val Asn Asn Val Gly Thr Ser Ile Phe Lys Pro Thr Thr
 100 105 110

Glu Tyr Thr Ala Glu Asp Phe Ser Phe Val Met Ala Thr Asn Leu Glu
 115 120 125

Ser Ala Phe His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala Ser

130		135		140											
Gly	Ser	Gly	Ser	Ile	Val	Leu	Ile	Ser	Ser	Ala	Ala	Gly	Val	Val	His
145					150					155					160
Val	Asn	Val	Gly	Ser	Ile	Tyr	Gly	Ala	Thr	Lys	Gly	Ala	Met	Asn	Gln
				165					170					175	
Leu	Ala	Arg	Asn	Leu	Ala	Cys	Glu	Trp	Ala	Ser	Asp	Asn	Ile	Arg	Thr
			180					185					190		
Asn	Ser	Val	Cys	Pro	Trp	Tyr	Ile	Thr	Thr	Pro	Leu	Ser	Asn	Asp	Phe
		195					200					205			
Phe	Asp	Glu	Glu	Phe	Lys	Lys	Glu	Ala	Val	Arg	Thr	Thr	Pro	Met	Gly
	210					215					220				
Arg	Val	Gly	Glu	Ala	Asn	Glu	Val	Ser	Pro	Leu	Val	Ala	Phe	Leu	Cys
225					230					235					240
Leu	Pro	Ser	Ala	Ser	Tyr	Ile	Thr	Gly	Gln	Thr	Ile	Cys	Val	Asp	Gly
				245					250					255	
Gly	Ala	Thr	Val	Asn	Gly	Phe	Ser	Phe	Lys	Thr	Met				
			260					265							

<210> 5
 <211> 789
 <212> DNA
 <213> Arabidopsis thaliana

<400> 5
 atggataaaa gatggagtct tcaaggtatg aatgctcttg tgaccggtgg cactaaaggc 60
 atcggggaag ctgttgtgga ggaactgtca attttgggag caagagtcca cacatgtgct 120
 agagacgaga ctcagcttca agaacgttta cgtgaatggc aagaaaaagg gtttcaggtc 180
 accacttcta tctgcgacgt ttctttgcgt gagcaacgag agaaactcat ggaaaccggt 240
 tcctctctct tccaaggaaa actcaacatc ctcgtgaaca atgtgggaac gttaatgctc 300
 aagccgacca cagagtatac agcagaagaa ttctcgtttc ttatggctac aaatctcgac 360
 tcagctttcc atatttcaca gcttgccgac cctttgttga aagcctctgg ctcagggagc 420
 attgtgctca tgtcctctat tgccggagtc gtgcatgtcg gtggttgatc catctatgga 480
 gcaacaaaag gagccatgaa tcagttggct agaaacttag cttgcgaatg ggcaagcgac 540
 aacataagga ctaacgctat ttgtccatgg ttaatcacia ctcctttgat tagtgatctt 600
 ctcagtggtg aggagatgaa aaaagaagca gaggaagga cacccatggg gcgtgtggga 660

gaggcaaatg aagtctcacc gcttgtggca tttctttgtc ttctgcagc ttcttatatt 720
 actgggtcaag tcatttgtgt tgatggaggt ctcaactgta atggcttctc ctatcagcca 780
 catgcttga 789

<210> 6
 <211> 262
 <212> PRT
 <213> Arabidopsis thaliana

<400> 6

Met Asp Lys Arg Trp Ser Leu Gln Gly Met Asn Ala Leu Val Thr Gly
 1 5 10 15
 Gly Thr Lys Gly Ile Gly Glu Ala Val Val Glu Glu Leu Ser Ile Leu
 20 25 30
 Gly Ala Arg Val His Thr Cys Ala Arg Asp Glu Thr Gln Leu Gln Glu
 35 40 45
 Arg Leu Arg Glu Trp Gln Glu Lys Gly Phe Gln Val Thr Thr Ser Ile
 50 55 60
 Cys Asp Val Ser Leu Arg Glu Gln Arg Glu Lys Leu Met Glu Thr Val
 65 70 75 80
 Ser Ser Leu Phe Gln Gly Lys Leu Asn Ile Leu Val Asn Asn Val Gly
 85 90 95
 Thr Leu Met Leu Lys Pro Thr Thr Glu Tyr Thr Ala Glu Glu Phe Ser
 100 105 110
 Phe Leu Met Ala Thr Asn Leu Asp Ser Ala Phe His Ile Ser Gln Leu
 115 120 125
 Ala His Pro Leu Leu Lys Ala Ser Gly Ser Gly Ser Ile Val Leu Met
 130 135 140
 Ser Ser Ile Ala Gly Val Val His Val Gly Val Gly Ser Ile Tyr Gly
 145 150 155 160
 Ala Thr Lys Gly Ala Met Asn Gln Leu Ala Arg Asn Leu Ala Cys Glu
 165 170 175
 Trp Ala Ser Asp Asn Ile Arg Thr Asn Ala Ile Cys Pro Trp Leu Ile
 180 185 190
 Thr Thr Pro Leu Ile Ser Asp Leu Leu Ser Val Glu Glu Met Lys Lys

195 200 205

Glu Ala Glu Glu Arg Thr Pro Met Gly Arg Val Gly Glu Ala Asn Glu
 210 215 220

Val Ser Pro Leu Val Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile
 225 230 235 240

Thr Gly Gln Val Ile Cys Val Asp Gly Gly Leu Thr Val Asn Gly Phe
 245 250 255

Ser Tyr Gln Pro His Ala
 260

<210> 7
 <211> 807
 <212> DNA
 <213> Arabidopsis thaliana

<400> 7

atgggctaaag cagaagaaaa cttgagagac aaatgtagat ggagccttgg aggcattgacc 60

gctctcgtta cgggtggctc caaaggcctc ggggaagctg tgggtggagga actagccatg 120

ttaggagcaa gaggccacac atgtgctaga aacgaaactc agcttcaaga atgcgtagct 180

gaatggcaag ctaaagggtt tgaggtcacc acttctgtat gcgacgtttc ttctcgtgac 240

caacgagaga aactcatgga aaacgttggc tctatcttcc aaggaaaact caacatcctt 300

gtaaacaatg cgggaacggg tataacaaag ccgaccacag agtatacagc acaagattac 360

tcatttctga tggctacaaa tctcgactca gcttttcatc tctcacagct cgcgcatcct 420

ttgttgaaag catctgggtc agggagcatc gtgctcatgt cctctactgc aggggttgta 480

catatcaatg ttggttccat ctatggagca actaaaggag ctatgaatca gctagcaaaa 540

aacttagcat gcgagtgggc aagggataac ataagggtta attctgtctg tccatgggtc 600

atagcaactc ctttatatct caatgatgaa gagttaaaga aagaagtgga gcgtaagaca 660

ccaatgggac gtgttgaaa cgcaaatgaa gtatcatcgc ttgtgcatt tctctgcttt 720

ccggctgctt cgtatataac aggtcagaca atctgcgtcg acggagggtt cacgggtcaac 780

tgcttttctt tcaagccagt gctttaa 807

<210> 8
 <211> 268
 <212> PRT
 <213> Arabidopsis thaliana

<400> 8

Met Ala Lys Ala Glu Glu Asn Leu Arg Asp Lys Cys Arg Trp Ser Leu
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Gly Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Leu Gly Glu
 20 25 30

Ala Val Val Glu Glu Leu Ala Met Leu Gly Ala Arg Val His Thr Cys
 35 40 45

Ala Arg Asn Glu Thr Gln Leu Gln Glu Cys Val Arg Glu Trp Gln Ala
 50 55 60

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

Gln Arg Glu Lys Leu Met Glu Asn Val Ala Ser Ile Phe Gln Gly Lys
 85 90 95

Leu Asn Ile Leu Val Asn Asn Ala Gly Thr Gly Ile Thr Lys Pro Thr
 100 105 110

Thr Glu Tyr Thr Ala Gln Asp Tyr Ser Phe Leu Met Ala Thr Asn Leu
 115 120 125

Asp Ser Ala Phe His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala
 130 135 140

Ser Gly Ser Gly Ser Ile Val Leu Met Ser Ser Thr Ala Gly Val Val
 145 150 155 160

His Ile Asn Val Gly Ser Ile Tyr Gly Ala Thr Lys Gly Ala Met Asn
 165 170 175

Gln Leu Ala Lys Asn Leu Ala Cys Glu Trp Ala Arg Asp Asn Ile Arg
 180 185 190

Val Asn Ser Val Cys Pro Trp Phe Ile Ala Thr Pro Leu Tyr Leu Asn
 195 200 205

Asp Glu Glu Leu Lys Lys Glu Val Glu Arg Lys Thr Pro Met Gly Arg
 210 215 220

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

Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp Gly Gly
 245 250 255

Phe Thr Val Asn Cys Phe Ser Phe Lys Pro Val Leu

260

265

<210> 9
 <211> 807
 <212> DNA
 <213> Arabidopsis thaliana

<400> 9
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 gctott₃₀gtta caggt₃₅ggctc caaagg₄₀cctc ggaga₄₅agctg tagt₅₀ggagga actag₅₅cctatg 120
 ttggg₆₀agcaa gagtt₆₅cacac atgt₇₀gccaga gacg₇₅aaactc agctt₈₀caaga acgct₈₅tacgt 180
 gaat₉₀ggcaag ccaa₉₅aggatt tgagg₁₀₀tcacc actt₁₀₅ctgtct gcg₁₁₀acgtctc ttct₁₁₅cg₁₂₀tgag 240
 caac₁₂₅gagaga aact₁₃₀catgga aacc₁₃₅gtttcc tctg₁₄₀tcttcc aagg₁₄₅aaaact caac₁₅₀atcctc 300
 gtaa₁₅₅ataatg cggg₁₆₀aacggg tata₁₆₅ataaag ccg₁₇₀agtacag agt₁₇₅atacagc aga₁₈₀agattac 360
 tcg₁₈₅tttctga tgg₁₉₀ctacaaa tct₁₉₅cgagtcc gct₂₀₀tttcatc tct₂₀₅cacagat cgc₂₁₀gcaccct 420
 ttgt₂₁₅tgaaag cct₂₂₀ctggttc cgg₂₂₅gagcatc gtg₂₃₀tcatgt cct₂₃₅ctgttgc tgg₂₄₀acttgtg 480
 cata₂₄₅ccggtg cat₂₅₀caatcta tgg₂₅₅agcatct aa₂₆₀aggagcta tga₂₆₅atcagct agga₂₇₀agaagc 540
 ttag₂₇₅catgcg agt₂₈₀gggcaag tgaca₂₈₅acata aggg₂₉₀ttaact ctg₂₉₅tgtgtcc atg₃₀₀ggtcata 600
 aca₃₀₅actcctt taac₃₁₀tagctt tatt₃₁₅ttcagt gat₃₂₀gaaaagt ta₃₂₅agaaaagc agt₃₃₀ggaggat 660
 aaga₃₃₅caccaa tgg₃₄₀gacgtgt tgg₃₄₅agaagca aat₃₅₀gaagtat cat₃₅₅cgcttgt cgc₃₆₀atttctc 720
 tgt₃₆₅tttccgg cag₃₇₀cttctta tatt₃₇₅acaggt cag₃₈₀actatct gcg₃₈₅tcgacgg agg₃₉₀tgcttcg 780
 gtga₃₉₅atggct tct₄₀₀ctttcaa gc₄₀₅cttag 807

<210> 10
 <211> 268
 <212> PRT
 <213> Arabidopsis thaliana

<400> 10

Met	Ala	Lys	Ala	Gly	Glu	Asn	Ser	Arg	Asp	Lys	Ser	Arg	Trp	Ser	Leu
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Glu	Gly	Met	Thr	Ala	Leu	Val	Thr	Gly	Gly	Ser	Lys	Gly	Leu	Gly	Glu
			20					25					30		
Ala	Val	Val	Glu	Glu	Leu	Ala	Met	Leu	Gly	Ala	Arg	Val	His	Thr	Cys
		35					40					45			
Ala	Arg	Asp	Glu	Thr	Gln	Leu	Gln	Glu	Arg	Leu	Arg	Glu	Trp	Gln	Ala
	50					55					60				
Lys	Gly	Phe	Glu	Val	Thr	Thr	Ser	Val	Cys	Asp	Val	Ser	Ser	Arg	Glu
65					70					75					80

Gln Arg Glu Lys Leu Met Glu Thr Val Ser Ser Val Phe Gln Gly Lys
 85 90 95

Leu Asn Ile Leu Val Asn Asn Ala Gly Thr Gly Ile Ile Lys Pro Ser
 100 105 110

Thr Glu Tyr Thr Ala Glu Asp Tyr Ser Phe Leu Met Ala Thr Asn Leu
 115 120 125

Glu Ser Ala Phe His Leu Ser Glu Ile Ala His Pro Leu Leu Lys Ala
 130 135 140

Ser Gly Ser Gly Ser Ile Val Phe Met Ser Ser Val Ala Gly Leu Val
 145 150 155 160

His Thr Gly Ala Ser Ile Tyr Gly Ala Ser Lys Gly Ala Met Asn Gln
 165 170 175

Leu Gly Arg Ser Leu Ala Cys Glu Trp Ala Ser Asp Asn Ile Arg Val
 180 185 190

Asn Ser Val Cys Pro Trp Val Ile Thr Thr Pro Leu Thr Ser Phe Ile
 195 200 205

Phe Ser Asp Glu Lys Leu Arg Lys Ala Val Glu Asp Lys Thr Pro Met
 210 215 220

Gly Arg Val Gly Glu Ala Asn Glu Val Ser Ser Leu Val Ala Phe Leu
 225 230 235 240

Cys Phe Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp
 245 250 255

Gly Gly Ala Ser Val Asn Gly Phe Ser Phe Lys Pro
 260 265

<210> 11
 <211> 807
 <212> DNA
 <213> Arabidopsis thaliana

<400> 11
 atggcaaaga gaggggaaag cttgagagac aaacctaaat ggagtcttga aggcattgact 60
 gctottgtta ccggtggatc taaaggcctc ggaaaagctg tgggtggagga actagccatg 120
 ttgggagcaa gagttcacac atgtgccaga gacgaaactc agcttcaaga aagcttacgt 180
 gaatggcaag caaaagggtt acaagtcacc acttctgttt gcgatgtttc ttctcgtgac 240

cagcgagaga aactcatgga aactgtttcc tctctcttcc aaggaaaact cagcatcctc 300
 gtaccaaatg tgggaatagg tgtactaaag cgcgacgactg agtgtacagc agaagagttc 360
 tcatttataa tagctacaaa tctggagtca actttccatt tctcgcaact cgcgcatcct 420
 ttattgaaag cctctgggtc agggaaacatt gtgctcatgt cttccgtggc tggagttgta 480
 aatttgggta atacatcaat ctatggagca accaaaggag ccatgaatca gctggcgaga 540
 aatttagcat gcgagtgggc gagtgataat ataagggcta attctgtttg tccatggttc 600
 attacaactc cgtcaactaa agatttctctc ggtgataaag atgtaaaaga aaaggtggag 660
 agtgtgacac cattgagacg tgttggagag gcaaatgaag tatcatcgct tgttgcattt 720
 ctatgtcttc cgcgagcttc ttatataaca ggtcaaacca tttgcgttga tggaggtttc 780
 actattaacg gcttctcttt gccttaa 807

<210> 12
 <211> 268
 <212> PRT
 <213> Arabidopsis thaliana

<400> 12

Met Ala Lys Arg Gly Glu Ser Leu Arg Asp Lys Pro Lys Trp Ser Leu
 1 5 10 15
 Glu Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Leu Gly Lys
 20 25 30
 Ala Val Val Glu Glu Leu Ala Met Leu Gly Ala Arg Val His Thr Cys
 35 40 45
 Ala Arg Asp Glu Thr Gln Leu Gln Glu Ser Leu Arg Glu Trp Gln Ala
 50 55 60
 Lys Gly Leu Gln Val Thr Thr Ser Val Cys Asp Val Ser Ser Arg Asp
 65 70 75 80
 Gln Arg Glu Lys Leu Met Glu Thr Val Ser Ser Leu Phe Gln Gly Lys
 85 90 95
 Leu Ser Ile Leu Val Pro Asn Val Gly Ile Gly Val Leu Lys Pro Thr
 100 105 110
 Thr Glu Cys Thr Ala Glu Glu Phe Ser Phe Ile Ile Ala Thr Asn Leu
 115 120 125
 Glu Ser Thr Phe His Phe Ser Gln Leu Ala His Pro Leu Leu Lys Ala
 130 135 140

Ser Gly Ser Gly Asn Ile Val Leu Met Ser Ser Val Ala Gly Val Val
 145 150 155 160

Asn Leu Gly Asn Thr Ser Ile Tyr Gly Ala Thr Lys Gly Ala Met Asn
 165 170 175

Gln Leu Ala Arg Asn Leu Ala Cys Glu Trp Ala Ser Asp Asn Ile Arg
 180 185 190

Ala Asn Ser Val Cys Pro Trp Phe Ile Thr Thr Pro Ser Thr Lys Asp
 195 200 205

Phe Leu Gly Asp Lys Asp Val Lys Glu Lys Val Glu Ser Val Thr Pro
 210 215 220

Leu Arg Arg Val Gly Glu Ala Asn Glu Val Ser Ser Leu Val Ala Phe
 225 230 235 240

Leu Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val
 245 250 255

Asp Gly Gly Phe Thr Ile Asn Gly Phe Ser Leu Pro
 260 265

<210> 13
 <211> 935
 <212> DNA
 <213> Arabidopsis thaliana

<400> 13
 aaagagagaa atggctggag cagagcaaag ccagagatgg agccttaagg ccaagaccgt 60
 acttgtaacc ggtggaacaa aaggcatcgg gcatgctata gtagaggaat ttgcaggatt 120
 tggagcagta atacatactt gtgctagaaa cgaatatgag cttaacgagt gtttaagcaa 180
 atggcaaaaag aagggttttc aagtcactgg ttcgggtctgt gacgcatctt tgagaccaga 240
 gagagaaaag ctaatgcaga ctgtgtcttc aatgtttggt ggaaagctcg atatcctcat 300
 aaacaatttg ggagcgattc ggtcaaaacc aacgctggat tacacagcag aagatttctc 360
 gtttcatatt tcgacaaact tagaatctgc ttatcatctg agccagcttg cgcatcctct 420
 gcttaaggct tcaggatgtg ggaacataat tttcatgtcc tctattgctg gagttgtatc 480
 tgctagtgtc ggctccatct actctgcaac gaaaggggca ttgaatcagc tagcaaggaa 540
 cttggcatgt gagtgggcta gtgatggcat tagggctaata gctgtggctc ctgcagtcac 600
 cgcaactcct ttggctgaag ctgtgtatga tgatgagttc aagaaagtgg tgatatcaag 660
 aaagccggtt gggcgtttcg gggagcctga agaagtgtcc tcgcttgggt cttttctatg 720

tatgcctgca gcttcttata taactgggtca aaccatttgt gtcgatggag gtctcactgt 780
 caatggcttc toctatcagc cacaaggtta aaatocaata tccaaaaaca aacacatttt 840
 caacttcatt actttgaaca ttgtcatctt gggctaagcc cggtttgaa gtatggattc 900
 tggccgtaac acttttagaaa ataaaaggaa acttc 935

<210> 14
 <211> 266
 <212> PRT
 <213> Arabidopsis thaliana

<400> 14

Met Ala Gly Ala Glu Gln Ser Gln Arg Trp Ser Leu Lys Ala Lys Thr
 1 5 10 15

Val Leu Val Thr Gly Gly Thr Lys Gly Ile Gly His Ala Ile Val Glu
 20 25 30

Glu Phe Ala Gly Phe Gly Ala Val Ile His Thr Cys Ala Arg Asn Glu
 35 40 45

Tyr Glu Leu Asn Glu Cys Leu Ser Lys Trp Gln Lys Lys Gly Phe Gln
 50 55 60

Val Thr Gly Ser Val Cys Asp Ala Ser Leu Arg Pro Glu Arg Glu Lys
 65 70 75 80

Leu Met Gln Thr Val Ser Ser Met Phe Gly Gly Lys Leu Asp Ile Leu
 85 90 95

Ile Asn Asn Leu Gly Ala Ile Arg Ser Lys Pro Thr Leu Asp Tyr Thr
 100 105 110

Ala Glu Asp Phe Ser Phe His Ile Ser Thr Asn Leu Glu Ser Ala Tyr
 115 120 125

His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala Ser Gly Cys Gly
 130 135 140

Asn Ile Ile Phe Met Ser Ser Ile Ala Gly Val Val Ser Ala Ser Val
 145 150 155 160

Gly Ser Ile Tyr Ser Ala Thr Lys Gly Ala Leu Asn Gln Leu Ala Arg
 165 170 175

Asn Leu Ala Cys Glu Trp Ala Ser Asp Gly Ile Arg Ala Asn Ala Val
 180 185 190

Ala Pro Ala Val Ile Ala Thr Pro Leu Ala Glu Ala Val Tyr Asp Asp
 195 200 205

Glu Phe Lys Lys Val Val Ile Ser Arg Lys Pro Leu Gly Arg Phe Gly
 210 215 220

Glu Pro Glu Glu Val Ser Ser Leu Val Ala Phe Leu Cys Met Pro Ala
 225 230 235 240

Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp Gly Gly Leu Thr
 245 250 255

Val Asn Gly Phe Ser Tyr Gln Pro Gln Gly
 260 265

<210> 15
 <211> 795
 <212> DNA
 <213> Arabidopsis thaliana

<400> 15
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 gtgtcaagga accaggaaga gctaaatgca tgcttgaatg attggaaagc gaatggttta 180
 gtcgtgtctg gttcggtttg tgatgcttcg gttagggatc agaggagaa gttgattcag 240
 gaagcttctt ctgccttcag tggcaagctc aacatcctta taaacaatgt tggaactaat 300
 gtcaggaaac caacagttga atactcaagc gaggaatatg ccaaatcat gtcgaccaac 360
 ttagaatccg ctttccattt atctcaaatt gctcatcctc ttctaaaagc atctggtgtc 420
 ggaagcattg tgttcatctc ctctgtagct ggcttgggtc atcttagcag tggatctatc 480
 tatggtgcaa ctaaaggagc acttaatcag cttacaagaa atctagcttg cgagtgggca 540
 agcgacaaca tcagaaccaa ttgctggtgc ccatggtaca tcaagacctc acttgtggaa 600
 acgctacttg agaagaaaga atttgtggag gctgtagttt caaggacccc acttgggctc 660
 gttggagaac cagaggaagt ctcatcgttg gttgcctttc tctgccttcc cgcacatctc 720
 tatattaccg gacaggtcat atccgtcgat ggaggattca ctgtcaacgg ttttagctat 780
 gctatgaagc cttaa 795

<210> 16
 <211> 264
 <212> PRT
 <213> Arabidopsis thaliana

<400> 16

Met Glu Thr Asp Lys Arg Trp Ser Leu Ala Gly Lys Thr Ala Leu Val
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 Thr Gly Gly Thr Arg Gly Ile Gly Arg Ala Val Val Glu Glu Leu Ala
 20 25 30
 Lys Phe Gly Ala Lys Val His Thr Cys Ser Arg Asn Gln Glu Glu Leu
 35 40 45
 Asn Ala Cys Leu Asn Asp Trp Lys Ala Asn Gly Leu Val Val Ser Gly
 50 55 60
 Ser Val Cys Asp Ala Ser Val Arg Asp Gln Arg Glu Lys Leu Ile Gln
 65 70 75 80
 Glu Ala Ser Ser Ala Phe Ser Gly Lys Leu Asn Ile Leu Ile Asn Asn
 85 90 95
 Val Gly Thr Asn Val Arg Lys Pro Thr Val Glu Tyr Ser Ser Glu Glu
 100 105 110
 Tyr Ala Lys Ile Met Ser Thr Asn Leu Glu Ser Ala Phe His Leu Ser
 115 120 125
 Gln Ile Ala His Pro Leu Leu Lys Ala Ser Gly Val Gly Ser Ile Val
 130 135 140
 Phe Ile Ser Ser Val Ala Gly Leu Val His Leu Ser Ser Gly Ser Ile
 145 150 155 160
 Tyr Gly Ala Thr Lys Gly Ala Leu Asn Gln Leu Thr Arg Asn Leu Ala
 165 170 175
 Cys Glu Trp Ala Ser Asp Asn Ile Arg Thr Asn Cys Val Ala Pro Trp
 180 185 190
 Tyr Ile Lys Thr Ser Leu Val Glu Thr Leu Leu Glu Lys Lys Glu Phe
 195 200 205
 Val Glu Ala Val Val Ser Arg Thr Pro Leu Gly Arg Val Gly Glu Pro
 210 215 220
 Glu Glu Val Ser Ser Leu Val Ala Phe Leu Cys Leu Pro Ala Ser Ser
 225 230 235 240
 Tyr Ile Thr Gly Gln Val Ile Ser Val Asp Gly Gly Phe Thr Val Asn
 245 250 255

Gly Phe Ser Tyr Ala Met Lys Pro
260

<210> 17
 <211> 789
 <212> DNA
 <213> Arabidopsis thaliana

<400> 17
 atggataaaa gatggagtct tcagggtatg accgctcttg taactggtgg agccagcgga 60
 atcgggcatg ctatagtaga ggagctagcc ggtcttgag ctagaatcta tgtatgcat 120
 atatctgaaa cactgctcaa tcaaagttta agtgaatggg aaaagaaagg gtttcaagta 180
 agtggttcaa tatgtgatgt atcctctcat tccgagaggg aaacacttat gcaaacagtc 240
 tcaaagatgt tcgatggcaa gctgaacatt cttgtgaaca atgttggcgt agttaatcca 300
 aagccaacaa tagaatatgt ggcagccgat ttctcgttca gtatttcaac aaacttgga 360
 tctgcttata accttagcca actttcacat cctctcctaa aagcttcaga atttggaagc 420
 atcatcttca tttcttctgt tggaggggtt gtgtcaatgg agtgtggatc tatctatagt 480
 ttaacgaaag gagctttgaa tcaactagca aaaactttgg catgtgaatg ggcaagagat 540
 ggcataagag ccaactctgt tgctcctaata tttatctaca ctgctatggc tcaacctttt 600
 ttcaaagacg ccgattacga gaagagtttg gttagtagaa ctccacttgg tcgcgctgga 660
 gagccaaatg aggtttcacc acttgtggct tttctgtgtc tacctgcagc ttcatatatt 720
 actggtcaga ccatttgtgt tgatggaggt ctcaactgtca atggtttctc ctataagcca 780
 caggcttga 789

<210> 18
 <211> 262
 <212> PRT
 <213> Arabidopsis thaliana

<400> 18

Met Asp Lys Arg Trp Ser Leu Gln Gly Met Thr Ala Leu Val Thr Gly
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Gly Ala Ser Gly Ile Gly His Ala Ile Val Glu Glu Leu Ala Gly Leu
 20 25 30

Gly Ala Arg Ile Tyr Val Cys Asp Ile Ser Glu Thr Leu Leu Asn Gln
 35 40 45

Ser Leu Ser Glu Trp Glu Lys Lys Gly Phe Gln Val Ser Gly Ser Ile
 50 55 60

Cys Asp Val Ser Ser His Ser Glu Arg Glu Thr Leu Met Gln Thr Val
65 70 75 80

Ser Lys Met Phe Asp Gly Lys Leu Asn Ile Leu Val Asn Asn Val Gly
85 90 95

Val Val Asn Pro Lys Pro Thr Ile Glu Tyr Val Ala Ala Asp Phe Ser
100 105 110

Phe Ser Ile Ser Thr Asn Leu Glu Ser Ala Tyr His Leu Ser Gln Leu
115 120 125

Ser His Pro Leu Leu Lys Ala Ser Glu Phe Gly Ser Ile Ile Phe Ile
130 135 140

Ser Ser Val Gly Gly Val Val Ser Met Glu Cys Gly Ser Ile Tyr Ser
145 150 155 160

Leu Thr Lys Gly Ala Leu Asn Gln Leu Ala Lys Thr Leu Ala Cys Glu
165 170 175

Trp Ala Arg Asp Gly Ile Arg Ala Asn Ser Val Ala Pro Asn Phe Ile
180 185 190

Tyr Thr Ala Met Ala Gln Pro Phe Phe Lys Asp Ala Asp Tyr Glu Lys
195 200 205

Ser Leu Val Ser Arg Thr Pro Leu Gly Arg Ala Gly Glu Pro Asn Glu
210 215 220

Val Ser Ser Leu Val Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile
225 230 235 240

Thr Gly Gln Thr Ile Cys Val Asp Gly Gly Leu Thr Val Asn Gly Phe
245 250 255

Ser Tyr Lys Pro Gln Ala
260

<210> 19

<211> 789

<212> DNA

<213> Arabidopsis thaliana

<400> 19

atggacaaaa gatggagtct caaaggtatg actgctcttg tgaccggtgg agccagtgga 60

atcggttatg ccatagtaga agagttggct ggttttggag ctagaatcca tgtatgtgac 120

atctctgaag ctaagctcaa tcaaagttta agcgaatggg aaaagaaagg ttttcaagta 180
 agtggctcag tttgtgatgt agcctctcgt cccgagagag aagaactgat gcaaaccgtc 240
 tcctcgcagt tcgatggcaa actcaacatt cttgtaagca atgtgggcgt aatccgctca 300
 aagccaacaa cagaatatac cgaagacgat ttcgcttttc atatatcatc aaacgtggaa 360
 gctgcttacc attttagcca gctttcacat cctctcctaa aggcttcagg ctatggaagc 420
 atcatctttg tttcctctat tgcaggggtt atatcttttg acgctggatc catttatggg 480
 ctaacaaaag gagctttgat tcagctagct aaaaatttgg catgtgaatg ggcaaaagac 540
 ggcataagag ccaacgctgt tgcgcctaata gtcacataata ctctctgtc tcaatcttat 600
 cttgaggacg tcagtttcaa gaaggcattg ttgagtagga ctccacttgg tcgtgttggg 660
 gagccaaatg aagttgcatc actagtggcc ttcttgtgtc tacctgcagc ttcttatatt 720
 actggtcaga ctatttgtgt tgatggagggt ctcaccgtta atggcttctc ctatcagcca 780
 gaggtttga 789

<210> 20
 <211> 262
 <212> PRT
 <213> Arabidopsis thaliana

<400> 20

Met Asp Lys Arg Trp Ser Leu Lys Gly Met Thr Ala Leu Val Thr Gly
 1 5 10 15

Gly Ala Ser Gly Ile Gly Tyr Ala Ile Val Glu Glu Leu Ala Gly Phe
 20 25 30

Gly Ala Arg Ile His Val Cys Asp Ile Ser Glu Ala Lys Leu Asn Gln
 35 40 45

Ser Leu Ser Glu Trp Glu Lys Lys Gly Phe Gln Val Ser Gly Ser Val
 50 55 60

Cys Asp Val Ala Ser Arg Pro Glu Arg Glu Glu Leu Met Gln Thr Val
 65 70 75 80

Ser Ser Gln Phe Asp Gly Lys Leu Asn Ile Leu Val Ser Asn Val Gly
 85 90 95

Val Ile Arg Ser Lys Pro Thr Thr Glu Tyr Thr Glu Asp Asp Phe Ala
 100 105 110

Phe His Ile Ser Ser Asn Val Glu Ala Ala Tyr His Phe Ser Gln Leu
 115 120 125

Ser His Pro Leu Leu Lys Ala Ser Gly Tyr Gly Ser Ile Ile Phe Val
 130 135 140

Ser Ser Ile Ala Gly Val Ile Ser Phe Asp Ala Gly Ser Ile Tyr Gly
 145 150 155 160

Leu Thr Lys Gly Ala Leu Ile Gln Leu Ala Lys Asn Leu Ala Cys Glu
 165 170 175

Trp Ala Lys Asp Gly Ile Arg Ala Asn Ala Val Ala Pro Asn Val Ile
 180 185 190

Asn Thr Pro Leu Ser Gln Ser Tyr Leu Glu Asp Val Ser Phe Lys Lys
 195 200 205

Ala Leu Leu Ser Arg Thr Pro Leu Gly Arg Val Gly Glu Pro Asn Glu
 210 215 220

Val Ala Ser Leu Val Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile
 225 230 235 240

Thr Gly Gln Thr Ile Cys Val Asp Gly Gly Leu Thr Val Asn Gly Phe
 245 250 255

Ser Tyr Gln Pro Glu Val
 260

<210> 21
 <211> 810
 <212> DNA
 <213> Arabidopsis thaliana

<400> 21
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 accggtgcag ccagcggaat cggttatgcc atagtagaag agttagctgg ttttggagct 120
 aaaatccaca tatgtgacat atccaaaact ttgctcaatc aaagtttaag cgaatgggaa 180
 aataaagggg ttcaagtgag tggttcagta tgtgatgtaa cctctcatcc tgagagagaa 240
 aaactgatgc aaaccgtctc ctcgattttc gatggcaaac tcaacattct tgtaaataac 300
 gtgggggtac ttcgcggaaa gccacaaca gaatatgtgg cagacgattt cacttttcat 360
 atatcaacta acttgaagc tgcttaccat ttttgtcaac tttcacatcc tctcttaaag 420
 gcttcaggct atggaagcat tgtattcctt tcctctgttg ctggggttgt atcactaatt 480
 gactgtggat ccatttatgg tctaacaaaa ggagctctaa atcaattagc tagaaacttg 540
 gcatgtgaat gggcaaaaga cggcataaga gccaatgctg ttgcacctaa tgtttgtcaag 600

actgctcagt ctcaatcggt tcttgaagac gtcagtaaaa aggagggatt gttgagtaga 660
 actccacttg gccgtggttg agagccgaat gaagtttcat cactagtggg cttcttgtgt 720
 ctacctgcag cttcttatat cacaggtcaa accatttgtg ttgatggagg tctcacgggt 780
 aacggtttct cctatcaacc acatgcttga 810

<210> 22
 <211> 269
 <212> PRT
 <213> Arabidopsis thaliana

<400> 22

Met Val Thr Arg Lys Met Asp Lys Arg Leu Trp Ser Leu Gln Gly Met
 1 5 10 15

Thr Ala Leu Val Thr Gly Ala Ala Ser Gly Ile Gly Tyr Ala Ile Val
 20 25 30

Glu Glu Leu Ala Gly Phe Gly Ala Lys Ile His Ile Cys Asp Ile Ser
 35 40 45

Lys Thr Leu Leu Asn Gln Ser Leu Ser Glu Trp Glu Asn Lys Gly Phe
 50 55 60

Gln Val Ser Gly Ser Val Cys Asp Val Thr Ser His Pro Glu Arg Glu
 65 70 75 80

Lys Leu Met Gln Thr Val Ser Ser Ile Phe Asp Gly Lys Leu Asn Ile
 85 90 95

Leu Val Asn Asn Val Gly Val Leu Arg Gly Lys Pro Thr Thr Glu Tyr
 100 105 110

Val Ala Asp Asp Phe Thr Phe His Ile Ser Thr Asn Leu Glu Ala Ala
 115 120 125

Tyr His Phe Cys Gln Leu Ser His Pro Leu Leu Lys Ala Ser Gly Tyr
 130 135 140

Gly Ser Ile Val Phe Leu Ser Ser Val Ala Gly Val Val Ser Leu Ile
 145 150 155 160

Asp Cys Gly Ser Ile Tyr Gly Leu Thr Lys Gly Ala Leu Asn Gln Leu
 165 170 175

Ala Arg Asn Leu Ala Cys Glu Trp Ala Lys Asp Gly Ile Arg Ala Asn
 180 185 190

Ala Val Ala Pro Asn Val Val Lys Thr Ala Gln Ser Gln Ser Phe Leu
 195 200 205

Glu Asp Val Ser Lys Lys Glu Gly Leu Leu Ser Arg Thr Pro Leu Gly
 210 215 220

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

Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp Gly
 245 250 255

Gly Leu Thr Val Asn Gly Phe Ser Tyr Gln Pro His Ala
 260 265

<210> 23
 <211> 969
 <212> DNA
 <213> Arabidopsis thaliana

<400> 23
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 cacaaatcct cttcctcttc cgtcttcgtc ccttacgcat cacagagttc cattgccatt 180
 acctccaagg aaagatggtc tctcaatgga atgtctgctc ttgtcaccgg cggcactcgt 240
 ggaattgggc gtgcgattgt ggaggaattg gctggctctg gagcagaagt tcacacttgc 300
 gctcggaatg agtacgagct ggagaattgt ttgagtgatt ggaaccgttc tggttttcga 360
 gttgctggat ccgtttgcca tgtctctgat cgatctcaga gagaggcttt gatggagacc 420
 gtgtcatctg tgtttgaagg gaagcttcat atccttgtaa acaatgttgg gacgaacatt 480
 aggaaaccaa tggtagagtt tactgctgga gaattttcga ctctgatgac tacgaatttc 540
 gaatcggttt tccatttatg tcaacttgct tatccattgc ttagagaatc taaagctgga 600
 agtgttgtgt tcatctcttc tgtttctggt tttgtttccc tcaagaatat gtcggtccaa 660
 tcttcaacca aaggagcaat taatcaactt acgagaagtc tggcttgcca gttggccaaa 720
 gacaatataa ggatcaatgc tgttgccccg tggatatca aaacatctat ggtggagcaa 780
 gtccttagca acaaagagta cctagaagaa gtctactcag taactcctct tggctgactt 840
 ggcgaaccga gagaggtgac ttctgcagtg gcttttttat gcctacctgc atcatcatat 900
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 catgattag 969

<210> 24

<211> 322

<212> PRT

<213> Arabidopsis thaliana

<400> 24

Met Val Leu Asp Met Ala Ser His Leu Tyr Thr Asn Pro Pro Gln Asn
 1 5 10 15

Leu His Phe Ile Ser Ser Ser Ser Ser Leu Lys Pro His Leu Cys Leu
 20 25 30

Ser Phe Lys Arg Ile Asn Pro Lys His Lys Ser Ser Ser Ser Ser Val
 35 40 45

Phe Val Pro Tyr Ala Ser Gln Ser Ser Ile Ala Ile Thr Ser Lys Glu
 50 55 60

Arg Trp Ser Leu Asn Gly Met Ser Ala Leu Val Thr Gly Gly Thr Arg
 65 70 75 80

Gly Ile Gly Arg Ala Ile Val Glu Glu Leu Ala Gly Leu Gly Ala Glu
 85 90 95

Val His Thr Cys Ala Arg Asn Glu Tyr Glu Leu Glu Asn Cys Leu Ser
 100 105 110

Asp Trp Asn Arg Ser Gly Phe Arg Val Ala Gly Ser Val Cys Asp Val
 115 120 125

Ser Asp Arg Ser Gln Arg Glu Ala Leu Met Glu Thr Val Ser Ser Val
 130 135 140

Phe Glu Gly Lys Leu His Ile Leu Val Asn Asn Val Gly Thr Asn Ile
 145 150 155 160

Arg Lys Pro Met Val Glu Phe Thr Ala Gly Glu Phe Ser Thr Leu Met
 165 170 175

Ser Thr Asn Phe Glu Ser Val Phe His Leu Cys Gln Leu Ala Tyr Pro
 180 185 190

Leu Leu Arg Glu Ser Lys Ala Gly Ser Val Val Phe Ile Ser Ser Val
 195 200 205

Ser Gly Phe Val Ser Leu Lys Asn Met Ser Val Gln Ser Ser Thr Lys
 210 215 220

Gly Ala Ile Asn Gln Leu Thr Arg Ser Leu Ala Cys Glu Trp Ala Lys

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 aaataaatat atcataatta cattatggat atacttaca tgt 1003

<210> 26
 <211> 262
 <212> PRT
 <213> Arabidopsis thaliana
 <400> 26

Met Asp Lys Arg Trp Ser Leu Gln Gly Met Thr Ala Leu Val Thr Gly
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Ala Ala Ser Gly Ile Gly Tyr Ala Ile Val Glu Glu Leu Ala Ser Phe
 20 25 30

Gly Ala Ile Ile His Ile Cys Asp Ile Ser Glu Thr Leu Leu Ser Gln
 35 40 45

Ser Leu Ser Glu Trp Glu Lys Lys Gly Phe Gln Val Ser Gly Ser Val
 50 55 60

Cys Asp Val Ala Ser Arg Pro Glu Arg Glu Lys Leu Met Gln Thr Val
 65 70 75 80

Ser Ser Leu Phe Asp Gly Lys Leu Asn Ile Leu Val Asn Asn Val Gly
 85 90 95

Val Ile Arg Gly Lys Pro Thr Thr Glu Tyr Val Ala Glu Asp Phe Ser
 100 105 110

Tyr His Ile Ser Thr Asn Leu Glu Pro Ala Phe His Phe Ser Gln Leu
 115 120 125

Ser His Pro Leu Leu Lys Ala Ser Gly Phe Gly Ser Ile Val Phe Met
 130 135 140

Ser Ser Ala Thr Gly Val Val Ser Val Gln Cys Gly Ser Ile Tyr Ser
 145 150 155 160

Leu Thr Lys Gly Ala Leu Asn Gln Leu Thr Arg Asn Leu Ala Cys Glu
 165 170 175

Trp Ala Lys Asp Gly Ile Arg Ala Asn Ala Val Ala Pro Asn Val Val
 180 185 190

Lys Thr Pro Leu Ser Gln Ser Tyr Leu Glu Asp Val Gly Phe Lys Glu
 195 200 205

Ala Leu Phe Ser Arg Thr Pro Leu Gly Arg Ala Gly Glu Pro Asn Glu
 210 215 220

Val Ala Ser Leu Val Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile
 225 230 235 240

Thr Gly Gln Thr Ile Cys Val Asp Gly Gly Leu Thr Val Asn Gly Phe
 245 250 255

Ser Tyr Gln Pro Glu Val
 260

<210> 27
 <211> 1040
 <212> DNA
 <213> Arabidopsis thaliana

<400> 27
 cggacgcgtg ggccggacgcg tggggcggacg cgtgggcgga cgcgtggccc gccgccgggg 60
 acgcgtgcct tgggttctgt ggttcataat cagaattggc taatacaggg gaaagcttga 120
 gagacaaacc tatatggagt cttgtaggca tgaccgctct tgtcaccggt ggctcaaaag 180
 gcatcggaca agctgtgggt gacgaactag ctacgttagg ggcaagaatc cacacatgtg 240
 ccagagacga aactcagctt caagaaagct tacatacgtg gcaagcagaa gggtttcagg 300
 ttaccacttc tgtctgacgac gtctcttctc gtgataaacg agagaagctc atggaaaccg 360
 tttccactat cttcgaccga aacctcaaca tacttgtcac caatgtggca acgtgtagag 420
 tcagtcccgc cctacaacat acagccgaag atttctcatt tacaatggca acgaatctcg 480
 agtcagcttt tcactctctg cagctcgcgc atcctttgtt gaaagcttct gggttcagggg 540
 gcatcgtgct catctcctcc gtatctggag ttgtacatgt caatggtgca tccatatatg 600
 gagtatctaa aggagctatg aatcagctag gaagaaactt agcgtgagag tgggcaagtg 660
 acaacataag gactaactct gtgtgtccat gggtcataga aactccttta gttaccgaaa 720
 gtcttagtaa tgaggagttt agaaaagaag tggagagtag ccccccaatg ggacgtgttg 780
 gagaagtaaa tgaagtatca tcgcttggg catttctttg tcttctgca gcttcttata 840
 ttacaggtca aaccatttgt gttgatggag gtttccctgt taatggtttt tctttcaagc 900
 ctctgcctta agacgtcgat tgttttcatt ttacgttact cccccccatt tacaatttgt 960
 ttagatatgt aaacataaac ccaattacc aaacaacaa aaaaaattgg cgtttggaat 1020
 gtaatccaac tctcccaat 1040

<210> 28
 <211> 253
 <212> PRT

<213> Arabidopsis thaliana

<400> 28

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

Val Asp Glu Leu Ala Thr Leu Gly Ala Arg Ile His Thr Cys Ala Arg
 20 25 30

Asp Glu Thr Gln Leu Gln Glu Ser Leu His Thr Trp Gln Ala Glu Gly
 35 40 45

Phe Gln Val Thr Thr Ser Val Cys Asp Val Ser Ser Arg Asp Lys Arg
 50 55 60

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

Ile Leu Val Thr Asn Val Ala Thr Cys Arg Val Ser Pro Ala Leu Gln
 85 90 95

His Thr Ala Glu Asp Phe Ser Phe Thr Met Ala Thr Asn Leu Glu Ser
 100 105 110

Ala Phe His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala Ser Gly
 115 120 125

Ser Gly Ser Ile Val Leu Ile Ser Ser Val Ser Gly Val Val His Val
 130 135 140

Asn Gly Ala Ser Ile Tyr Gly Val Ser Lys Gly Ala Met Asn Gln Leu
 145 150 155 160

Gly Arg Asn Leu Ala Cys Glu Trp Ala Ser Asp Asn Ile Arg Thr Asn
 165 170 175

Ser Val Cys Pro Trp Phe Ile Glu Thr Pro Leu Val Thr Glu Ser Leu
 180 185 190

Ser Asn Glu Glu Phe Arg Lys Glu Val Glu Ser Ser Pro Pro Met Gly
 195 200 205

Arg Val Gly Glu Val Asn Glu Val Ser Ser Leu Val Ala Phe Leu Cys
 210 215 220

Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp Gly
 225 230 235 240

Gly Phe Pro Val Asn Gly Phe Ser Phe Lys Pro Leu Pro
 245 250

<210> 29
 <211> 970
 <212> DNA
 <213> Arabidopsis thaliana

<400> 29
 agcccacgcg tccgtacggg gcgagaagac gacagaaggg gactagaaa aatggacaaa 60
 agattatgga gtcttcaagg tatgactgct cttgtgaccg gtgcagccag cggaatcggg 120
 tatgccatag tagaagagtt agctggtttt ggagctagaa tccacatatg tgacatatcc 180
 aaaactttgc tcaatcaaag ttaagcgaa tgggaaaata aagggtttca agtgagtggg 240
 tcagtatgtg atgtaacctc tcctcctgag agagaaaaac tgatgcaaac cgtctcctcg 300
 attttcgatg gcaaactcaa cattcttgta aataacgtgg ggtacttcg cggaaagcca 360
 acaacagaat atgtggcaga cgatttcact tttcatatat caactaactt ggaagctgct 420
 taccactttt gtcaactttc acatcctctc ttaaagactt caggctatgg aagcatcgtg 480
 ttcctttcct ctgtttctgg ggttgtgtca ataacttgcc tgggatcact ttatggctca 540
 ccaaaaggag ctctaaatca gctagctaga aatttggcat gtgaatggcc aaaagacggc 600
 ataagagcca atgcggttgc acctaatggt gtcaagactg ctcagtctca attctttctt 660
 caagacgtca gtaaaaagga gggattgttt agtagaactc cacttggtcg gtctggagag 720
 ccgaatgaag tagcatcact agtggctctc ttgtgtcttc ctgcagcttc ttatatcaca 780
 ggtcaaacca tttgtattga tggaggcctc acggtttacg gtttctcctc tcaaccacag 840
 gcttgaaaat ggtcttctcg tttcttctct gtttaaataat tctctcccta ataaaatctc 900
 tcctcagct atgagtttat gtcccttgta caaaaacaca caatgtcaaa ttgttgctct 960
 ttagtcttgt 970

<210> 30
 <211> 264
 <212> PRT
 <213> Arabidopsis thaliana

<400> 30

Met Asp Lys Arg Leu Trp Ser Leu Gln Gly Met Thr Ala Leu Val Thr
 1 5 10 15

Gly Ala Ala Ser Gly Ile Gly Tyr Ala Ile Val Glu Glu Leu Ala Gly
 20 25 30

Phe Gly Ala Arg Ile His Ile Cys Asp Ile Ser Lys Thr Leu Leu Asn
 35 40 45

Gln Ser Leu Ser Glu Trp Glu Asn Lys Gly Phe Gln Val Ser Gly Ser
 50 55 60

Val Cys Asp Val Thr Ser His Pro Glu Arg Glu Lys Leu Met Gln Thr
 65 70 75 80

Val Ser Ser Ile Phe Asp Gly Lys Leu Asn Ile Leu Val Asn Asn Val
 85 90 95

Gly Val Leu Arg Gly Lys Pro Thr Thr Glu Tyr Val Ala Asp Asp Phe
 100 105 110

Thr Phe His Ile Ser Thr Asn Leu Glu Ala Ala Tyr His Phe Cys Gln
 115 120 125

Leu Ser His Pro Leu Leu Lys Thr Ser Gly Tyr Gly Ser Ile Val Phe
 130 135 140

Leu Ser Ser Val Ser Gly Val Val Ser Ile Thr Cys Leu Gly Ser Leu
 145 150 155 160

Tyr Gly Leu Pro Lys Gly Ala Leu Asn Gln Leu Ala Arg Asn Leu Ala
 165 170 175

Cys Glu Trp Pro Lys Asp Gly Ile Arg Ala Asn Ala Val Ala Pro Asn
 180 185 190

Val Val Lys Thr Ala Gln Ser Gln Phe Phe Leu Gln Asp Val Ser Lys
 195 200 205

Lys Glu Gly Leu Phe Ser Arg Thr Pro Leu Gly Arg Ser Gly Glu Pro
 210 215 220

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

Tyr Ile Thr Gly Gln Thr Ile Cys Ile Asp Gly Gly Leu Thr Val Tyr
 245 250 255

Gly Phe Ser Ser Gln Pro Gln Ala
 260

<210> 31
 <211> 943
 <212> DNA
 <213> Arabidopsis thaliana

<400> 31
 ccggaattcc ttggtcgacc cacgcgtccg catacaagga ctatggataa aaggtggagt 60
 ctccaagggtt tgactgctct tgtgaccggt ggagccagcg gaatcgggtca tgctatagta 120
 gaagaactcg ccggttttgg ggccaaaatc catgtgtgtg acatatcgaa aactctgctc 180
 aatcaaagtt tatccgaatg ggagaagaaa gggtttcaag tgagtggttc agtctgcgat 240
 gcatccaatc gtctcgaaag agaaacactt atgcaaactg tcaccacaat atttgatggc 300
 aagcttaaca ttcttgtgaa caatgttggc acaattcgca caaagccaac aatagaatat 360
 gaggcagaag atttttcgtt ccttatttca acaaacttgg aatctgctta tcatctaagc 420
 caactttcac atccactcct aaaggcttca ggcaacggaa ttattacttt tatttcctct 480
 gctgcagga tcgtatcatt tgatgctgca tccatttatg gcctaacgaa aggagctttg 540
 aatcagctag cacgaaattt ggcgtgtgaa tgggcaaaaag acggcattcg agccaacgcg 600
 gttgcgccta attttatcac cactgctctg gctaaacctt ttctcgaaga cgctggtttt 660
 aacgagattt tgtcgagtag aactccactt ggtcgcgctg gagaaccaag agaggttgcc 720
 tcacttggg cttttctgtg tctacctgct gcttcatata ttactgggtca gaccatttgt 780
 gttgatggag gtctcactgt taatggcttc tcatatcagc cataggcttg agccatgtct 840
 tgtcttgtct tgtgtgttgt ggagtatggt catatgggtca gtatctccat aatctaaatc 900
 catagatatg tgagttgtgg agtagacaca atttttcaat aat 943

<210> 32

<211> 260

<212> PRT

<213> Arabidopsis thaliana

<400> 32

Met Asp Lys Arg Trp Ser Leu Gln Gly Leu Thr Ala Leu Val Thr Gly
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Gly Ala Ser Gly Ile Gly His Ala Ile Val Glu Glu Leu Ala Gly Phe
 20 25 30

Gly Ala Lys Ile His Val Cys Asp Ile Ser Lys Thr Leu Leu Asn Gln
 35 40 45

Ser Leu Ser Glu Trp Glu Lys Lys Gly Phe Gln Val Ser Gly Ser Val
 50 55 60

Cys Asp Ala Ser Asn Arg Leu Glu Arg Glu Thr Leu Met Gln Thr Val
 65 70 75 80

Thr Thr Ile Phe Asp Gly Lys Leu Asn Ile Leu Val Asn Asn Val Gly
 85 90 95

Thr Ile Arg Thr Lys Pro Thr Ile Glu Tyr Glu Ala Glu Asp Phe Ser
 100 105 110

Phe Leu Ile Ser Thr Asn Leu Glu Ser Ala Tyr His Leu Ser Gln Leu
 115 120 125

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

Ser Ser Ala Ala Gly Ile Val Ser Phe Asp Ala Ala Ser Ile Tyr Gly
 145 150 155 160

Leu Thr Lys Gly Ala Leu Asn Gln Leu Ala Arg Asn Leu Ala Cys Glu
 165 170 175

Trp Ala Lys Asp Gly Ile Arg Ala Asn Ala Val Ala Pro Asn Phe Ile
 180 185 190

Thr Thr Ala Leu Ala Lys Pro Phe Leu Glu Asp Ala Gly Phe Asn Glu
 195 200 205

Ile Leu Ser Ser Arg Thr Pro Leu Gly Arg Ala Gly Glu Pro Arg Glu
 210 215 220

Val Ala Ser Leu Val Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile
 225 230 235 240

Thr Gly Gln Thr Ile Cys Val Asp Gly Gly Leu Thr Val Asn Gly Phe
 245 250 255

Ser Tyr Gln Pro
 260

- <210> 33
- <211> 1002
- <212> DNA
- <213> Glycine max

<400> 33
 aagcaaacia caatggogaa ccoctgaaggc agtagcagag gctctagatg gtcaccttaag 60
 ggaacctactg ctctcgttac tggaggaacg cgtggaattg ggcacgctgt ggtggaggaa 120
 ctagcggagt ttggtgccac agtgtacact tgttcgagga atgaagaaga gctgaatgca 180
 tgcttgaagg agtggaaaga gaagggattt tcggtttctg ggttggtttg tgatgctct 240
 tctccacccc atagagagaa cctcattcaa caagtggcct ctgctttcaa cggcaagctc 300
 aacatacttg taaacaatgt tggaacaaat gtgaggaagc cgacaattga gtatacagcc 360

gaagaatatt caaaattgat ggcaactaac ttggactcca cataccattt gtgccaactt 420
 gcatacctc ttcttaaagc atctggaaat ggaagtattg tgtccatttc ctctgttgca 480
 agtcagacaa gcgtaggttc tggagccatt tacgcagcaa ctaaagctgc tattgatcag 540
 cttaccaa attttgcttg tgaatgggca aaagacaata taaggagcaa cgggtgttgca 600
 ccctggtata ccataacttc acttgtggaa cctttgcttg cgaacaaaca gcttggttagt 660
 gagataatat ctggaacgcc gataaagcgg atggcagaaa cacatgaagt ttcacacctg 720
 gtgactttcc tttgcctgcc agcagcatcc tacatcactg gacagattgt ttcagttgat 780
 ggaggattca ctgctaattg atttcaacc agcatgagaa tttcttaaag caaaatggcc 840
 atttttattg ttggattgtg tgaacaaaagt ggtggccact tgagttttgg accatctatg 900
 atcaacttgt tttgattggt caataaatta ttttagtatt ttagaaattt gcattgtcat 960
 tttcaacggc tatcattgtg agatgatggg tgctcttgga ac 1002

<210> 34
 <211> 271
 <212> PRT
 <213> Glycine max

<400> 34

Met Ala Asn Pro Glu Gly Ser Ser Arg Gly Ser Arg Trp Ser Leu Lys
 1 5 10 15

Gly Thr Thr Ala Leu Val Thr Gly Gly Thr Arg Gly Ile Gly His Ala
 20 25 30

Val Val Glu Glu Leu Ala Glu Phe Gly Ala Thr Val Tyr Thr Cys Ser
 35 40 45

Arg Asn Glu Glu Glu Leu Asn Ala Cys Leu Lys Glu Trp Lys Glu Lys
 50 55 60

Gly Phe Ser Val Ser Gly Leu Val Cys Asp Ala Ser Ser Pro Pro His
 65 70 75 80

Arg Glu Asn Leu Ile Gln Gln Val Ala Ser Ala Phe Asn Gly Lys Leu
 85 90 95

Asn Ile Leu Val Asn Asn Val Gly Thr Asn Val Arg Lys Pro Thr Ile
 100 105 110

Glu Tyr Thr Ala Glu Glu Tyr Ser Lys Leu Met Ala Thr Asn Leu Asp
 115 120 125

Ser Thr Tyr His Leu Cys Gln Leu Ala Tyr Pro Leu Leu Lys Ala Ser
 130 135 140

Gly Asn Gly Ser Ile Val Ser Ile Ser Ser Val Ala Ser Gln Thr Ser
 145 150 155 160

Val Gly Ser Gly Ala Ile Tyr Ala Ala Thr Lys Ala Ala Ile Asp Gln
 165 170 175

Leu Thr Lys Tyr Phe Ala Cys Glu Trp Ala Lys Asp Asn Ile Arg Ser
 180 185 190

Asn Gly Val Ala Pro Trp Tyr Thr Ile Thr Ser Leu Val Glu Pro Leu
 195 200 205

Leu Ala Asn Lys Gln Leu Val Ser Glu Ile Ile Ser Arg Thr Pro Ile
 210 215 220

Lys Arg Met Ala Glu Thr His Glu Val Ser Ser Leu Val Thr Phe Leu
 225 230 235 240

Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Ile Val Ser Val Asp
 245 250 255

Gly Gly Phe Thr Ala Asn Gly Phe Gln Pro Ser Met Arg Ile Ser
 260 265 270

<210> 35
 <211> 1084
 <212> DNA
 <213> Glycine max

<400> 35
 atggaaaggc gttttaaaaa aagagtggaa gatggcaatg gcaggaagca gcattaacag 60
 aggagaaaga tggctctctca atggaatgac tgctcttgtc accggcggca ctctgtgggat 120
 cgggcactcc atagtgagtg acttggctgc gtttggcgct gcagtgcaca cctgttccag 180
 gacccaaaca gagctcaaca aatgcttaca agagtggcag agtcaggggt ttcaggtaac 240
 tgggtcgctc tgtgacgtgt cctcaccacc ccagagagag aagctcattc aggaagttgc 300
 atccaccttc aatggcaagc ttaacatcta cgtgaacaat gttggaataa acattagaaa 360
 gcccaaccatt gactacactg ctgaagaata ttcacagatt atgacagtta atttagactc 420
 ctcatccat ctgtgccagc ttgcatatcc tcttctgaaa gcatctgaaa aggaagcat 480
 tgtgttcatt tcatctgttg ctggtgtagt gagtttaggt actggagctg tctttgcagc 540
 aagtaaagct gcaattaatc agcttataaa aaacctggct tgtgactggg ccaaagacaa 600
 catacggagc aactgtgttg taccatgggc aaccagaacc ccggtttag aacatttgtt 660

caaagaccaa aagtttgtgg atgatattat gtctcggact ccaattaaac gtatagcaga 720
 accagaagaa gtgtcatcgt tggatgaattt tctttgcttg cctgctgctt cattcatcac 780
 tggacaggtt atttgtgttg atggaggatt aactgtgaat ggatttcaac ccagcatgag 840
 aattacctga aattacttca ctgctttctc tttgtggaac aagcgattct catttgattt 900
 caacatgttt gtgacaagtg atgcgcaatt tcgttttatac tgccttctag ttgcatatga 960
 aatgaatgg cctcatttaa ttccatatga aaaacttata taataaatgg aacatcctat 1020
 ttaatagcgt tgatgtttct tttttccttt tataataata ataataataa tacatggcta 1080
 ttgt 1084

<210> 36
 <211> 272
 <212> PRT
 <213> Glycine max
 <400> 36

Met Ala Met Ala Gly Ser Ser Ile Asn Arg Gly Glu Arg Trp Ser Leu
 1 5 10 15

Asn Gly Met Thr Ala Leu Val Thr Gly Gly Thr Arg Gly Ile Gly His
 20 25 30

Ser Ile Val Ser Asp Leu Ala Ala Phe Gly Ala Ala Val His Thr Cys
 35 40 45

Ser Arg Thr Gln Thr Glu Leu Asn Lys Cys Leu Gln Glu Trp Gln Ser
 50 55 60

Gln Gly Phe Gln Val Thr Gly Ser Leu Cys Asp Val Ser Ser Pro Pro
 65 70 75 80

Gln Arg Glu Lys Leu Ile Gln Glu Val Ala Ser Thr Phe Asn Gly Lys
 85 90 95

Leu Asn Ile Tyr Val Asn Asn Val Gly Ile Asn Ile Arg Lys Pro Thr
 100 105 110

Ile Glu Tyr Thr Ala Glu Glu Tyr Ser Gln Ile Met Thr Val Asn Leu
 115 120 125

Asp Ser Ser Phe His Leu Cys Gln Leu Ala Tyr Pro Leu Leu Lys Ala
 130 135 140

Ser Glu Lys Gly Ser Ile Val Phe Ile Ser Ser Val Ala Gly Val Val
 145 150 155 160

Ser Leu Gly Thr Gly Ala Val Phe Ala Ala Ser Lys Ala Ala Ile Asn
 165 170 175

Gln Leu Thr Lys Asn Leu Ala Cys Asp Trp Ala Lys Asp Asn Ile Arg
 180 185 190

Ser Asn Cys Val Val Pro Trp Ala Thr Arg Thr Pro Val Val Glu His
 195 200 205

Leu Phe Lys Asp Gln Lys Phe Val Asp Asp Ile Met Ser Arg Thr Pro
 210 215 220

Ile Lys Arg Ile Ala Glu Pro Glu Glu Val Ser Ser Leu Val Asn Phe
 225 230 235 240

Leu Cys Leu Pro Ala Ala Ser Phe Ile Thr Gly Gln Val Ile Cys Val
 245 250 255

Asp Gly Gly Leu Thr Val Asn Gly Phe Gln Pro Ser Met Arg Ile Thr
 260 265 270

<210> 37
 <211> 1133
 <212> DNA
 <213> Glycine max

<400> 37
 cccacgattc ggctcgagcg gatatctgac aggttcatca tcataacctt caatgaaaag 60
 aaatggctga ggcaagcatt ggagcaaaa gcagcagatg gtctttacag ggaatgacag 120
 ctctcgtcac cggtgatcc aaaggaatcg gatatgctat cgtggaggag ttggcacagc 180
 ttggagccac tgtgcacact tgcgctcggg acgaagctga actcaatgaa tccttaaatg 240
 aatggaacac aaaaggatac agagtaactg gttccgtctg tgacgtggcg tctcgtgcag 300
 aaagacaaga cctcatagct agagtctcca atgagtttaa tggcaaactc aatatccttg 360
 taaacaacgt gggaacaaac gtaccgaaac atacccttga tgttacggag gaagacttct 420
 catttctgat aaatacaaat cttgaatctg cttaccacct aagccagctt gcacatcctc 480
 tcctgaaagc ttcagaggct gcaaaccatca tttttatata ctccattgct ggtgtgctat 540
 caataggtat aggatccact tatggtgcaa caaaggagc aatgaaccaa ctgactaaaa 600
 atttggcatg tgaatgggcc aaagacaata taaggactaa ttgcgttgca ccagggccaa 660
 ttagaacccc tottgggtgac aagcacttta aagaggaaaa acttaataat agtctgattg 720
 cgcaacccc tcttgacgg attggagagg cagaggaggt ttcttcgttg gtggcattcc 780
 tctgcttacc tgcagcatct tacataacag gacaaacctt ttgtgttgat ggcggcttca 840

ccgtgaacgg tctctacata agctaggcctt attgtgacag tttcaatggt ttgttttagtc 900
 tgttacactc tacttgtatt agtttctctt taagttgtgg agagtggaga ccctatatct 960
 gtttttaagg acgtgataat aacagtactc agtgataagt ttgtttttct tgttttgatt 1020
 tagtcacatg tcccatgatt ctaggagcag tataatcgtg ggcagtttat cctatttccc 1080
 gttggttgta aggctttcaa taataataat aataatagcc atgtttgctt ttc 1133

<210> 38
 <211> 267
 <212> PRT
 <213> Glycine max

<400> 38

Met Ala Glu Ala Ser Ile Gly Ser Lys Ser Ser Arg Trp Ser Leu Gln
 1 5 10 15

Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Tyr Ala
 20 25 30

Ile Val Glu Glu Leu Ala Gln Leu Gly Ala Thr Val His Thr Cys Ala
 35 40 45

Arg Asn Glu Ala Glu Leu Asn Glu Ser Leu Asn Glu Trp Asn Thr Lys
 50 55 60

Gly Tyr Arg Val Thr Gly Ser Val Cys Asp Val Ala Ser Arg Ala Glu
 65 70 75 80

Arg Gln Asp Leu Ile Ala Arg Val Ser Asn Glu Phe Asn Gly Lys Leu
 85 90 95

Asn Ile Leu Val Asn Asn Val Gly Thr Asn Val Pro Lys His Thr Leu
 100 105 110

Asp Val Thr Glu Glu Asp Phe Ser Phe Leu Ile Asn Thr Asn Leu Glu
 115 120 125

Ser Ala Tyr His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala Ser
 130 135 140

Glu Ala Ala Asn Ile Ile Phe Ile Ser Ser Ile Ala Gly Val Leu Ser
 145 150 155 160

Ile Gly Ile Gly Ser Thr Tyr Gly Ala Thr Lys Gly Ala Met Asn Gln
 165 170 175

Leu Thr Lys Asn Leu Ala Cys Glu Trp Ala Lys Asp Asn Ile Arg Thr
 180 185 190

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

Phe Lys Glu Glu Lys Leu Asn Asn Ser Leu Ile Ala Arg Thr Pro Leu
 210 215 220

Gly Arg Ile Gly Glu Ala Glu Glu Val Ser Ser Leu Val Ala Phe Leu
 225 230 235 240

Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp
 245 250 255

Gly Gly Phe Thr Val Asn Gly Leu Tyr Ile Ser
 260 265

<210> 39
 <211> 1113
 <212> DNA
 <213> Glycine max

<400> 39
 cggatatctc aaaggtttat ccttcagtga agagaaatgg cagctgaggc aaatdddgat 60
 agcaaaagca gcaggtggtc tttgcagggg atgacagctc tcgttaccgg tggttccaaa 120
 ggaatcggat atgctatcgt ggaggagttg gcacaacttg gagccactgt acacacttgc 180
 gctcgcaacg aagctgaact caataaatcc ttaaatgaat ggaacacaaa aggatacaga 240
 gtaactgggt cggtcctgta cgtggcgtct cgtgcagaaa gacaagacct catagctaga 300
 gtctcaaatg agtttaaatg caaactcaat atccttghta acaatgtggg aacaaacata 360
 cagaaagaga ccttggattt cacagaggaa gatttcacat ttctgggtgaa tacgaatctt 420
 gaatcttggt tccacctaag ccagcttgea catcctctcc taaaagcttc agaagctgca 480
 aacatcattc tcatatcctc cattgctggt gtggtagctt caaatatagt atccgttgtg 540
 tatggtgcaa caaaaggagc aatgaaccaa atgacaaaac atttggcatg tgaatgggcc 600
 aaagacaata taaggactaa ttgcgttgca ccagggccaa ttagaacccc tcttggtgac 660
 aagcacttta aagaggaaaa acttaataat agtctgattg cgccaacccc tcttggacgg 720
 attggagagg cagaggaggt ttcttcgttg gtggcattcc tctgcttacc tgcagcatct 780
 tacataacag gacaaacat ttgtgttgat ggcggcttca ccgtgaacgg tctctacata 840
 agctaggctt attgtgacag tttcaatggt ttgttttagtc tgttacactc tacttgtatt 900
 agtttctctt taagttgtgg agagtggaga ccctatatct gtttttaagg acgtgataat 960
 aacagtactc agtgataagt ttgtttttct tgttttgatt tagtcacatg tcccatgatt 1020

ctaggagcag tataatcgtg ggcagtttat cctatttccc gttggttgta aggcattcaa 1080

taataataat aataatagcc atgtttgctt ttc 1113

<210> 40

<211> 269

<212> PRT

<213> Glycine max

<400> 40

Met Ala Ala Glu Ala Asn Phe Asp Ser Lys Ser Ser Arg Trp Ser Leu
1 5 10 15

Gln Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Tyr
20 25 30

Ala Ile Val Glu Glu Leu Ala Gln Leu Gly Ala Thr Val His Thr Cys
35 40 45

Ala Arg Asn Glu Ala Glu Leu Asn Lys Ser Leu Asn Glu Trp Asn Thr
50 55 60

Lys Gly Tyr Arg Val Thr Gly Ser Val Arg Asp Val Ala Ser Arg Ala
65 70 75 80

Glu Arg Gln Asp Leu Ile Ala Arg Val Ser Asn Glu Phe Asn Gly Lys
85 90 95

Leu Asn Ile Leu Val Asn Asn Val Gly Thr Asn Ile Gln Lys Glu Thr
100 105 110

Leu Asp Phe Thr Glu Glu Asp Phe Thr Phe Leu Val Asn Thr Asn Leu
115 120 125

Glu Ser Cys Phe His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala
130 135 140

Ser Glu Ala Ala Asn Ile Ile Leu Ile Ser Ser Ile Ala Gly Val Val
145 150 155 160

Ala Ser Asn Ile Val Ser Val Val Tyr Gly Ala Thr Lys Gly Ala Met
165 170 175

Asn Gln Met Thr Lys His Leu Ala Cys Glu Trp Ala Lys Asp Asn Ile
180 185 190

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

Lys His Phe Lys Glu Glu Lys Leu Asn Asn Ser Leu Ile Ala Arg Thr
 210 215 220

Pro Leu Gly Arg Ile Gly Glu Ala Glu Glu Val Ser Ser Leu Val Ala
 225 230 235 240

Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys
 245 250 255

Val Asp Gly Gly Phe Thr Val Asn Gly Leu Tyr Ile Ser
 260 265

<210> 41
 <211> 1157
 <212> DNA
 <213> Glycine max

<400> 41
 caaatccata tcacagaaaa ccattttggc ttctcaaaac cccactcatc atccatatca 60
 caaaaaacta ttttggcttc tgcctttcct tttccaggga gctcattgac aatggctgaa 120
 acaaagtggg tcatgaagga caaaagatgg tctctccatg gaatgacagc tctagtcaca 180
 ggaggcaccg gaggcatagg gcatgccatt gttgaagagt tagctgagtt tggagcaact 240
 gttcatatat gtgcacgtaa tcaagatgat attgataaat gtttagaaga gtggaaaaac 300
 aagggactta atgtgactgg ttcagtatgt gatttactat gttctgacca acgtaaaaga 360
 ttaatggaaa ttgttggctc catctttcat ggaaagctca atattctagt gaacaatgct 420
 gctacaaata taacaaagaa gataacagat tacacagcag aggatataatc agccataatg 480
 ggcaccaatt ttgagtccgt ttaccatttg tgtcaagttg cacaccact tctaaaagat 540
 tctgggaatg ggagcatagt atttatttct tccgtagcag gtttaaaagc tcttctgtg 600
 ttctctgttt atgcagcctc taaaggagcc atgaatcaat tcaccaaaaaa cttggcattg 660
 gaatgggcaa aggataatat tcgtgcaaat gctgttgccc ctggacctgt taagactaaa 720
 cttttggagt gtatcgtgaa ttcttcggaa gggaatgagt ctataaatgg aatagtgctc 780
 caaacatttg ttggtcgcac gggagaaact aaagagatat cagcattagt tgcttttctt 840
 tgccttcggg ctgcatcata catcactgga caggttatat gtgtagatgg gggtttcaca 900
 acttagattg ttgttaatga ttttgttgcg gaattaaatc ttgtgtttcg tgttcctcca 960
 aacgactcat gcttatgtcg ttggttaggc ttatctcctg ttcttaaaca taaccatgtg 1020
 ttcttgttta gtttcgtgtg tctttctctc agaattcaac aaggcagaac aattgttaat 1080
 tgctgctcc ccatgactcc atgttatctt atttgttctt tagatgaaac ataacttcaa 1140
 ttctcgtaa aaaaaca 1157

<210> 42

<211> 264

<212> PRT

<213> Glycine max

<400> 42

Met Ala Glu Thr Lys Trp Val Met Lys Asp Lys Arg Trp Ser Leu His
1 5 10 15

Gly Met Thr Ala Leu Val Thr Gly Gly Thr Arg Gly Ile Gly His Ala
20 25 30

Ile Val Glu Glu Leu Ala Glu Phe Gly Ala Thr Val His Ile Cys Ala
35 40 45

Arg Asn Gln Asp Asp Ile Asp Lys Cys Leu Glu Glu Trp Lys Asn Lys
50 55 60

Gly Leu Asn Val Thr Gly Ser Val Cys Asp Leu Leu Cys Ser Asp Gln
65 70 75 80

Arg Lys Arg Leu Met Glu Ile Val Gly Ser Ile Phe His Gly Lys Leu
85 90 95

Asn Ile Leu Val Asn Asn Ala Ala Thr Asn Ile Thr Lys Lys Ile Thr
100 105 110

Asp Tyr Thr Ala Glu Asp Ile Ser Ala Ile Met Gly Thr Asn Phe Glu
115 120 125

Ser Val Tyr His Leu Cys Gln Val Ala His Pro Leu Leu Lys Asp Ser
130 135 140

Gly Asn Gly Ser Ile Val Phe Ile Ser Ser Val Ala Gly Leu Lys Ala
145 150 155 160

Leu Pro Val Phe Ser Val Tyr Ala Ala Ser Lys Gly Ala Met Asn Gln
165 170 175

Phe Thr Lys Asn Leu Ala Leu Glu Trp Ala Lys Asp Asn Ile Arg Ala
180 185 190

Asn Ala Val Ala Pro Gly Pro Val Lys Thr Lys Leu Leu Glu Cys Ile
195 200 205

Val Asn Ser Ser Glu Gly Asn Glu Ser Ile Asn Gly Ile Val Ser Gln
210 215 220

Thr Phe Val Gly Arg Met Gly Glu Thr Lys Glu Ile Ser Ala Leu Val
 225 230 235 240

Ala Phe Leu Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Val Ile
 245 250 255

Cys Val Asp Gly Gly Phe Thr Thr
 260

<210> 43
 <211> 1366
 <212> DNA
 <213> Zea mays

<400> 43
 atccacttgc ctcggccgct atcgtgaccg ggaacaacaa ctgcgagatg gccacggtgg 60
 agacctcggg cacggcgata gggtcctccg ggagatgggc actacacggc aagacagccc 120
 tcgtcaccgg cggcaccggc ggcatcgggc gtgctgtagt ggaggagctg gcggcgctgg 180
 gggcgggccgt gcacacatgc tcccggaagg cggaggagct cggcgagcgc atcaaggagt 240
 gggaggccag gggattcagc gttaccgggt ccgtctgcga cctctccgag agggaccagc 300
 gggagcggtt gctccgcgag gttgcccacc gcttcggcgg caagctcaac atcctcgtaa 360
 acaatgtagg aacaaacata aggaaaccaa ctactgagtt tactgcagag gaatactcgt 420
 ttctgatggc tactaatcct gaatctgcat atcacttgtg ccaaattgca catcctcttt 480
 tgaaattatc tgggtcaggc agcattatat tcatatcadc tgttgctgga gcgataggaa 540
 tctttagtgg aactatataat gctatgacta aagggtgccat taaccagcta accaagaatt 600
 tagcttgtga atgggctaag gacaacataa gagccaactc tgtcgcctcc tgggtacatca 660
 ccacttcaact tacggaagga attttgcaa ataagaactt tgaggaacaa gttgtgagtc 720
 gaactccgct tggacgtgtc ggagaacctg gagaagtatc ggcacttggt gcttttcttt 780
 gcatgccggg ttccacttat attagcggcc agacgattgc ggtcgacgga ggtatgactg 840
 tgaacggggt ttaccctccc aagccctagg cggcagctgc catgtttctt tgtgctgagg 900
 aaacgacaat agctgttggt cttggttggt gagaaaaaaaa atctaataaa atgtatgtaa 960
 aggtaagggt gtcatgttga tactgttcat tacattacat aattctatca gggcaatgtg 1020
 aagaacgttg tgttttagatc aataaaaaat atagggtggt ggcataatit ttggacaaag 1080
 acggaaactt aggaaacaaa ttttctgatg attgtgaacg gtttctgtaa agcattcagt 1140
 gcccttttac tactgtattg ttgtggttgt gtttttttgt aatggttgga tttttgtggc 1200
 tgattgttct tttttttttt ttcgatctcc actatctttt ctttggttgt ttccogtgtt 1260
 ttttgctttt ggttgttttt ttggtttttt tttccttgtt ttgtattgta tgtcttgttt 1320

ttgctttttt ttttggtttt ttgtgttggt gatttttttt cctttc

1366

<210> 44
 <211> 273
 <212> PRT
 <213> Zea mays

<400> 44

Met Ala Thr Val Glu Thr Ser Gly Thr Ala Ile Gly Ser Ser Gly Arg
 1 5 10 15

Trp Ala Leu His Gly Lys Thr Ala Leu Val Thr Gly Gly Thr Arg Gly
 20 25 30

Ile Gly Arg Ala Val Val Glu Glu Leu Ala Ala Leu Gly Ala Ala Val
 35 40 45

His Thr Cys Ser Arg Lys Ala Glu Glu Leu Gly Glu Arg Ile Lys Glu
 50 55 60

Trp Glu Ala Arg Gly Phe Ser Val Thr Gly Ser Val Cys Asp Leu Ser
 65 70 75 80

Glu Arg Asp Gln Arg Glu Arg Leu Leu Arg Glu Val Ala Asp Arg Phe
 85 90 95

Gly Gly Lys Leu Asn Ile Leu Val Asn Asn Val Gly Thr Asn Ile Arg
 100 105 110

Lys Pro Thr Thr Glu Phe Thr Ala Glu Glu Tyr Ser Phe Leu Met Ala
 115 120 125

Thr Asn Leu Glu Ser Ala Tyr His Leu Cys Gln Ile Ala His Pro Leu
 130 135 140

Leu Lys Leu Ser Gly Ser Gly Ser Ile Ile Phe Ile Ser Ser Val Ala
 145 150 155 160

Gly Ala Ile Gly Ile Phe Ser Gly Thr Ile Tyr Ala Met Thr Lys Gly
 165 170 175

Ala Ile Asn Gln Leu Thr Lys Asn Leu Ala Cys Glu Trp Ala Lys Asp
 180 185 190

Asn Ile Arg Ala Asn Ser Val Ala Pro Trp Tyr Ile Thr Thr Ser Leu
 195 200 205

Thr Glu Gly Ile Leu Ala Asn Lys Asn Phe Glu Glu Gln Val Val Ser
 210 215 220

Arg Thr Pro Leu Gly Arg Val Gly Glu Pro Gly Glu Val Ser Ala Leu
 225 230 235 240

Val Ala Phe Leu Cys Met Pro Gly Ser Thr Tyr Ile Ser Gly Gln Thr
 245 250 255

Ile Ala Val Asp Gly Gly Met Thr Val Asn Gly Phe Tyr Pro Pro Lys
 260 265 270

Pro

- <210> 45
- <211> 1295
- <212> DNA
- <213> Zea mays

<400> 45
 caaatatggg caggcactag ctagctaggc tgcaaagagc caaagactgc cggcaccgct 60
 gctagcttgc tagcatactt ccggtcggca agcagtcgat cgtcgcgatg cgccaggagg 120
 cagcggcgag cgggtggagcc tggccggcgc gacagcgcgtg gtcaccgggtg gcagcaaggg 180
 gatcgggcaa gccgtcgtgg aggagctggc caggctcggc gcgcgcgtgc acacgtgcgc 240
 ccgcagcgcg gcggacctgg aggagtgccg ccggcgggtg gccgagaagg ggctccgcgt 300
 caccgtctcc gtgtgcgacg tcgccgtgcg ccgccaccgg gagaggctcg tcctggacac 360
 ggtcagcggg gccttcgacg gcaagctcga tatcctggtc aacaacgctg cgctgctgct 420
 gctcaagccg gcggcggagt gggcggcgga ggactacgcg cggatcatgg cgaccaacct 480
 ggagtcgtgc ttgcacatct cccagctcgc gcaccgcgtg ctctcaacg cctccgtcgc 540
 cggaggggcg agcatcgtca atgtctctc catcgcacg gtccttggct tcccgcagga 600
 agtcatgtac agcgtcacca aaggaggact gaatcagatg acgaggagcc tagctgtgga 660
 gtgggcctgc gataggatcc gtgtgaactg cgtcgcgccg ggcgtgatca tgacggacat 720
 gggtaaagag ctaccggcgg cgttggtgga gcaggagcgg tcacgcatcc cgctgcggcg 780
 gaccggcgag ccggaggagg tggcgtccct ggtgtcgttc ctctgcatgc cggcggcgtc 840
 ctacgtcacc gggcagggtca tcttcgtcga cggcggccgg accattagtg gcgcctgatc 900
 gacgacgacg gccagctata tatatagcga agaataaatg tggatctatt ctctattcta 960
 ctgtatattga atttgttttc gttcttgtct tcttgatcgg ccatgtatca ccgtcagacg 1020
 tcagcggcaa taatccggcc gccgtcggct tgccgggagg gagataccca actagcgaca 1080
 tgcaagctgc ccgatcagtc gtctcgtctc gcctcgcggc ccggcaaggt gcagccactg 1140

tcgttgacctg cagttctgtt cctttttcta ctatggtttc ttgtattcac tgcccactgg 1200
 tgatgtttct gtgataacga cttcgacaga ggggatacat tttaaaatcc ttgtgctaata 1260
 cattttttgt gcacatcaga gttttttgag aacag 1295

<210> 46
 <211> 263
 <212> PRT
 <213> Zea mays

<400> 46

Met Ala Pro Gly Gly Ser Gly Glu Arg Trp Ser Leu Ala Gly Ala Thr
 1 5 10 15

Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Gln Ala Val Val Glu
 20 25 30

Glu Leu Ala Arg Leu Gly Ala Arg Val His Thr Cys Ala Arg Ser Ala
 35 40 45

Ala Asp Leu Glu Glu Cys Arg Arg Arg Trp Ala Glu Lys Gly Leu Arg
 50 55 60

Val Thr Val Ser Val Cys Asp Val Ala Val Arg Ala Asp Arg Glu Arg
 65 70 75 80

Leu Val Leu Asp Thr Val Ser Gly Ala Phe Asp Gly Lys Leu Asp Ile
 85 90 95

Leu Val Asn Asn Ala Ala Leu Leu Leu Leu Lys Pro Ala Ala Glu Trp
 100 105 110

Ala Ala Glu Asp Tyr Ala Arg Ile Met Ala Thr Asn Leu Glu Ser Cys
 115 120 125

Leu His Ile Ser Gln Leu Ala His Pro Leu Leu Leu Asn Ala Ser Val
 130 135 140

Ala Gly Gly Ala Ser Ile Val Asn Val Ser Ser Ile Ala Ser Val Leu
 145 150 155 160

Gly Phe Pro Gln Glu Val Met Tyr Ser Val Thr Lys Gly Gly Leu Asn
 165 170 175

Gln Met Thr Arg Ser Leu Ala Val Glu Trp Ala Cys Asp Arg Ile Arg
 180 185 190

Val Asn Cys Val Ala Pro Gly Val Ile Met Thr Asp Met Gly Lys Glu
 195 200 205

Leu Pro Ala Ala Leu Val Glu Gln Glu Arg Ser Arg Ile Pro Leu Arg
 210 215 220

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

Met Pro Ala Ala Ser Tyr Val Thr Gly Gln Val Ile Phe Val Asp Gly
 245 250 255

Gly Arg Thr Ile Ser Gly Ala
 260

- <210> 47
- <211> 1259
- <212> DNA
- <213> Oryza sativa

<400> 47
 ataacgccgt atactgggga cgagccagga acccgaattg ggagtttttc ccgaagaact 60
 gataatacgg attttctcgc aaaattattc ccagccgcac cctcgaggcc gatggccgag 120
 gcggtcgcaa gcggcgcagc ggggagatgg accctccgcg gcaagacggc cctcgtcacc 180
 ggcggcacc cgggcatcgg gcatgcgggtg gtggacgagc tggcggcgct gggggcggcc 240
 gtgcacacct gctccaggaa ggaggccgag ctgggcgagc gcctcagggg gtgggagggc 300
 aagggtctcc gcgtcaccgg ctcagtctgc gacgtctccg tgcgggagca gcgggaacgc 360
 atgctccgag aggtcgccgg cctctacggt ggcaagctcg acatcctcgt gaacaacgtg 420
 ggaacaaact tttcaaaaca aaccactgaa tactctgcag atgattactc gttcataatg 480
 gccaccaatc ttgaatctgc gtatcatctg tgccaacttg ctcatcctct tctcaaatca 540
 tctgggtcag gaagcgttgt cttcatatcg tcagtctctg gagtgggtggc tgtaagttct 600
 ggctctgtct atgcgatgac aaaaggtgcc atgaaccagc tggccaagaa cctggcatgt 660
 gagtgggcga aagacaatat aagaaccaat tctgttgacac catggtacat gaagacttca 720
 cttgtggaag atgaattggc aaggaaggac ttcgctgaca gcgtcgtgcg tcgaacagcg 780
 ctgaagcgcg tgggagaacc agaagaggtg tcatcgctgg tcgcttctct gtgcatgccc 840
 ggcgcttctc acatcaccgg ccagacgac tcggctcgagc gcggcatgac cataaacgga 900
 ctgtatccac ctcaggacta gaccgaatag gcaaaccgta tggccgtatg gggatcgaag 960
 aactgtgttt ggagaggtca gaggtgaggg tcgaacgtca gtaaaactca actatcccaa 1020
 acaaacagga gatcgcaaat agaccgctct gaacttgatt ggtttgcaact tgggtatgca 1080
 ccgcatgcta ctgtactaaa ttaaaaaaaaa acgtttggtg cgagcaacaa aacgaaatcc 1140

tgcgtggata gcgggcgctac tgcatagtgt tgtgtcgttc gtatgcgcca tagacaaacg 1200

gaatgccctt gaaaaaaagt ttaattggat gcatcgtgca gcctattaca atttcttcc 1259

<210> 48

<211> 269

<212> PRT

<213> *Oryza sativa*

<400> 48

Met Ala Glu Ala Val Ala Ser Gly Ala Ala Gly Arg Trp Thr Leu Arg
1 5 10 15

Gly Lys Thr Ala Leu Val Thr Gly Gly Thr Arg Gly Ile Gly His Ala
20 25 30

Val Val Asp Glu Leu Ala Ala Leu Gly Ala Ala Val His Thr Cys Ser
35 40 45

Arg Lys Glu Ala Glu Leu Gly Glu Arg Leu Arg Glu Trp Glu Gly Lys
50 55 60

Gly Phe Arg Val Thr Gly Ser Val Cys Asp Val Ser Val Arg Glu Gln
65 70 75 80

Arg Glu Arg Met Leu Arg Glu Val Ala Gly Leu Tyr Gly Gly Lys Leu
85 90 95

Asp Ile Leu Val Asn Asn Val Gly Thr Asn Phe Ser Lys Gln Thr Thr
100 105 110

Glu Tyr Ser Ala Asp Asp Tyr Ser Phe Ile Met Ala Thr Asn Leu Glu
115 120 125

Ser Ala Tyr His Leu Cys Gln Leu Ala His Pro Leu Leu Lys Ser Ser
130 135 140

Gly Ser Gly Ser Val Val Phe Ile Ser Ser Val Ser Gly Val Val Ala
145 150 155 160

Val Ser Ser Gly Ser Val Tyr Ala Met Thr Lys Gly Ala Met Asn Gln
165 170 175

Leu Ala Lys Asn Leu Ala Cys Glu Trp Ala Lys Asp Asn Ile Arg Thr
180 185 190

Asn Ser Val Ala Pro Trp Tyr Met Lys Thr Ser Leu Val Glu Asp Glu
195 200 205

Leu Ala Arg Lys Asp Phe Ala Asp Ser Val Val Arg Arg Thr Ala Leu
 210 215 220

Lys Arg Val Gly Glu Pro Glu Glu Val Ser Ser Leu Val Ala Phe Leu
 225 230 235 240

Cys Met Pro Gly Ala Ser Tyr Ile Thr Gly Gln Thr Ile Ser Val Asp
 245 250 255

Gly Gly Met Thr Ile Asn Gly Leu Tyr Pro Pro Gln Asp
 260 265

<210> 49
 <211> 804
 <212> DNA
 <213> Nostoc sp. PCC 7120

<400> 49
 gtggaaaatt ttgttacccc tccagctttt ggggaaaaaa tccgggaacg ctggacatta 60
 gccggacgaa aagctttgat taccggagct acaaaaggaa ttggtttagc gatcgctcaa 120
 gagttccttg ctttaggtgc agaagtcgtc attgtggctc gcaatgctga agcgattgag 180
 cagcagatga aggcttggca ttctgcgggg aaagttcacg gagttgcagc cgacgtttcc 240
 acttctgagg gtcgtcaa at gatgcttgat tacgtttagca agacctttgg agaacttgac 300
 attttggtga acaacgttgg cacaaatatt cgtaagaaag cgactgatta cacagaagaa 360
 gaatttgotg cgatatttca aattaatcta acttctatat ttgagctttc ccgactgttt 420
 tatcccttac tcaaaacaag taaaacagc agtatcggtta acatcgggtc tgttgctgga 480
 ttaatttccg ttcggactgg cgcaccctat ggcacgacca aagctgcact cgtgcagtta 540
 acgcgatcgc tggcggtaga atgggcagat gatggtatc gcgttaatgc gatcgcacct 600
 tggtttatcc aaactcctct gaccgagcct ctactcaaca atcccgaac ttaagcgca 660
 gtcctttcac gcacaccaat gaaacgcgtg ggtcaaccgg aagaagtagc cagcctgacg 720
 gcttttctct gtatgcctac cgcacccctac atcacaggac aatgtattgc tgttgatgga 780
 ggatttctgg catttggttt ttaa 804

<210> 50
 <211> 267
 <212> PRT
 <213> Nostoc sp. PCC 7120

<400> 50

Met Glu Asn Phe Val Thr Pro Pro Ala Phe Gly Glu Lys Ile Arg Glu
 1 5 10 15

Arg Trp Thr Leu Ala Gly Arg Lys Ala Leu Ile Thr Gly Ala Thr Lys
 20 25 30

Gly Ile Gly Leu Ala Ile Ala Gln Glu Phe Leu Ala Leu Gly Ala Glu
 35 40 45

Val Val Ile Val Ala Arg Asn Ala Glu Ala Ile Glu Gln Gln Met Lys
 50 55 60

Ala Trp His Ser Ala Gly Lys Val His Gly Val Ala Ala Asp Val Ser
 65 70 75 80

Thr Ser Glu Gly Arg Gln Met Met Leu Asp Tyr Val Ser Lys Thr Phe
 85 90 95

Gly Glu Leu Asp Ile Leu Val Asn Asn Val Gly Thr Asn Ile Arg Lys
 100 105 110

Lys Ala Thr Asp Tyr Thr Glu Glu Glu Phe Ala Ala Ile Phe Gln Ile
 115 120 125

Asn Leu Thr Ser Ile Phe Glu Leu Ser Arg Leu Phe Tyr Pro Leu Leu
 130 135 140

Lys Thr Ser Lys Asn Ser Ser Ile Val Asn Ile Gly Ser Val Ala Gly
 145 150 155 160

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

Leu Val Gln Leu Thr Arg Ser Leu Ala Val Glu Trp Ala Asp Asp Gly
 180 185 190

Ile Arg Val Asn Ala Ile Ala Pro Trp Phe Ile Gln Thr Pro Leu Thr
 195 200 205

Glu Pro Leu Leu Asn Asn Pro Glu Thr Leu Ser Ala Val Leu Ser Arg
 210 215 220

Thr Pro Met Lys Arg Val Gly Gln Pro Glu Glu Val Ala Ser Leu Thr
 225 230 235 240

Ala Phe Leu Cys Met Pro Thr Ala Ser Tyr Ile Thr Gly Gln Cys Ile
 245 250 255

Ala Val Asp Gly Gly Phe Leu Ala Phe Gly Phe
 260 265

<210> 51
 <211> 777
 <212> DNA
 <213> Xanthomonas campestris

<400> 51
 gtgacagcgc accgctggcg gctggatgga cagaccgccc tgatcaccgg cgccagtgcc 60
 ggcaattggcc tggccatcgc acgcgaattg ctcggttcg gcgccgacct gctgatggtg 120
 gcgcgcgatg ccgatgcgct ggcgcaggcc cgcgacgagc tggccgaaga gtttcccag 180
 cgcgaactgc acggcctggc cgcggatggt tccgacgacg aggagcgccg cgcgattctg 240
 gattgggtcg aagatcacgc cgatggctta cacctgctga tcaacaatgc cggtggaac 300
 atcaccgcg cgccatcga ctacaccgaa gacgaatggc gcggcatctt cgaaaccaac 360
 gtgttttccg cgttcgaact ctcgcgctat gcgcacccat tgctgaccgg gcacgccc 420
 tcggcgatcg tcaacgtggg cagcgtgtcg gggatcacgc atgtgcgcag cggcgcgccc 480
 tatggcatga ccaaggccgc gctgcagcag atgacgcgca acctggcggt ggaatgggcc 540
 gaagacggca tccgcgtcaa cgcggtggcg ccgtgtgaca tccgcacccg gcgcacctcc 600
 ggcccgttgt ccgatccgga ctactatgag caggtgatcg aacgcacgcc gatgcgccgc 660
 atcggcgagc cggaagaagt cgccgccgca gtgggtttcc tgtgcctgcc ggcggccagc 720
 tacatcaccg gtgaatgcat cgcggtggat ggcggcttcc tgcgtcatgg gttctag 777

<210> 52
 <211> 258
 <212> PRT
 <213> Xanthomonas campestris

<400> 52

Met Thr Ala His Arg Trp Arg Leu Asp Gly Gln Thr Ala Leu Ile Thr
 1 5 10 15
 Gly Ala Ser Ala Gly Ile Gly Leu Ala Ile Ala Arg Glu Leu Leu Gly
 20 25 30
 Phe Gly Ala Asp Leu Leu Met Val Ala Arg Asp Ala Asp Ala Leu Ala
 35 40 45
 Gln Ala Arg Asp Glu Leu Ala Glu Glu Phe Pro Glu Arg Glu Leu His
 50 55 60
 Gly Leu Ala Ala Asp Val Ser Asp Asp Glu Glu Arg Arg Ala Ile Leu
 65 70 75 80
 Asp Trp Val Glu Asp His Ala Asp Gly Leu His Leu Leu Ile Asn Asn

	85		90		95
Ala Gly Gly Asn Ile Thr Arg Ala Ala Ile Asp Tyr Thr Glu Asp Glu	100		105		110
Trp Arg Gly Ile Phe Glu Thr Asn Val Phe Ser Ala Phe Glu Leu Ser	115		120		125
Arg Tyr Ala His Pro Leu Leu Thr Arg His Ala Ala Ser Ala Ile Val	130		135		140
Asn Val Gly Ser Val Ser Gly Ile Thr His Val Arg Ser Gly Ala Pro	145		150		155
Tyr Gly Met Thr Lys Ala Ala Leu Gln Gln Met Thr Arg Asn Leu Ala	165		170		175
Val Glu Trp Ala Glu Asp Gly Ile Arg Val Asn Ala Val Ala Pro Trp	180		185		190
Tyr Ile Arg Thr Arg Arg Thr Ser Gly Pro Leu Ser Asp Pro Asp Tyr	195		200		205
Tyr Glu Gln Val Ile Glu Arg Thr Pro Met Arg Arg Ile Gly Glu Pro	210		215		220
Glu Glu Val Ala Ala Ala Val Gly Phe Leu Cys Leu Pro Ala Ala Ser	225		230		235
Tyr Ile Thr Gly Glu Cys Ile Ala Val Asp Gly Gly Phe Leu Arg His	245		250		255

Gly Phe

<210> 53

<211> 777

<212> DNA

<213> Xylella fastidiosa

<400> 53

atgacccaat cccggttgccg tttggatggc cgtaaggcctt tgatcacggg ggctagtaca 60

ggatcgggtc tggctggttc tctgagttg ttgggcctgg gtgccgatgt gctggttgta 120

gcgcgtaatg ctgatttgct ggggcgagta cgcgatgaac tggcagagga attcctgag 180

cgtgagttac atggattggc tgotgatggt gctgacgatg tggaccgccg tgcgattctg 240

gattgggtcg aagattgcag taatggtttg catgtgttga tcaacaatgc tgggtgtaat 300

gtgacgcgtg cagcgccttga atatacagaa gacgagtggc gtgagatfff cgagatcaat 360
 ctgttctctg cgttcagatt gtgtcgttac acgcagccgt tgctggcgtg tcatgcaagt 420
 acggcgattg tcaatattgg cagtgtatct ggcttgacgc atgtgcgcag tggagtgccg 480
 tatgggatga gcaaggcggc gttgcaccaa atgacccgca atttggctgt ggaatgggct 540
 gaggatggga tccgtgtcaa tgcggtagcg ccttgggtata tccgtacgcg tcgtacgtcg 600
 gagccgctct ctgatgtgta ctattacgag caagtcattg aacgtacgcc gatgcgccgc 660
 attggagagc cggaggaagt agctgcggca gtggcatttc tttgtttgcc agcgtccagt 720
 tatgtgaccg gcgaatgtat tgcggtggat ggcggtttta tgcgctatgg tttctga 777

<210> 54
 <211> 258
 <212> PRT
 <213> Xylella fastidiosa

<400> 54

Met Thr Gln Ser Arg Trp Arg Leu Asp Gly Arg Thr Ala Leu Ile Thr
 1 5 10 15
 Gly Ala Ser Thr Gly Ile Gly Leu Ala Val Ala Arg Glu Leu Leu Gly
 20 25 30
 Leu Gly Ala Asp Val Leu Leu Val Ala Arg Asn Ala Asp Leu Leu Gly
 35 40 45
 Arg Val Arg Asp Glu Leu Ala Glu Glu Phe Pro Glu Arg Glu Leu His
 50 55 60
 Gly Leu Ala Ala Asp Val Ala Asp Asp Val Asp Arg Arg Ala Ile Leu
 65 70 75 80
 Asp Trp Val Glu Asp Cys Ser Asn Gly Leu His Val Leu Ile Asn Asn
 85 90 95
 Ala Gly Gly Asn Val Thr Arg Ala Ala Leu Glu Tyr Thr Glu Asp Glu
 100 105 110
 Trp Arg Glu Ile Phe Glu Ile Asn Leu Phe Ser Ala Phe Glu Leu Cys
 115 120 125
 Arg Tyr Thr Gln Pro Leu Leu Ala Cys His Ala Ser Thr Ala Ile Val
 130 135 140
 Asn Ile Gly Ser Val Ser Gly Leu Thr His Val Arg Ser Gly Val Pro
 145 150 155 160

Tyr Gly Met Ser Lys Ala Ala Leu His Gln Met Thr Arg Asn Leu Ala
 165 170 175

Val Glu Trp Ala Glu Asp Gly Ile Arg Val Asn Ala Val Ala Pro Trp
 180 185 190

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

Tyr Glu Gln Val Ile Glu Arg Thr Pro Met Arg Arg Ile Gly Glu Pro
 210 215 220

Glu Glu Val Ala Ala Ala Val Ala Phe Leu Cys Leu Pro Ala Ser Ser
 225 230 235 240

Tyr Val Thr Gly Glu Cys Ile Ala Val Asp Gly Gly Phe Met Arg Tyr
 245 250 255

Gly Phe

<210> 55
 <211> 25
 <212> DNA
 <213> Arabidopsis thaliana

<400> 55
 atattaaca agccatggca aagga 25

<210> 56
 <211> 27
 <212> DNA
 <213> Arabidopsis thaliana

<400> 56
 atatgtgitt gaattcatag tcttgaa 27

<210> 57
 <211> 27
 <212> DNA
 <213> Arabidopsis thaliana

<400> 57
 gtaagatatc atatggcaaa ggaaggg 27

<210> 58
 <211> 36
 <212> DNA
 <213> arabidopsis thaliana

<400> 58
 gcgcgaattc tcgagtcata gtcttgaagg aaaaac 36

<210> 59
<211> 21
<212> DNA
<213> arabidopsis thaliana

<400> 59
ggctacaaat ctcgagtcag c 21

<210> 60
<211> 21
<212> DNA
<213> arabidopsis thaliana

<400> 60
gctgactcga gatttgtagc c 21

<210> 61
<211> 31
<212> DNA
<213> Nostoc sp. PCC 7120

<400> 61
gtaagatatc atatgagtga tgcctacggt g 31

<210> 62
<211> 34
<212> DNA
<213> Nostoc sp. PCC 7120

<400> 62
gtaagatctg ccacocatggg tgatgcctac ggtg 34

<210> 63
<211> 31
<212> DNA
<213> Nostoc sp. PCC 7120

<400> 63
gagaagcttc tcgagttaaa aaccgaaagc c 31

<210> 64
<211> 19
<212> DNA
<213> Nostoc sp. PCC 7120

<400> 64
cctctactgc aaaataggg 19

/

<210> 65
<211> 19
<212> DNA
<213> Nostoc sp. PCC 7120

<400> 65
ccctatthttg cagtagagg 19

<210> 66
 <211> 901
 <212> DNA
 <213> Nostoc sp.

<400> 66
 cattgggcac ttacaagata aactcttaaa ttgatacctt gaaaataaaa aatccgcaaa 60
 attacaaatg agtgatecct acggtgggct aagtctacgc tggacactga gaggaagaa 120
 agcactgatt acaggtgcta ccaaaggcat tggactagct gttgccaatg agttcttattc 180
 tctaggtgca gaagtcataa ttgtggcacg gaattctcaa gatgttgacc aacaactaat 240
 tatctggcga gaattaggat tacctgctta tggaattacg gcagatgttg caactgctga 300
 ggggcgacaa gccatttttg agcaggttgg caaacctgg gataaattag acatcctggt 360
 taataatgtg ggaaccaata tcagcaaaaa agtactagac tacacggctg ctgaatatca 420
 gttcatcatc cagacgaacc agatatcaat ttttgagatg tgccgtctgt tttaccctct 480
 actgcaaaat agggaaaata gcagcattgt gaatataagt tctgtggcgg gtttagtctc 540
 aaaccgtaca ggcgctcctt atggtatgac taaagcggct ataaatcaat tgacgcgatc 600
 gctatcagtt gaatgggctt gcgatcaaat tagggtaaata accgtagcac cttgggctat 660
 tcgcactcct ctaaccgagt ctgtactcga taaccaagat tttctcaaat tagttctgtc 720
 ccaaacacct atgggaagag ttggtcaacc agaagaagta gcgggttag tagcatttct 780
 ttgtatgcct gcggcatcct tcatcactgg acaatgtatc actgtcgatg gtggtttttt 840
 ggctttoggt ttttaagtct caaagccaga tccatccacg aggtagaaga aactacttac 900
 a 901

<210> 67
 <211> 262
 <212> PRT
 <213> Nostoc sp.

<400> 67
 Met Ser Asp Ala Tyr Gly Gly Leu Ser Leu Arg Trp Thr Leu Arg Gly
 1 5 10 15
 Lys Lys Ala Leu Ile Thr Gly Ala Thr Lys Gly Ile Gly Leu Ala Val
 20 25 30
 Ala Asn Glu Phe Leu Ser Leu Gly Ala Glu Val Ile Ile Val Ala Arg
 35 40 45
 Asn Ser Gln Asp Val Asp Gln Gln Leu Ile Ile Trp Arg Glu Leu Gly
 50 55 60

Leu Pro Ala Tyr Gly Ile Thr Ala Asp Val Ala Thr Ala Glu Gly Arg
65 70 75 80

Gln Ala Ile Phe Glu Gln Val Gly Lys Thr Trp Asp Lys Leu Asp Ile
85 90 95

Leu Val Asn Asn Val Gly Thr Asn Ile Ser Lys Lys Val Leu Asp Tyr
100 105 110

Thr Ala Ala Glu Tyr Gln Phe Ile Ile Gln Thr Asn Gln Ile Ser Ile
115 120 125

Phe Glu Met Cys Arg Leu Phe Tyr Pro Leu Leu Gln Asn Arg Glu Asn
130 135 140

Ser Ser Ile Val Asn Ile Ser Ser Val Ala Gly Leu Val Ser Asn Arg
145 150 155 160

Thr Gly Ala Pro Tyr Gly Met Thr Lys Ala Ala Ile Asn Gln Leu Thr
165 170 175

Arg Ser Leu Ser Val Glu Trp Ala Cys Asp Gln Ile Arg Val Asn Thr
180 185 190

Val Ala Pro Trp Ala Ile Arg Thr Pro Leu Thr Glu Ser Val Leu Asp
195 200 205

Asn Gln Asp Phe Leu Lys Leu Val Leu Ser Gln Thr Pro Met Gly Arg
210 215 220

Val Gly Gln Pro Glu Glu Val Ala Gly Leu Val Ala Phe Leu Cys Met
225 230 235 240

Pro Ala Ala Ser Phe Ile Thr Gly Gln Cys Ile Thr Val Asp Gly Gly
245 250 255

Phe Leu Ala Phe Gly Phe
260

<210> 68

<211> 810

<212> DNA

<213> Nostoc sp.

<400> 68

tcatacatgg ccaactgccag ctctataaac aagaggttta gcctcgccgg tgccacggcg 60

cttgtcaccg gtggcagcaa aggcacgcgc cgcgccatcg tggaagagct ggcgagcttc 120

ggcgcgacgg tgcacacctg cgctcggaac caggcggagc tgagcagatg ccaggaggag 180
 tggacggcca agggcctcgc cgtcaccgtc tccgtctgcg acgtggcggg gcgcgccgac 240
 agggaggcgc tcgccggcag ggtgtccgcc atgttcgacg gcaagctcag catcctgggtg 300
 aacaacgctg ggacggcgta cctgaagccg gcggcggacc tgacgccgga ggagacgtcg 360
 aggttgatga cgaccaactt cgagtcgtgc ttccacctga gccagctggt ctaccctctc 420
 ctcaaggact ccggaagagg cagtatcgtt aatatctcgt ccgtcgtctt tgtcctcgcg 480
 ttccattctc ttcctatcta ctcggctgcc aaaggagcaa tgaaccaagt cacaaggaac 540
 ctggcttgtg agtgggagcag cgatgggatc agagtcaatt ccgttgacc gggctacatc 600
 cagactccac tcctaactgc cttcgtggcc ggcaacgatt tcgcacaggt tgagttcaac 660
 cgtcttcctt tgggccgtct cggtaaacct gaggacatct cgtcgttggg ggcgttcctg 720
 tgcatgcctg cggcctccta catcactggc cagattatat gcgtcgatgg gggccgatg 780
 ctttcttaat aaactctaca tatacaaaat 810

<210> 69
 <211> 260
 <212> PRT
 <213> Nostoc sp.

<400> 69

Met Ala Thr Ala Ser Ser Ile Asn Lys Arg Phe Ser Leu Ala Gly Ala
 1 5 10 15

Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Arg Ala Ile Val
 20 25 30

Glu Glu Leu Ala Ser Phe Gly Ala Thr Val His Thr Cys Ala Arg Asn
 35 40 45

Gln Ala Glu Leu Ser Arg Cys Gln Glu Glu Trp Thr Ala Lys Gly Leu
 50 55 60

Ala Val Thr Val Ser Val Cys Asp Val Ala Val Arg Ala Asp Arg Glu
 65 70 75 80

Ala Leu Ala Gly Arg Val Ser Ala Met Phe Asp Gly Lys Leu Ser Ile
 85 90 95

Leu Val Asn Asn Ala Gly Thr Ala Tyr Leu Lys Pro Ala Ala Asp Leu
 100 105 110

Thr Pro Glu Glu Thr Ser Arg Leu Met Thr Thr Asn Phe Glu Ser Cys
 115 120 125

Phe His Leu Ser Gln Leu Phe Tyr Pro Leu Leu Lys Asp Ser Gly Arg
 130 135 140

Gly Ser Ile Val Asn Ile Ser Ser Val Ala Ser Val Leu Ala Phe His
 145 150 155 160

Ser Leu Pro Ile Tyr Ser Ala Ala Lys Gly Ala Met Asn Gln Val Thr
 165 170 175

Arg Asn Leu Ala Cys Glu Trp Ala Ser Asp Gly Ile Arg Val Asn Ser
 180 185 190

Val Ala Pro Gly Tyr Ile Gln Thr Pro Leu Leu Thr Ala Phe Val Ala
 195 200 205

Gly Asn Asp Phe Ala Gln Val Glu Phe Asn Arg Leu Pro Leu Gly Arg
 210 215 220

Leu Gly Lys Pro Glu Asp Ile Ser Ser Leu Val Ala Phe Leu Cys Met
 225 230 235 240

Pro Ala Ala Ser Tyr Ile Thr Gly Gln Ile Ile Cys Val Asp Gly Gly
 245 250 255

Arg Met Leu Ser
 260

<210> 70
 <211> 1253
 <212> DNA
 <213> Glycine max

<400> 70
 gaggaagtat ccgatatttg acaggttcat catcataacc ttcaatgaaa agaaatggct 60
 gaggaagca ttggcagcaa aagcagcaga tggcttttac agggaatgac agctctcgtc 120
 accggtggat ccaaaggaat cggatatgct atcgtggagg agttggcaca gcttggagcc 180
 actgtgcaca cttgcgctcg gaacgaagct gaactcaatg aatccttaaa tgaatggaac 240
 acaaaaggat acagagtaac tggttccgtc tgtgacgtgg cgtctcgtgc agaaagacaa 300
 gacctcatag ctagagtctc caatgagttt aatggcaaac tcaatcctt tgtaaacaaac 360
 gtgggaacaa acgtaccgaa acataccctt gatgttacgg aggaagactt ctcatctctg 420
 ataaatacaa atcttgaatc tgcttaccac ctaagccagc ttgcacatcc tctcctgaaa 480
 gcttcagagg ctgcaaacat catttttata tcctccattg ctggtgtgct atcaataggt 540
 ataggatcca cttatggtgc aacaaaagga gcaatgaacc aactgactaa aaatttggca 600

tgtgagtggg ccaaagacaa tataaggact aattgcgttg caccagggcc aatcaaaacc 660
 cctctcgggtg acaagcattt taaaaatgaa aaacttctta atgctttcat ttcgcaaacc 720
 cccttggac ggattggaga agcagaggaa gtgtcttcat tgggtggcatt cctctgctta 780
 cctgcagcct cttacataac aggacagacc atttgtgttg atggtggatt aacagtgaat 840
 ggtctctata taaattagag aagtactacc aacacactct ctaacacact tctcttgaca 900
 catattctac cataagttac aaataataaa aaattacaaa atcatgaaaa taagatctag 960
 gacccatata tttttgtgat atctaaaaaa tttcaagcaa taaaaaaaaat gtattgaaaa 1020
 gagtgataaa gataatgtgt tgtttagcatt tttatataaa ctatgttaag agcaaggctc 1080
 gatgcatacg ttgctagttt ttatattagt attgagaata aatagctgaa agtcccttgg 1140
 ttcaatctat atcatttggg agggggaatt tgttgggaat aaatttgtat gtccacgatc 1200
 caatcatcat gtaatcaatt ttaattttaa taataaagta ttattttatt ttc 1253

<210> 71
 <211> 267
 <212> PRT
 <213> Glycine max

<400> 71

Met Ala Glu Ala Ser Ile Gly Ser Lys Ser Ser Arg Trp Ser Leu Gln
 1 5 10 15

Gly Met Thr Ala Leu Val Thr Gly Gly Ser Lys Gly Ile Gly Tyr Ala
 20 25 30

Ile Val Glu Glu Leu Ala Gln Leu Gly Ala Thr Val His Thr Cys Ala
 35 40 45

Arg Asn Glu Ala Glu Leu Asn Glu Ser Leu Asn Glu Trp Asn Thr Lys
 50 55 60

Gly Tyr Arg Val Thr Gly Ser Val Cys Asp Val Ala Ser Arg Ala Glu
 65 70 75 80

Arg Gln Asp Leu Ile Ala Arg Val Ser Asn Glu Phe Asn Gly Lys Leu
 85 90 95

Asn Ile Leu Val Asn Asn Val Gly Thr Asn Val Pro Lys His Thr Leu
 100 105 110

Asp Val Thr Glu Glu Asp Phe Ser Phe Leu Ile Asn Thr Asn Leu Glu
 115 120 125

Ser Ala Tyr His Leu Ser Gln Leu Ala His Pro Leu Leu Lys Ala Ser

130
Glu Ala Ala Asn Ile Ile Phe Ile Ser Ser Ile Ala Gly Val Leu Ser
145 150 155 160

Ile Gly Ile Gly Ser Thr Tyr Gly Ala Thr Lys Gly Ala Met Asn Gln
165 170 175

Leu Thr Lys Asn Leu Ala Cys Glu Trp Ala Lys Asp Asn Ile Arg Thr
180 185 190

Asn Cys Val Ala Pro Gly Pro Ile Lys Thr Pro Leu Gly Asp Lys His
195 200 205

Phe Lys Asn Glu Lys Leu Leu Asn Ala Phe Ile Ser Gln Thr Pro Leu
210 215 220

Gly Arg Ile Gly Glu Ala Glu Glu Val Ser Ser Leu Val Ala Phe Leu
225 230 235 240

Cys Leu Pro Ala Ala Ser Tyr Ile Thr Gly Gln Thr Ile Cys Val Asp
245 250 255

Gly Gly Leu Thr Val Asn Gly Leu Tyr Ile Asn
260 265