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(54) Title: PLANT POLYNUCLEOTIDES ENCODING NOVEL Na⁺/H⁺ ANTIPORTERS

(57) Abstract: The present invention provides novel polynucleotides encoding plant Na⁺/H⁺ antiporter polypeptides, fragments and homologs thereof. Also provided are vectors, host cells, antibodies, and recombinant methods for producing said polypeptides. The invention further relates to methods of applying these novel plants polypeptides to the identification, prevention, and/or conferment of resistance to various plant diseases and/or disorders, particularly those associated with modulating environmental stress responses, such as drought and salt tolerance.

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**PLANT POLYNUCLEOTIDES ENCODING NOVEL
Na⁺/H⁺ ANTIPORTERS**

FIELD OF THE INVENTION

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The present invention provides novel polynucleotides encoding plant Na⁺/H⁺ antiporter polypeptides, fragments and homologs thereof. Also provided are vectors, host cells, antibodies, and recombinant methods for producing said polypeptides. The invention further relates to methods of applying these novel plant polypeptides to the identification, prevention, and/or conferment of resistance to various plant diseases and/or disorders, particularly those associated with modulating environmental stress responses, such as drought and salt tolerance.

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BACKGROUND OF THE INVENTION

Environmental stress due to salinity is one of the most serious factors limiting the productivity of agricultural crops, which are predominantly sensitive to the presence of high concentrations of salts in soil. It is estimated that 35-45% of the 279 million hectares of land irrigation is presently affected by salinity. This is exclusive of the regions classified as arid and desert lands. In this century, more areas including vast regions of Australia, Europe, Southwest USA, the Canadian prairies and others have seen considerable declines in crop productivity due to salinity in lands. The consequence represents a significant economic and political factor and contributes to food shortages in many undeveloped countries.

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Although there is engineering technology available to overcome this problem, though drainage and supply of high quality water, these measures are extremely costly. In most of the cases, due to the increased need for extensive agriculture, neither improved irrigation efficiency nor the installation of drainage systems is applicable. Moreover, in the arid and semi-arid regions of the world water evaporation exceeds precipitation. These soils are inherently high in salt and require vast amounts of irrigation to become productive. Since irrigation water contains dissolved salts and minerals, an application of water is also an application of salt that compounds the salinity problem.

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5 Conventional breeding strategies for salt tolerance have been attempted for a long time. These breeding practices have been based mainly on the following strategies: a) the use of wide crosses between crop plants and their more salt-tolerant wild relatives (Rush, PW and Epstein, E, 1981 J.Amer. Soc. Hort. Sci. 106, 699-704) (b) screening and selecting for variation within a particular phenotype (Norlyn, JD. 10 1980. In: Genetic Engineering of Osmoregulation (Eds. DW Rains, RC Valentine and A Hollaender) pp.293-309. Plenum press: New York.), c) designing new phenotypes through recurrent selection (Tal, M. 1985. Plant and Soil 89, 199-226). The lack of success in generating tolerant varieties would suggest that conventional breeding practices are not enough and that in order to succeed a breeding program should 15 include the engineering of transgenic crops which allows one to generate salt stress-tolerant crops (Flowers TJ and Yeo, AR, 1995. Aust. J. Plant. Physiol. 22., 875-884., Bohnert HJ and Jensen, RG. 1996. Aust. J. Plant. Physiol. 23., 661-667.)

Plant cells are structurally well suited to the compartmentation of ions. Large membrane-bound vacuoles are the site for a considerable amount of sequestration of 20 ions and other osmotically active substances. Transport mechanisms could actively move ions into the vacuole, removing the potentially harmful ions from cytosols. Thus, at the cellular level both specific transport systems for sodium accumulation in the vacuole and sodium extrusion out of the cell are correlated with salt tolerance.

Several sodium transport systems associated with salt tolerance have been 25 characterized in different organisms and a few of the genes involved in this process have been identified and in some cases the predicted role of protein has been investigated in transgenic/overexpression experiments. A single gene (*sod2*) coding for a Na^+/H^+ antiport has been shown to confer sodium tolerance in fission yeast (Jia, ZP, et al. 1992 EMBO J. 11, 1631-1640., Young, PG and Zheng, PJ. 30 Patent#WO9106651), although the role of this plasma membrane-bound protein appears to be only limited to yeast. One of the main disadvantages of using this gene for transformation of plants is associated with the typical problems encountered in heterologous gene expression. Two homologues of sodium antiporter, *AtNhx1* and *SOS1* from salt-sensitive plants, *Arabidopsis thaliana* have been identified and 35 characterized (Apse, MP et al., 1999, Science 285, 1256-1258., Shi, H et al., 2000 Proc. Natl. Aca. Sci.USA 97, 6896-6901). Overexpression of *AtNhx1* in *Arabidopsis* as well as in fission yeast shows increased salt tolerance due to better performance of salt compartmentation into the vacuole (Gaxiola, RA., et al. 1999. Proc. Natl. Acad.

5 Sci. USA. 96, 1480-1485., Apse, MP et al., 1999, Science 285, 1256-1258.,).
However, a comparison of ion distribution in cells and tissues of various plant species
indicates that a primary characteristic of salt-tolerant plants is their ability to exclude
sodium out of the cell and to take up sodium and sequester it in the cell vacuoles (Niu,
X., et al., 1995 Plant Physiol. 109, 735-742). This strongly suggests that Na⁺/H⁺
10 antiporter from salt-tolerant plants have functionally more effective sodium transport
systems compared with salt-sensitive plants such as Arabidopsis.

Therefore, elucidating the function of sodium transport genes in salt tolerant
plants will not only advance our understanding of plant adaptation and tolerance to
salinity stress, but also may provide important information for designing new
15 strategies for crop improvement. Newly generated salt tolerant plants will have many
advantages, such as increasing the range that crop plants can be cultivated in salinity
lands. This invention fulfills this need by providing the sequences of plant Na⁺/H⁺
antiporter genes that are expressed in the halophyte *Physcomitrella patens*, which can
therefore provide a basis of increasing the salt tolerance of non-halophytic plants.

20 Moreover, the present invention provides novel polynucleotides encoding
plant Na⁺/H⁺ antiporter polypeptides, fragments and homologs thereof. Also provided
are vectors, host cells, antibodies, and recombinant methods for producing said
polypeptides. The invention further relates to methods of applying these novel plant
polypeptides to the identification, prevention, and/or conferment of resistance to
25 various plant diseases and/or disorders, particularly environmental stress tolerance in
plants.

Due to the commercial consequences of environmental damage to crops, there
is an interest in understanding how to improve a plant's response to environmental
damage. By improving a plant's performance or survival in response to cold, drought
30 and salinity the environment stress-related risks of farming can be reduced. This
invention fulfills in part the need to identify new Na⁺/H⁺ antiporters capable of
conferring drought, freezing and salt tolerance to plants upon over-expression.
Namely, we describe Na⁺/H⁺ antiporters (PpNHX1 and PpNHX2) from
Physcomitrella patens.

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BRIEF SUMMARY OF THE INVENTION

This invention provides novel nucleic acid molecules encoding for either fully active polypeptides or portions thereof from the enzymes Na^+/H^+ antiporter from *Physcomitrella patens*.

10 The present invention further provides a general method for engineering salt-tolerant plants. Moreover, this method should be applicable to all plants of interest. Moreover, this method should be applicable to other environmental stresses.

The present invention describes the introduction of a Na^+/H^+ antiporter activity from the halophyte *Physcomitrella patens* into plants. This method should be very well suited to enhance salt tolerance as the antiporter from a halophyte is more
15 efficient to protect the cell from salt stress than is the antiporter from non-halophyte sources.

Further described in this invention are vectors used to transform Arabidopsis, Rapeseed, soybeans and corn plants.

In a preferred aspect of this invention, the engineering of salt-tolerant plants is
20 described.

Moreover, the invention provides methods of applying the polynucleotides and polypeptides of the invention for creating transgenic plants with desirable traits, which include, but are not limited, to enhanced plant defense, drought tolerance, salt tolerance, ultraviolet(uv) tolerance, enhanced flower development and terpene
25 synthesis. The invention provides antibodies specific to polypeptides of the present invention. Further, the invention provides methods of using antibodies of the present invention to modulate plant phenotype, both functionally and morphologically. The invention also provides methods for more particularly refining the function of the polynucleotides and/or polypeptides of the present invention.

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BRIEF DESCRIPTION OF THE FIGURES/DRAWINGS

Figure 1 - Diagram of the plant expression vector pGMSG containing the plant promoter controlling the expression of the *Physcomitrella* Na^+/H^+ Antiporter (PpNHX1 and PpNHX2). Diagram of the plant expression vector pGMSG containing
35 the plant promoter controlling the expression of the *Physcomitrella* Na^+/H^+ Antiporter gene.

The components are: aacCI gentamycin resistance gene (Hajdukiewicz et al., 1994 (Hajdukiweicz, P., Svab, Z. and Maliga, P. (1994). The small, versatile pPZP family

5 of *Agrobacterium* binary vectors for plant transformation. *Plant Molecular Biology* 25, 989-994)), nosP NOS promoter (Becker et al., 1992 (Becker, D., E. Kemper, J.Schell, and R. Masterson. 1992. New plant binary vectors with selectable markers located proximal to the left T-DNA border. *Plant Mol. Biol.* 20(6): 1195-1197)), g7T terminator (Becker et al., 1992), NOSpA terminator (Jefferson et al., 1987 (Jefferson, R.A., T.A. Kavanagh, and M.W. Bevan. 1987. GUS fusions: β -Glucuronidase as a sensitive and versatile gene fusion marker in higher plants. *EMBO J.* 6(13): 3901-7)). In pBPSNVT001 and pBPSNVT003, the expression of the Na^+/H^+ Antiporter (PpNHX1 and PpNHX2) gene is under control of the constitutive super-promoter. (Ni et al., 1995 (Ni, M., Cui, D., Einstein, J., Narasimhulu, S., Vergara, C. E., and Gelvin, S. B. (1995) Strength and tissue specificity of chimeric promoters derived from the octopine and mannopine synthase gene. *The Plant Journal* 7(4): 661-676)).

20 Figures 2A-D - Na^+/H^+ Antiporters. Nucleotide sequence of the full-length PpNHX1 (SEQ ID NO: 1); and the partial and full-length PpNHX2 (SEQ ID NO: 3 and NO: 4, respectively) from *Physcomitrella patens*.

Figure 3 - Deduced amino acid sequence of the full length Na^+/H^+ antiporters NHX1 (SEQ ID NO: 2); and NHX2 (SEQ ID NO: 5) from *Physcomitrella patens*

25 Figures 4A-C - Amino acid multiple sequence alignment (CLUSTAL W algorithm, blosum62 scoring matrix) between the full-length Vacuole-type Na^+/H^+ Antiporter (SEQ ID NO:2) and sequences of *Oryza sativa* (Q9SXJ8), *Caenorhabditis elegans* (Q9X14), *Drosophila melanogaster* (Q9VP1) and *Arabidopsis AtNHX1* (AAD 16946). The amiloride binding site is highlighted.

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Figure 5 - Amino acid pile up comparison (PARAMETERS) of SEQ ID NO: 5 and sequences of *Oryza sativa* (Swiss plot # :Q9SXJ8), *Arabidopsis thaliana* (Swiss plot # :Q9SUQ7, Q9SS27, Q9LUN4).

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Figure 6 - Amino acid sequence comparison (fasta: Pairwise alignment, blosum 62 scoring matrix, gap opening penalty: 10, gap extension penalty: 0.1) of the full-length Na⁺/H⁺ Antiporter (SEQ ID NO:2 and SEQ ID NO: 5), *Oryza sativa* (Q9SXJ8),
10 *Caenorhabditis elegans* (Q9X14), *Drosophila melanogaster* (Q9VP1), and *Arabidopsis AtNHX1* (AAD 16946) and Na⁺/H⁺ exchanging protein-like (BAB02053). The percent identify and percent similarity values are shown in parenthesis.

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DETAILED DESCRIPTION OF THE INVENTION

The present invention may be understood more readily by reference to the following detailed description of the preferred embodiments of the invention and the Examples included herein.

20 One aspect of this invention pertains to isolated nucleic acid molecules that encode fully active polypeptides the PpNHX1 and PpNHX2 Na⁺/H⁺ antiporter from *Physcomitrella patens*. Moreover, this invention pertains to nucleic acid fragments originated from the clones mentioned above, as well as other nucleic acid fragments from other organisms that can be isolated using the described nucleic acid fragments
25 as probes in hybridization experiments.

One embodiment of the present invention relates to increasing stress resistance in a plant by transforming a plant using the polynucleotides and polypeptides, including agonists and/or fragments thereof. This method has herein been demonstrated for *Arabidopsis thaliana*, Rapeseed, soybeans, and corn but its
30 application is not restricted to these plants. The method relies on the increased sodium transport efficiency of the PpNHX1 and PpNHX2 polypeptides as compared to endogenous plant sodium antiporters. The method is most advantageous when the PpNHX1 and PpNHX2 polynucleotides are expressed constitutively in a plant. However, it may be advantageous to have the PpNHX1 and PpNHX2 polynucleotides
35 expressed respectively in specific tissues and/or under the control of an inducible promoter using promoters disclosed herein. Alternatively, the method is also advantageous when the expression of the PpNHX1 and PpNHX2 polynucleotides are

5 under the control of a stress-responsive promoter, such as a drought and/or salt activatable promoter.

The present invention can make a significant contribution to the art by providing new strategies to engineer salt-and drought-tolerance in crop plants, especially the use of the previously unknown PpNHX1 and PpNHX2 clone,
10 respectively. Over-expression of these Na^+/H^+ antiporter (PpNHX1 and PpNHX2) in Arabidopsis confers a high degree of salt tolerance to this plant. It is noteworthy that the analyses of these transgenic lines were performed with T2 plants. The results will be better when a homozygous, strong expression variant is found. The predicted proteins of PpNHX1 and PpNHX2 are homologous to several plant Na^+/H^+ antiporter.
15 The PpNHX1 and PpNHX2 proteins have 27% identity each other at the amino acid level, suggesting that these genes could function differentially as a Na^+/H^+ antiporter (e.g. stress specific or tissue/organ specific).

The present invention also encompasses polynucleotides of the present invention comprising a sodium binding site (amilolide binding site).

20 In another embodiment of the present invention, the polynucleotides and polypeptides of the present invention, in addition to agonists and/or fragments thereof, are useful in modulating the stress tolerance in a plant, preferably modulating a plant's tolerance to the following, non-limiting conditions: limited or inadequate water availability, excess salt or osmotic conditions, excess temperature conditions,
25 excess metal concentrations in soil or water, chemical stress, in addition to oxidative stress. Moreover, the polynucleotides and polypeptides of the present invention are useful in modulating multiple stress responses in a plant, preferably at least one, two, three, four, or more stress responses.

The invention also encompasses the use of the polynucleotides and
30 polypeptides of the present invention, including fragments thereof, in modulating the activity of the sodium antiporter of the present invention, in addition to other sodium antiporters, through protein-protein interactions of the C-terminal regulatory domain of the present invention. Proteins that bind to the C-terminal regulatory domain of the present invention are encompassed by the present invention.

35 The polynucleotides can also be used to express recombinant proteins for analysis, characterization and agronomic use, to express recombinant proteins to raise antibodies directed against polypeptides of the present invention, as markers for tissues in which the corresponding protein is expressed (e.g., preferentially, or non-

5 preferentially), as hybridization markers on Southern gels, as genetic markers for breeding assistance, as RFLP markers, as markers for genotyping (varieties, etc.), and the encoded protein, can, at the very least, be used as a molecular weight marker.

The polynucleotides of the present invention are also useful as chromosome markers or tags (when labeled) to identify chromosomes, to map related gene
10 positions within a chromosome, or as a comparative reference to endogenous DNA sequences of mutant plants to identify allelic variants, and/or spontaneous or biotic mutations.

The polynucleotides of the present invention are also useful for genetic fingerprinting, for selecting and making oligomers for attachment to a “gene chip” or
15 other support, including for examination of expression patterns for particular genes, to differentiate intron and/or exon boundaries, to identify splice and/or allelic variants, and as diagnostic tools for identification of developmental stages, disease states, and/or nutrient levels.

The present invention encompasses polynucleotides that hybridize to the
20 polynucleotides of the present invention under either stringent or non-stringent conditions. Such hybridization may be used to identify orthologs, homologs, allelic variants, variants, and/or mutants of the polynucleotides of the present invention. Additionally, the polynucleotides of the present invention may be used to clone orthologs, homologs, allelic variants, variants, and/or mutants of the polynucleotides
25 of the present by using oligonucleotides directed to polynucleotide sequences of the present invention, and performing PCR on plant cell or tissue samples.

The present invention encompasses the identification of proteins, nucleic acids, or other molecules, that bind to polypeptides and polynucleotides of the present invention (for example, in a receptor-ligand interaction). The polynucleotides of the
30 present invention can also be used in interaction trap assays (such as, for example, that described by Ozenberger and Young (*Mol Endocrinol.*, 9(10):1321-9, (1995); and *Ann N Y Acad Sci.*, 7;766:279-81, (1995)).

Potential uses of polynucleotides and polypeptides of the present invention include nutritional (e.g., as an amino acid supplement), as a carbon source, as nitrogen
35 source, as a carbohydrate source, modulating plant defense activity, modulating signal transduction, modulating metabolite transport (e.g., carbon, nitrogen fluxes, etc.), conferring abiotic stress tolerance and/or resistance (water, drought, cold, salt, etc.),

5 conferring xenobiotic stress tolerance and/or resistance, and development control (for example, yield, flowering time, etc.).

The polynucleotide and polypeptides of the present invention are useful as probes for the identification and isolation of full-length cDNAs and/or genomic DNA which correspond to the polynucleotides of the present invention, as probes to hybridize and
10 discover novel, related DNA sequences, as probes for positional cloning of this or a related sequence, as probe to "subtract-out" known sequences in the process of discovering other novel polynucleotides, as probes to quantify gene expression, and as probes for microarrays.

In addition, polynucleotides and polypeptides of the present invention may
15 comprise one, two, three, four, five, six, seven, eight, or more membrane domains.

Also, in preferred embodiments the present invention provides methods for further refining the biological function of the polynucleotides and/or polypeptides of the present invention.

Specifically, the invention provides methods for using the polynucleotides and
20 polypeptides of the invention to identify orthologs, homologs, variants, and/or allelic variants of the invention. Also provided are methods of using the polynucleotides and polypeptides of the invention to identify the entire coding region of the invention, non-coding regions of the invention, regulatory sequences of the invention, and secreted, mature, pro-, prepro-, forms of the invention.

In preferred embodiments, the invention provides methods for identifying the
25 glycosylation sites inherent in the polynucleotides and polypeptides of the invention, and the subsequent alteration, deletion, and/or addition of said sites for a number of desirable characteristics which include, but are not limited to, augmentation of protein folding, inhibition of protein aggregation, regulation of intracellular trafficking to
30 organelles, increasing resistance to proteolysis, modulation of protein antigenicity, and mediation of intercellular adhesion.

In further preferred embodiments, methods are provided for evolving the
polynucleotides and polypeptides of the present invention using molecular evolution in an effort to create and identify novel variants with desired structural, functional,
35 and/or physical characteristics.

The present invention in further preferred embodiments provides for other experimental methods and procedures currently available to derive functional assignments. These procedures include but are not limited to spotting of clones on

5 arrays, micro-array technology, PCR based methods and other procedures that could use sequence information from the clones to build a primer or a hybrid partner.

As used herein, the term "environmental stress" refers to any sub-optimal growing condition and includes, but is not limited to, sub-optimal conditions associated with salinity, drought, temperature, metal, chemical, pathogenic and
10 oxidative stresses, or combinations thereof. In preferred embodiments, the environmental stress can be salinity, drought, or temperature, or combinations thereof, and in particular, can be high salinity, low water content or low temperature. It is also to be understood that as used in the specification and in the claims, "a" or "an" can mean one or more, depending upon the context in which it is used. Thus, for
15 example, reference to "a cell" can mean that at least one cell can be utilized.

As also used herein, the term "nucleic acid" and "polynucleotide" refer to RNA or DNA that is linear or branched, single or double stranded, or a hybrid thereof. The term also encompasses RNA/DNA hybrids. These terms also encompass untranslated sequence located at both the 3' and 5' ends of the coding region of the gene: at least
20 about 1000 nucleotides of sequence upstream from the 5' end of the coding region and at least about 200 nucleotides of sequence downstream from the 3' end of the coding region of the gene. Less common bases, such as inosine, 5-methylcytosine, 6-methyladenine, hypoxanthine and others can also be used for antisense, dsRNA and ribozyme pairing. For example, polynucleotides that contain C-5 propyne analogues
25 of uridine and cytidine have been shown to bind RNA with high affinity and to be potent antisense inhibitors of gene expression. Other modifications, such as modification to the phosphodiester backbone, or the 2'-hydroxy in the ribose sugar group of the RNA can also be made. The antisense polynucleotides and ribozymes can consist entirely of ribonucleotides, or can contain mixed ribonucleotides and
30 deoxyribonucleotides. The polynucleotides of the invention may be produced by any means, including genomic preparations, cDNA preparations, in vitro synthesis, RT-PCR and *in vitro* or *in vivo* transcription.

An "isolated" nucleic acid molecule is one that is substantially separated from other nucleic acid molecules which are present in the natural source of the nucleic
35 acid (i.e., sequences encoding other polypeptides). Preferably, an "isolated" nucleic acid is free of some of the sequences which naturally flank the nucleic acid (i.e., sequences located at the 5' and 3' ends of the nucleic acid) in its naturally occurring replicon. For example, a cloned nucleic acid is considered isolated. In various

5 embodiments, the isolated NA^+/H^+ ANTIporter nucleic acid molecule can
contain less than about 5 kb, 4 kb, 3 kb, 2 kb, 1 kb, 0.5 kb or 0.1 kb of nucleotide
sequences which naturally flank the nucleic acid molecule in genomic DNA of the
cell from which the nucleic acid is derived. A nucleic acid is also considered isolated
if it has been altered by human intervention, or placed in a locus or location that is not
10 its natural site, or if it is introduced into a cell by agroinfection. Moreover, an
“isolated” nucleic acid molecule, such as a cDNA molecule, can be free from some of
the other cellular material with which it is naturally associated, or culture medium
when produced by recombinant techniques, or chemical precursors or other chemicals
when chemically synthesized.

15 Specifically excluded from the definition of “isolated nucleic acids” are:
naturally-occurring chromosomes (such as chromosome spreads), artificial
chromosome libraries, genomic libraries, and cDNA libraries that exist either as an in
vitro nucleic acid preparations or as a transfected/transformed host cell preparation,
wherein the host cells are either an in vitro heterogeneous preparation or plated as a
20 heterogeneous population of single colonies. Also specifically excluded are the above
libraries wherein a specified nucleic acid makes up less than 5% of the number of
nucleic acid inserts in the vector molecules. Further specifically excluded are whole
cell genomic DNA or whole cell RNA preparations (including whole cell preparations
that are mechanically sheared or enzymatically digested). Even further specifically
25 excluded are the whole cell preparations found as either an in vitro preparation or as a
heterogeneous mixture separated by electrophoresis wherein the nucleic acid of the
invention has not further been separated from the heterologous nucleic acids in the
electrophoresis medium (e.g., further separating by excising a single band from a
heterogeneous band population in an agarose gel or nylon blot).

30

Preferred Polynucleotides and Polypeptides of the Invention

Features of the Polypeptides

35 The polypeptides of genes provided as SEQ ID NO:2 and SEQ ID NO:5 (Figure
3), encoded by the polynucleotide sequence according to SEQ ID NO:1 and SEQ ID
NO:3, and SEQ ID NO:4 (Figures 2A-D), and/or encoded by the polynucleotide
contained within the clones, PpNHX1 and PpNHX2, have significant homology at the
nucleotide and amino acid level to sodium anti-porters in *Oryza sativa* (Q9SXJ8,

5 Q9SXJ8#2), *Caenorhabditis elegans* (Q9X14), *Drosophila melanogaster* (Q9VP1),
and *Arabidopsis* (AtNHX1; BAB02053) (see Figures 4A-C, Figure 5 and Figure 8).
Based upon the homology, the polypeptide of the present invention may share at least
some biological activity with Na⁺/H⁺ antiporters. The polynucleotides and
polypeptides of the present invention, including agonists and/or fragments thereof,
10 have uses that include, but are not limited to conferring drought tolerance and/or salt
tolerance to plants. Moreover, the polynucleotides and polypeptides of the present
invention may also be useful in increasing the cultivatable range of crops within high
salinity soils.

The polypeptide of the present invention has been shown to comprise a sodium
15 binding site (See Figure 4). Such a binding site is also known as an amiloride binding
site.

In preferred embodiments, the following polypeptide is encompassed by the
present invention: ILEFDEELFFIYLLPPIIFNAGFQVKKKEFFRNFFITIMFF (SEQ
ID NO:13). Polynucleotide encoding this polypeptide is also provided.

20 The polypeptide of the present invention has also been shown to comprise a
FKBP-type peptidyl-prolyl cis-trans isomerase domain. FKBP [1,2,3] is the major
high-affinity binding protein, in vertebrates, for the immunosuppressive drug FK506.
It exhibits peptidyl-prolyl cis-trans isomerase activity (EC 5.2.1.8) (PPIase or
rotamase). PPIase is an enzyme that accelerates protein folding by catalyzing the cis-
25 trans isomerization of proline imidic peptide bonds in oligopeptides [4]. At least three
different forms of FKBP are known in mammalian species: FKBP-12, which is
cytosolic and inhibited by both FK506 and rapamycin; FKBP-13, which is membrane
associated and inhibited by both FK506 and rapamycin; and FKBP-25, which is
preferentially inhibited by rapamycin. The following publications are referenced
30 above and are hereby incorporated herein in their entirety: [1] Tropschug M., Wachter
E., Mayer S., Schoenbrunner E.R., Schmid F.X., *Nature* 346:674-677(1990); [2] Stein
R.L., *Curr. Biol.* 1:234-236(1991); [3] Siekierka J.J., Widerrecht G., Greulich H.,
Boulton D., Hung S.H.Y., Cryan J., Hodges P.J., Sigal N.H., *J. Biol. Chem.*
265:21011-21015(1990); and [4] Fischer G., Schmid F.X., *Biochemistry* 29:2205-
35 2212(1990).

Based upon the presence of the FKBP-type peptidyl-prolyl cis-trans isomerase
domain, the polypeptide of the present invention may share at least some biological

5 activity with FKBP-type peptidyl-prolyl cis-trans isomerase domain containing proteins.

In preferred embodiments, the following polypeptide is encompassed by the present invention: QDETPLLYSLVFGEGVVNDATSVVLS (SEQ ID NO:14). Polynucleotide encoding this polypeptide is also provided.

10 Although it is believed the encoded polypeptides may share at least some biological activities with Na⁺/H⁺ antiporters, a number of methods of determining the exact biological function of this clone are either known in the art or are described elsewhere herein. Briefly, the function of these clones may be construed by applying microarray methodology. The PpNHX1 and PpNHX2 clones in addition to other
15 clones of the present invention, may be arrayed on microchips for expression profiling. Depending on which polynucleotide probe is used to hybridize to the slides, a change in expression of a specific gene may provide additional insight into the function of this gene based upon the conditions being studied. For example, an observed increase or decrease in expression levels when the polynucleotide probe
20 used comes from tissue that has been cold treated might indicate a function in modulating cold tolerance, for example. In the case of PpNHX1 and PpNHX2, tissue deprived of water or stressed by other biotic or abiotic stresses (heat, drought, high light, high salt, etc.) should be used to extract RNA to prepare the probe.

In addition, the function of the protein may be assessed by applying
25 quantitative PCR methodology, for example. Real time quantitative PCR would provide the capability of following the expression of specific genes throughout the plant development cycle, for example. Quantitative PCR methodology requires only a nominal amount of tissue from each developmentally important step (3 days germinated seedlings, 1 week old seedlings [roots, shoots, and stems]; roots, leaves
30 and stems before the onset of flowering, flowers [different parts]; and/or developing embryos) is needed to perform such experiments. Therefore, the application of quantitative PCR methodology to refining the biological function of this polypeptide is encompassed by the present invention. Also encompassed by the present invention are quantitative PCR probes corresponding to the polynucleotide sequence provided
35 as SEQ ID NO:1 and SEQ ID NO: 3 and 4 (Figures 2A-D).

The function of the protein may also be assessed through complementation assays in yeast. For example, in the case of the PpNHX1 and PpNHX2 clones, transforming yeast deficient in sodium antiporter activity and assessing their ability to

5 grow in high salinity conditions would provide additional evidence the PpNHX1 and PpNHX2 clones have sodium antiporter activity. Alternatively, transforming wild type yeast with the PpNHX1 and PpNHX2 clones under increasing salinity conditions and assessing the ability of the transformed yeast to grow on media containing higher concentrations than non-transformed yeast could also establish the sodium antiporter
10 activity of PpNHX1 and PpNHX2 clones, in addition to its increased sodium transport efficiency. Additional assay conditions and methods that may be used in assessing the function of the polynucleotides and polypeptides of the present invention are known in the art, some of which are disclosed elsewhere herein.

Alternatively, a biological function of the encoded polypeptide may be
15 determined by disrupting a homologue of these polypeptides in *Synechosystis*. Cyanobacteria (blue-green algae) is considered a precursor to the plant chloroplast. It possesses both photosynthetic systems and many other metabolic processes reminiscent to those of plants. These processes are often targets for many commercial herbicides, and this organism has been widely used in the study of the mode of action
20 of many classes of herbicides. *Synechocystis* is one of the best-studied cyanobacteria. In addition to most of the features common to cyanobacteria, it offers many other added advantages. *Synechocystis* has a naturally occurring genetic transformation system, thus entailing vigorous and sophisticated genetic and molecular manipulation (e.g. targeted-gene disruption, gene replacement, etc.) applicable to some of the well-
25 characterized systems (*S. cerevisiae*, *E. coli*). Most importantly, the availability of the complete genomic sequence information of the *Synechocystis* affords an avenue for the rapid identification and cloning of gene(s) of interest, and elucidation of gene function through genetic and molecular means.

Moreover, a biological function of these polypeptides may be determined by
30 the application of antisense and/or sense methodology and the resulting generation of transgenic plants. Expressing a particular gene in either sense or antisense orientation in a transgenic plant can lead to respectively higher or lower expression levels of that particular gene. Altering the endogenous expression levels of a gene can lead to the observation of a particular phenotype that can then be used to derive indications on the
35 function of the gene. These genes can be either over-expressed or under expressed in every cell of the plant at all times using a strong ubiquitous promoter, or it can be expressed in one or more discrete parts of the plant using a well characterized tissue-specific promoter (i.e., a root promoter or a flower specific promoter), or it can be

5 expressed at a specified time of development using an inducible and/or a developmentally regulated promoter.

In the case of PpNHX1 and PpNHX2 transgenic plants, if no phenotype is apparent in normal growth conditions, observing the plants under stress conditions (deprivation of water, presence of high salt, or other biotic or abiotic stresses, such as cold, heat, drought, high light, etc.) may lead to understanding the function of the
10 gene. Therefore, the application of antisense and/or sense methodology to the creation of transgenic plants to refine the biological function of the polypeptide is encompassed by the present invention.

In preferred embodiments, the following N-terminal deletion mutants are
15 encompassed by the present invention: M1-S558, A2-S558, T3-S558, N4-S558, D5-S558, V6-S558, V7-S558, S8-S558, V9-S558, S10-S558, H11-S558, S12-S558, M13-S558, L14-S558, L15-S558, K16-S558, A17-S558, T18-S558, D19-S558, L20-S558, K21-S558, D22-S558, D23-S558, R24-S558, I25-S558, D26-S558, V27-S558, I28-S558, S29-S558, I30-S558, C31-S558, L32-S558, F33-S558, V34-S558, F35-S558,
20 L36-S558, L37-S558, C38-S558, A39-S558, C40-S558, I41-S558, V42-S558, L43-S558, G44-S558, H45-S558, L46-S558, L47-S558, E48-S558, E49-S558, N50-S558, R51-S558, W52-S558, M53-S558, N54-S558, E55-S558, S56-S558, I57-S558, T58-S558, A59-S558, L60-S558, L61-S558, L62-S558, G63-S558, L64-S558, F65-S558, T66-S558, G67-S558, S68-S558, I69-S558, V70-S558, L71-S558, I72-S558, S73-S558, S74-S558, K75-S558, G76-S558, Q77-S558, G78-S558, S79-S558, H80-S558,
25 I81-S558, L82-S558, E83-S558, F84-S558, D85-S558, E86-S558, E87-S558, L88-S558, F89-S558, F90-S558, I91-S558, Y92-S558, L93-S558, L94-S558, P95-S558, P96-S558, I97-S558, I98-S558, F99-S558, N100-S558, A101-S558, G102-S558, F103-S558, Q104-S558, V105-S558, K106-S558, K107-S558, K108-S558, E109-S558, F110-S558, F111-S558, R112-S558, N113-S558, F114-S558, I115-S558, T116-S558, I117-S558, M118-S558, F119-S558, F120-S558, G121-S558, V122-S558, I123-S558, G124-S558, V125-S558, F126-S558, I127-S558, S128-S558, F129-S558, G130-S558, I131-S558, I132-S558, S133-S558, T134-S558, G135-S558, S136-S558, W137-S558, Y138-S558, F139-S558, F140-S558, S141-S558, K142-S558,
35 F143-S558, G144-S558, L145-S558, K146-S558, N147-S558, L148-S558, P149-S558, I150-S558, R151-S558, D152-S558, I153-S558, L154-S558, A155-S558, I156-S558, G157-S558, V158-S558, I159-S558, F160-S558, S161-S558, A162-S558, T163-S558, D164-S558, S165-S558, V166-S558, C167-S558, T168-S558, L169-

5 S558, Q170-S558, V171-S558, L172-S558, N173-S558, Q174-S558, D175-S558,
E176-S558, T177-S558, P178-S558, L179-S558, L180-S558, Y181-S558, S182-
S558, L183-S558, V184-S558, F185-S558, G186-S558, E187-S558, G188-S558,
V189-S558, V190-S558, N191-S558, D192-S558, A193-S558, T194-S558, S195-
S558, V196-S558, V197-S558, L198-S558, S199-S558, R200-S558, A201-S558,
10 V202-S558, Q203-S558, T204-S558, Y205-S558, N206-S558, F207-S558, D208-
S558, N209-S558, F210-S558, T211-S558, S212-S558, L213-S558, E214-S558,
G215-S558, L216-S558, Q217-S558, I218-S558, G219-S558, G220-S558, S221-
S558, F222-S558, L223-S558, Y224-S558, L225-S558, F226-S558, F227-S558,
S228-S558, S229-S558, C230-S558, I231-S558, L232-S558, G233-S558, I234-S558,
15 A235-S558, S236-S558, G237-S558, L238-S558, I239-S558, S240-S558, A241-
S558, Y242-S558, I243-S558, I244-S558, K245-S558, T246-S558, M247-S558,
Y248-S558, F249-S558, G250-S558, R251-S558, H252-S558, S253-S558, T254-
S558, D255-S558, R256-S558, E257-S558, I258-S558, A259-S558, I260-S558,
M261-S558, T262-S558, L263-S558, M264-S558, A265-S558, Y266-S558, L267-
20 S558, S268-S558, Y269-S558, V270-S558, F271-S558, A272-S558, E273-S558,
L274-S558, F275-S558, Y276-S558, L277-S558, S278-S558, G279-S558, I280-S558,
L281-S558, S282-S558, V283-S558, F284-S558, F285-S558, C286-S558, G287-
S558, I288-S558, V289-S558, M290-S558, S291-S558, H292-S558, Y293-S558,
T294-S558, W295-S558, H296-S558, N297-S558, V298-S558, T299-S558, E300-
25 S558, N301-S558, S302-S558, R303-S558, I304-S558, T305-S558, S306-S558,
K307-S558, H308-S558, S309-S558, F310-S558, A311-S558, T312-S558, M313-
S558, S314-S558, F315-S558, I316-S558, A317-S558, E318-S558, T319-S558,
F320-S558, I321-S558, F322-S558, L323-S558, Y324-S558, V325-S558, G326-
S558, M327-S558, D328-S558, A329-S558, L330-S558, D331-S558, F332-S558,
30 E333-S558, K334-S558, W335-S558, K336-S558, M337-S558, M338-S558, Q339-
S558, S340-S558, S341-S558, F342-S558, T343-S558, E344-S558, S345-S558,
A346-S558, G347-S558, L348-S558, F349-S558, G350-S558, S351-S558, L352-
S558, L353-S558, F354-S558, L355-S558, V356-S558, L357-S558, L358-S558,
G359-S558, R360-S558, A361-S558, A362-S558, F363-S558, V364-S558, F365-
35 S558, P366-S558, L367-S558, S368-S558, A369-S558, L370-S558, S371-S558,
N372-S558, Y373-S558, S374-S558, T375-S558, K376-S558, S377-S558, P378-
S558, D379-S558, A380-S558, K381-S558, I382-S558, N383-S558, L384-S558,
R385-S558, Q386-S558, M387-S558, V388-S558, I389-S558, I390-S558, W391-

5 S558, W392-S558, A393-S558, G394-S558, L395-S558, M396-S558, R397-S558,
G398-S558, A399-S558, V400-S558, S401-S558, I402-S558, A403-S558, L404-
S558, A405-S558, F406-S558, N407-S558, Q408-S558, G409-S558, G410-S558,
D411-S558, A412-S558, K413-S558, D414-S558, S415-S558, N416-S558, Q417-
10 T424-S558, I425-S558, I426-S558, I427-S558, V428-S558, L429-S558, F430-S558,
S431-S558, T432-S558, I433-S558, V434-S558, F435-S558, G436-S558, T437-S558,
A438-S558, T439-S558, K440-S558, P441-S558, L442-S558, I443-S558, S444-S558,
W445-S558, L446-S558, L447-S558, P448-S558, P449-S558, H450-S558, F451-
S558, R452-S558, S453-S558, N454-S558, Y455-S558, S456-S558, D457-S558,
15 S458-S558, A459-S558, S460-S558, L461-S558, S462-S558, P463-S558, K464-
S558, A465-S558, S466-S558, L467-S558, D468-S558, A469-S558, D470-S558,
F471-S558, H472-S558, I473-S558, P474-S558, L475-S558, L476-S558, M477-
S558, D478-S558, T479-S558, E480-S558, R481-S558, E482-S558, E483-S558,
L484-S558, E485-S558, A486-S558, N487-S558, D488-S558, R489-S558, S490-
20 S558, T491-S558, I492-S558, N493-S558, Q494-S558, I495-S558, L496-S558,
N497-S558, G498-S558, L499-S558, P500-S558, C501-S558, P502-S558, Q503-
S558, S504-S558, I505-S558, G506-S558, M507-S558, L508-S558, L509-S558,
T510-S558, A511-S558, P512-S558, R513-S558, S514-S558, T515-S558, I516-S558,
H517-S558, H518-S558, V519-S558, W520-S558, R521-S558, K522-S558, F523-
25 S558, D524-S558, D525-S558, S526-S558, Y527-S558, M528-S558, R529-S558,
P530-S558, T531-S558, F532-S558, G533-S558, G534-S558, R535-S558, G536-
S558, Y537-S558, V538-S558, R539-S558, L540-S558, V541-S558, S542-S558,
R543-S558, R544-S558, D545-S558, M546-S558, D547-S558, I548-S558, Q549-
S558, E550-S558, D551-S558, E552-S558, of SEQ ID NO:2. Polynucleotide
30 sequences encoding these polypeptides are also provided.

In preferred embodiments, the following C-terminal deletion mutants are encompassed by the present invention: M1-S558, M1-H557, M1-D556, M1-E555, M1-L554, M1-I553, M1-E552, M1-D551, M1-E550, M1-Q549, M1-I548, M1-D547, M1-M546, M1-D545, M1-R544, M1-R543, M1-S542, M1-V541, M1-L540, M1-
35 R539, M1-V538, M1-Y537, M1-G536, M1-R535, M1-G534, M1-G533, M1-F532, M1-T531, M1-P530, M1-R529, M1-M528, M1-Y527, M1-S526, M1-D525, M1-D524, M1-F523, M1-K522, M1-R521, M1-W520, M1-V519, M1-H518, M1-H517, M1-I516, M1-T515, M1-S514, M1-R513, M1-P512, M1-A511, M1-T510, M1-L509,

5 M1-L508, M1-M507, M1-G506, M1-I505, M1-S504, M1-Q503, M1-P502, M1-C501,
M1-P500, M1-L499, M1-G498, M1-N497, M1-L496, M1-I495, M1-Q494, M1-N493,
M1-I492, M1-T491, M1-S490, M1-R489, M1-D488, M1-N487, M1-A486, M1-E485,
M1-L484, M1-E483, M1-E482, M1-R481, M1-E480, M1-T479, M1-D478, M1-
M477, M1-L476, M1-L475, M1-P474, M1-I473, M1-H472, M1-F471, M1-D470,
10 M1-A469, M1-D468, M1-L467, M1-S466, M1-A465, M1-K464, M1-P463, M1-
S462, M1-L461, M1-S460, M1-A459, M1-S458, M1-D457, M1-S456, M1-Y455,
M1-N454, M1-S453, M1-R452, M1-F451, M1-H450, M1-P449, M1-P448, M1-L447,
M1-L446, M1-W445, M1-S444, M1-I443, M1-L442, M1-P441, M1-K440, M1-T439,
M1-A438, M1-T437, M1-G436, M1-F435, M1-V434, M1-I433, M1-T432, M1-S431,
15 M1-F430, M1-L429, M1-V428, M1-I427, M1-I426, M1-I425, M1-T424, M1-I423,
M1-V422, M1-M421, M1-L420, M1-T419, M1-A418, M1-Q417, M1-N416, M1-
S415, M1-D414, M1-K413, M1-A412, M1-D411, M1-G410, M1-G409, M1-Q408,
M1-N407, M1-F406, M1-A405, M1-L404, M1-A403, M1-I402, M1-S401, M1-V400,
M1-A399, M1-G398, M1-R397, M1-M396, M1-L395, M1-G394, M1-A393, M1-
20 W392, M1-W391, M1-I390, M1-I389, M1-V388, M1-M387, M1-Q386, M1-R385,
M1-L384, M1-N383, M1-I382, M1-K381, M1-A380, M1-D379, M1-P378, M1-S377,
M1-K376, M1-T375, M1-S374, M1-Y373, M1-N372, M1-S371, M1-L370, M1-
A369, M1-S368, M1-L367, M1-P366, M1-F365, M1-V364, M1-F363, M1-A362,
M1-A361, M1-R360, M1-G359, M1-L358, M1-L357, M1-V356, M1-L355, M1-
25 F354, M1-L353, M1-L352, M1-S351, M1-G350, M1-F349, M1-L348, M1-G347,
M1-A346, M1-S345, M1-E344, M1-T343, M1-F342, M1-S341, M1-S340, M1-Q339,
M1-M338, M1-M337, M1-K336, M1-W335, M1-K334, M1-E333, M1-F332, M1-
D331, M1-L330, M1-A329, M1-D328, M1-M327, M1-G326, M1-V325, M1-Y324,
M1-L323, M1-F322, M1-I321, M1-F320, M1-T319, M1-E318, M1-A317, M1-I316,
30 M1-F315, M1-S314, M1-M313, M1-T312, M1-A311, M1-F310, M1-S309, M1-
H308, M1-K307, M1-S306, M1-T305, M1-I304, M1-R303, M1-S302, M1-N301,
M1-E300, M1-T299, M1-V298, M1-N297, M1-H296, M1-W295, M1-T294, M1-
Y293, M1-H292, M1-S291, M1-M290, M1-V289, M1-I288, M1-G287, M1-C286,
M1-F285, M1-F284, M1-V283, M1-S282, M1-L281, M1-I280, M1-G279, M1-S278,
35 M1-L277, M1-Y276, M1-F275, M1-L274, M1-E273, M1-A272, M1-F271, M1-
V270, M1-Y269, M1-S268, M1-L267, M1-Y266, M1-A265, M1-M264, M1-L263,
M1-T262, M1-M261, M1-I260, M1-A259, M1-I258, M1-E257, M1-R256, M1-D255,
M1-T254, M1-S253, M1-H252, M1-R251, M1-G250, M1-F249, M1-Y248, M1-

5 M247, M1-T246, M1-K245, M1-I244, M1-I243, M1-Y242, M1-A241, M1-S240,
M1-I239, M1-L238, M1-G237, M1-S236, M1-A235, M1-I234, M1-G233, M1-L232,
M1-I231, M1-C230, M1-S229, M1-S228, M1-F227, M1-F226, M1-L225, M1-Y224,
M1-L223, M1-F222, M1-S221, M1-G220, M1-G219, M1-I218, M1-Q217, M1-L216,
M1-G215, M1-E214, M1-L213, M1-S212, M1-T211, M1-F210, M1-N209, M1-
10 D208, M1-F207, M1-N206, M1-Y205, M1-T204, M1-Q203, M1-V202, M1-A201,
M1-R200, M1-S199, M1-L198, M1-V197, M1-V196, M1-S195, M1-T194, M1-
A193, M1-D192, M1-N191, M1-V190, M1-V189, M1-G188, M1-E187, M1-G186,
M1-F185, M1-V184, M1-L183, M1-S182, M1-Y181, M1-L180, M1-L179, M1-P178,
M1-T177, M1-E176, M1-D175, M1-Q174, M1-N173, M1-L172, M1-V171, M1-
15 Q170, M1-L169, M1-T168, M1-C167, M1-V166, M1-S165, M1-D164, M1-T163,
M1-A162, M1-S161, M1-F160, M1-I159, M1-V158, M1-G157, M1-I156, M1-A155,
M1-L154, M1-I153, M1-D152, M1-R151, M1-I150, M1-P149, M1-L148, M1-N147,
M1-K146, M1-L145, M1-G144, M1-F143, M1-K142, M1-S141, M1-F140, M1-F139,
M1-Y138, M1-W137, M1-S136, M1-G135, M1-T134, M1-S133, M1-I132, M1-I131,
20 M1-G130, M1-F129, M1-S128, M1-I127, M1-F126, M1-V125, M1-G124, M1-I123,
M1-V122, M1-G121, M1-F120, M1-F119, M1-M118, M1-I117, M1-T116, M1-I115,
M1-F114, M1-N113, M1-R112, M1-F111, M1-F110, M1-E109, M1-K108, M1-
K107, M1-K106, M1-V105, M1-Q104, M1-F103, M1-G102, M1-A101, M1-N100,
M1-F99, M1-I98, M1-I97, M1-P96, M1-P95, M1-L94, M1-L93, M1-Y92, M1-I91,
25 M1-F90, M1-F89, M1-L88, M1-E87, M1-E86, M1-D85, M1-F84, M1-E83, M1-L82,
M1-I81, M1-H80, M1-S79, M1-G78, M1-Q77, M1-G76, M1-K75, M1-S74, M1-S73,
M1-I72, M1-L71, M1-V70, M1-I69, M1-S68, M1-G67, M1-T66, M1-F65, M1-L64,
M1-G63, M1-L62, M1-L61, M1-L60, M1-A59, M1-T58, M1-I57, M1-S56, M1-E55,
M1-N54, M1-M53, M1-W52, M1-R51, M1-N50, M1-E49, M1-E48, M1-L47, M1-
30 L46, M1-H45, M1-G44, M1-L43, M1-V42, M1-I41, M1-C40, M1-A39, M1-C38,
M1-L37, M1-L36, M1-F35, M1-V34, M1-F33, M1-L32, M1-C31, M1-I30, M1-S29,
M1-I28, M1-V27, M1-D26, M1-I25, M1-R24, M1-D23, M1-D22, M1-K21, M1-L20,
M1-D19, M1-T18, M1-A17, M1-K16, M1-L15, M1-L14, M1-M13, M1-S12, M1-
H11, M1-S10, M1-V9, M1-S8, M1-V7, of SEQ ID NO:2. Polynucleotide sequences
35 encoding these polypeptides are also provided.

Many polynucleotide sequences, such as EST sequences, are publicly available and accessible through sequence databases. Some of these sequences are related to SEQ ID NO: 1 and SEQ ID NO: 3 and SEQ ID NO: 4 and may have been

5 publicly available prior to conception of the present invention. Preferably, such related polynucleotides are specifically excluded from the scope of the present invention. To list every related sequence would be cumbersome. Accordingly, preferably excluded from the present invention are one or more polynucleotides comprising a nucleotide sequence described by the general formula of a-b, where a is
10 any integer between 1 to 1677 of SEQ ID NO: 1, b is an integer between 15 to 1677, where both a and b correspond to the positions of nucleotide residues shown in SEQ ID NO:1, and where b is greater than or equal to a+14. Accordingly, preferably excluded from the present invention are one or more polynucleotides comprising a nucleotide sequence described by the general formula of a-b, where a is any integer
15 between 1 to 2624 of SEQ ID NO: 4, b is an integer between 15 to 2624, where both a and b correspond to the positions of nucleotide residues shown in SEQ ID NO: 4, and where b is greater than or equal to a+14.

Table 1

Gene No.	CDNA Clone ID	ATCC Deposit NO:Z	Vector	NT SEQ ID. No. X	Total NT Seq of Clone	5' NT of Start Codon of ORF	3' NT of ORF	AA Seq ID No. Y	Total AA of ORF
1.	PpNHX1	Acc. No xxx	pCR2.1	1	1677	1	1674	2	558
2.	PpNHX2	Acc. No xxx	pCR2.1	4	2624	14	2564	5	849

5 Table 1 summarizes the information corresponding to each "Gene No." described above. The nucleotide sequence identified as "NT SEQ ID NO:X" was assembled from partially homologous ("overlapping") sequences obtained from the "cDNA clone ID" identified in Table 1 and, in some cases, from additional related DNA clones. The overlapping sequences were assembled into a single contiguous
10 sequence of high redundancy (usually several overlapping sequences at each nucleotide position), resulting in a final sequence identified as SEQ ID NO:X.

The cDNA Clone ID was deposited and given the accession number and date provided in Table 1, referred to collectively herein as ATCC Deposit No:Z. "Vector" refers to the type of vector contained in the cDNA Clone ID. pCR2.1 was obtained
15 from Invitrogen, Inc.

"Total NT Seq. Of Clone" refers to the total number of nucleotides in the clone contig identified by "Gene No." The deposited clone may contain all or most of the sequence of SEQ ID NO:X. The nucleotide position of SEQ ID NO:X of the putative start codon (methionine) is identified as "5' NT of Start Codon of ORF."

20 The translated amino acid sequence, beginning with the methionine, is identified as "AA SEQ ID NO:Y," although other reading frames can also be easily translated using known molecular biology techniques. The polypeptides produced by these alternative open reading frames are specifically contemplated by the present invention.

25 The total number of amino acids within the open reading frame of SEQ ID NO:Y is identified as "Total AA of ORF".

SEQ ID NO:X (where X may be any of the polynucleotide sequences disclosed in the sequence listing) and the translated SEQ ID NO:Y (where Y may be any of the polypeptide sequences disclosed in the sequence listing) are sufficiently
30 accurate and otherwise suitable for a variety of uses well known in the art and described further below. For instance, SEQ ID NO:X is useful for designing nucleic acid hybridization probes that will detect nucleic acid sequences contained in SEQ ID NO:X or the cDNA contained in the deposited clone. These probes will also hybridize to nucleic acid molecules in biological samples, thereby enabling a variety
35 of forensic and diagnostic methods of the invention. Similarly, polypeptides identified from SEQ ID NO:Y may be used, for example, to generate antibodies which bind specifically to proteins containing the polypeptides and the proteins encoded by the cDNA clones identified in Table 1.

5 Nevertheless, DNA sequences generated by sequencing reactions can contain sequencing errors. The errors exist as misidentified nucleotides, or as insertions or deletions of nucleotides in the generated DNA sequence. The erroneously inserted or deleted nucleotides may cause frame shifts in the reading frames of the predicted amino acid sequence. In these cases, the predicted amino acid sequence diverges
10 from the actual amino acid sequence, even though the generated DNA sequence may be greater than 99.9% identical to the actual DNA sequence (for example, one base insertion or deletion in an open reading frame of over 1000 bases).

 Accordingly, for those applications requiring precision in the nucleotide sequence or the amino acid sequence, the present invention provides not only the
15 generated nucleotide sequence identified as SEQ ID NO:X and the predicted translated amino acid sequence identified as SEQ ID NO:Y, but also a sample of plasmid DNA containing a cDNA of the invention as set forth in Table 1. The nucleotide sequence of each clone can readily be determined by sequencing the clone in accordance with known methods. The predicted amino acid sequence can then be
20 verified. Moreover, the amino acid sequence of the protein encoded by a particular clone can also be directly determined by peptide sequencing or by expressing the protein in a suitable host cell containing the cDNA, collecting the protein, and determining its sequence.

 The present invention also relates to the genes corresponding to SEQ ID
25 NO:X, SEQ ID NO:Y, or the clone. The corresponding gene can be isolated in accordance with known methods using the sequence information disclosed herein. Such methods include preparing probes or primers from the disclosed sequence and identifying or amplifying the corresponding gene from appropriate sources of genomic material.

30 Also provided in the present invention are species homologs, allelic variants, and/or orthologs. The skilled artisan could, using procedures well-known in the art, obtain the polynucleotide sequence corresponding to full-length genes (including, but not limited to the full-length coding region), allelic variants, splice variants, orthologs, and/or species homologs of genes corresponding to SEQ ID NO:X, SEQ ID NO:Y, or
35 a deposited clone, relying on the sequence from the sequences disclosed herein or the clones deposited with the ATCC. For example, allelic variants and/or species homologs may be isolated and identified by making suitable probes or primers which correspond to the 5', 3', or internal regions of the sequences provided herein and

5 screening a suitable nucleic acid source for allelic variants and/or the desired
homologue.

The polypeptides of the invention can be prepared in any suitable manner. Such polypeptides include isolated naturally occurring polypeptides, recombinantly produced polypeptides, synthetically produced polypeptides, or polypeptides
10 produced by a combination of these methods. Means for preparing such polypeptides are well understood in the art.

The polypeptides may be in the form of the full-length protein, including the mature form (if applicable), or may be a part of a larger protein, such as a fusion protein (see below). It is often advantageous to include an additional amino acid
15 sequence which contains secretory or leader sequences, pro-sequences, sequences which aid in purification, such as multiple histidine residues, or an additional sequence for stability during recombinant production.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced
20 version of a polypeptide, including the full-length polypeptide, can be substantially purified using techniques described herein or otherwise known in the art, such as, for example, by the one-step method described in Smith and Johnson, Gene 67:31-40 (1988). Polypeptides of the invention also can be purified from natural, synthetic or recombinant sources using protocols described herein or otherwise known in the art,
25 such as, for example, antibodies of the invention raised against the mature and/or full-length form of the protein.

The present invention provides a polynucleotide comprising, or alternatively consisting of, the sequence identified as SEQ ID NO:X, and/or a cDNA provided in ATCC Deposit No. Z:. The present invention also provides a polypeptide comprising,
30 or alternatively consisting of, the sequence identified as SEQ ID NO:Y, and/or a polypeptide encoded by the cDNA provided in ATCC Deposit NO:Z. The present invention also provides polynucleotides encoding a polypeptide comprising, or alternatively consisting of the polypeptide sequence of SEQ ID NO:Y, and/or a polypeptide sequence encoded by the cDNA contained in ATCC Deposit No:Z.

35 Preferably, the present invention is directed to a polynucleotide comprising, or alternatively consisting of, the sequence identified as SEQ ID NO:X, and/or a cDNA provided in ATCC Deposit No.:Z that is less than, or equal to, a polynucleotide

5 sequence that is 5 mega basepairs, 1 mega basepairs, 0.5 mega basepairs, 0.1 mega basepairs, 50,000 basepairs, 20,000 basepairs, or 10,000 basepairs in length.

The present invention encompasses polynucleotides with sequences complementary to those of the polynucleotides of the present invention disclosed herein. Such sequences may be complementary to the sequence disclosed as SEQ ID
10 NO:X, the sequence contained in a deposit, and/or the nucleic acid sequence encoding the sequence disclosed as SEQ ID NO:Y.

The present invention also encompasses polynucleotides capable of hybridizing, preferably under reduced stringency conditions, more preferably under stringent conditions, and most preferably under highly stringent conditions, to
15 polynucleotides described herein. Examples of stringency conditions are shown in Table 2 below: highly stringent conditions are those that are at least as stringent as, for example, conditions A-F; stringent conditions are at least as stringent as, for example, conditions G-L; and reduced stringency conditions are at least as stringent as, for example, conditions M-R.

20

5 **TABLE 2**

Stringency Condition	Polynucleotide Hybrid±	Hybrid Length (bp) ‡	Hyridization Temperature and Buffer†	Wash Temperature and Buffer †
A	DNA:DNA	> or equal to 50	65°C; 1xSSC – or- 42°C; 1xSSC, 50% formamide	65°C; 0.3xSSC
B	DNA:DNA	< 50	Tb*; 1xSSC	Tb*; 1xSSC
C	DNA:RNA	> or equal to 50	67°C; 1xSSC –or- 45°C; 1xSSC, 50% formamide	67°C; 0.3xSSC
D	DNA:RNA	< 50	Td*; 1xSSC	Td*; 1xSSC
E	RNA:RNA	> or equal to 50	70°C; 1xSSC –or- 50°C; 1xSSC, 50% formamide	70°C; 0.3xSSC
F	RNA:RNA	< 50	Tf*; 1xSSC	Tf*; 1xSSC
G	DNA:DNA	> or equal to 50	65°C; 4xSSC –or- 45°C; 4xSSC, 50% formamide	65°C; 1xSSC
H	DNA:DNA	< 50	Th*; 4xSSC	Th*; 4xSSC
I	DNA:RNA	> or equal to 50	67°C; 4xSSC –or- 45°C; 4xSSC, 50% formamide	67°C; 1xSSC
J	DNA:RNA	< 50	Tj*; 4xSSC	Tj*; 4xSSC
K	RNA:RNA	> or equal to 50	70°C; 4xSSC	67°C; 1xSSC

			-or- 40°C; 6xSSC, 50% formamide	
L	RNA:RNA	< 50	Tl*; 2xSSC	Tl*; 2xSSC
M	DNA:DNA	> or equal to 50	50°C; 4xSSC -or- 40°C 6xSSC, 50% formamide	50°C; 2xSSC
N	DNA:DNA	< 50	Tn*; 6xSSC	Tn*; 6xSSC
O	DNA:RNA	> or equal to 50	55°C; 4xSSC -or- 42°C; 6xSSC, 50% formamide	55°C; 2xSSC
P	DNA:RNA	< 50	Tp*; 6xSSC	Tp*; 6xSSC
Q	RNA:RNA	> or equal to 50	60°C; 4xSSC -or- 45°C; 6xSSC, 50% formamide	60°C; 2xSSC
R	RNA:RNA	< 50	Tr*; 4xSSC	Tr*; 4xSSC

5

5 ‡: The “hybrid length” is the anticipated length for the hybridized region(s) of the hybridizing polynucleotides. When hybridizing a polynucleotide of unknown sequence, the hybrid is assumed to be that of the hybridizing polynucleotide of the present invention. When polynucleotides of known sequence are hybridized, the hybrid length can be determined by aligning the sequences of the polynucleotides and identifying the region or regions of optimal sequence complementarity. Methods of aligning two or more polynucleotide sequences and/or determining the percent identity between two polynucleotide sequences are well known in the art (e.g., MegAlign program of the DNA*Star suite of programs, etc).

15 †: SSPE (1xSSPE is 0.15M NaCl, 10mM NaH₂PO₄, and 1.25mM EDTA, pH 7.4) can be substituted for SSC (1xSSC is 0.15M NaCl and 15mM sodium citrate) in the hybridization and wash buffers; washes are performed for 15 minutes after hybridization is complete. The hybridizations and washes may additionally include 5X Denhardt’s reagent, .5-1.0% SDS, 100ug/ml denatured, 20 fragmented salmon sperm DNA, 0.5% sodium pyrophosphate, and up to 50% formamide.

*T_b – T_r: The hybridization temperature for hybrids anticipated to be less than 50 base pairs in length should be 5-10°C less than the melting temperature T_m of the hybrids there T_m is determined according to the following equations. For hybrids less than 18 base pairs in length, $T_m(^{\circ}\text{C}) = 2(\# \text{ of A} + \text{T bases}) + 4(\# \text{ of G} + \text{C bases})$. For hybrids between 18 and 49 base pairs in length, $T_m(^{\circ}\text{C}) = 81.5 + 16.6(\log_{10}[\text{Na}^+]) + 0.41(\% \text{G+C}) - (600/\text{N})$, where N is the number of bases in the hybrid, and [Na⁺] is the concentration of sodium ions in the hybridization buffer ([Na⁺] for 1xSSC = .165 M).

±: The present invention encompasses the substitution of any one, or more DNA or RNA hybrid partners with either a PNA, or a modified polynucleotide. Such modified polynucleotides are known in the art and are more particularly described elsewhere herein.

5 Additional examples of stringency conditions for polynucleotide hybridization are provided, for example, in Sambrook, J., E.F. Fritsch, and T.Maniatis, 1989, Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, NY, chapters 9 and 11, and Current Protocols in Molecular Biology, 1995, F.M., Ausubel et al., eds, John Wiley and Sons, Inc., sections 2.10 and
10 6.3-6.4, which are hereby incorporated by reference herein.

Preferably, such hybridizing polynucleotides have at least 70% sequence identity (more preferably, at least 80% identity; and most preferably at least 90% or 95% identity) with the polynucleotide of the present invention to which they hybridize, where sequence identity is determined by comparing the sequences of the
15 hybridizing polynucleotides when aligned so as to maximize overlap and identity while minimizing sequence gaps. The determination of identity is well known in the art, and discussed more specifically elsewhere herein.

The invention encompasses the application of PCR methodology to the polynucleotide sequences of the present invention, the clone deposited with the
20 ATCC, and/or the cDNA encoding the polypeptides of the present invention. PCR techniques for the amplification of nucleic acids are described in US Patent No. 4, 683, 195 and Saiki et al., Science, 239:487-491 (1988). PCR, for example, may include the following steps, of denaturation of template nucleic acid (if double-stranded), annealing of primer to target, and polymerization. The nucleic acid probed
25 or used as a template in the amplification reaction may be genomic DNA, cDNA, RNA, or a PNA. PCR may be used to amplify specific sequences from genomic DNA, specific RNA sequence, and/or cDNA transcribed from mRNA. References for the general use of PCR techniques, including specific method parameters, include Mullis et al., Cold Spring Harbor Symp. Quant. Biol., 51:263, (1987), Ehrlich (ed),
30 PCR Technology, Stockton Press, NY, 1989; Ehrlich et al., Science, 252:1643-1650, (1991); and "PCR Protocols, A Guide to Methods and Applications", Eds., Innis et al., Academic Press, New York, (1990).

Signal Sequences

35 The present invention also encompasses mature forms of the polypeptide comprising, or alternatively consisting of, the polypeptide sequence of SEQ ID NO:Y, the polypeptide encoded by the polynucleotide described as SEQ ID NO:X, and/or the polypeptide sequence encoded by a cDNA in the deposited clone. The present

5 invention also encompasses polynucleotides encoding mature forms of the present invention, such as, for example the polynucleotide sequence of SEQ ID NO:X, and/or the polynucleotide sequence provided in a cDNA of the deposited clone.

According to the signal hypothesis, proteins secreted by eukaryotic cells have a signal or secretary leader sequence which is cleaved from the mature protein once
10 export of the growing protein chain across the rough endoplasmic reticulum has been initiated. Most eukaryotic cells cleave secreted proteins with the same specificity. However, in some cases, cleavage of a secreted protein is not entirely uniform, which results in two or more mature species of the protein. Further, it has long been known that cleavage specificity of a secreted protein is ultimately determined by the primary
15 structure of the complete protein, that is, it is inherent in the amino acid sequence of the polypeptide.

Methods for predicting whether a protein has a signal sequence, as well as the cleavage point for that sequence, are available. For instance, the method of McGeoch, *Virus Res.* 3:271-286 (1985), uses the information from a short N-terminal
20 charged region and a subsequent uncharged region of the complete (uncleaved) protein. The method of von Heinje, *Nucleic Acids Res.* 14:4683-4690 (1986) uses the information from the residues surrounding the cleavage site, typically residues -13 to +2, where +1 indicates the amino terminus of the secreted protein. The accuracy of predicting the cleavage points of known mammalian secretory proteins for each of
25 these methods is in the range of 75-80%. (von Heinje, supra.) However, the two methods do not always produce the same predicted cleavage point(s) for a given protein.

The established method for identifying the location of signal sequences, in addition, to their cleavage sites has been the SignalP program (v1.1) developed by
30 Henrik Nielsen et al., *Protein Engineering* 10:1-6 (1997). The program relies upon the algorithm developed by von Heinje, though provides additional parameters to increase the prediction accuracy.

More recently, a hidden Markov model has been developed (H. Neilson, et al., *Ismb* 1998;6:122-30), which has been incorporated into the more recent SignalP
35 (v2.0). This new method increases the ability to identify the cleavage site by discriminating between signal peptides and uncleaved signal anchors. The present invention encompasses the application of the method disclosed therein to the

5 prediction of the signal peptide location, including the cleavage site, to any of the polypeptide sequences of the present invention.

As one of ordinary skill would appreciate, however, cleavage sites sometimes vary from organism to organism and cannot be predicted with absolute certainty. Accordingly, the polypeptide of the present invention may contain a signal sequence.

10 Polypeptides of the invention which comprise a signal sequence have an N-terminus beginning within 5 residues (i.e., + or - 5 residues, or preferably at the -5, -4, -3, -2, -1, +1, +2, +3, +4, or +5 residue) of the predicted cleavage point. Similarly, it is also recognized that in some cases, cleavage of the signal sequence from a secreted protein is not entirely uniform, resulting in more than one secreted species. These

15 polypeptides, and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Moreover, the signal sequence identified by the above analysis may not necessarily predict the naturally occurring signal sequence. For example, the naturally occurring signal sequence may be further upstream from the predicted signal

20 sequence. However, it is likely that the predicted signal sequence will be capable of directing the secreted protein to the ER. Nonetheless, the present invention provides the mature protein produced by expression of the polynucleotide sequence of SEQ ID NO:X and/or the polynucleotide sequence contained in the cDNA of a deposited clone, in a mammalian cell (e.g., COS cells, as described below). These polypeptides,

25 and the polynucleotides encoding such polypeptides, are contemplated by the present invention.

Polynucleotide and Polypeptide Variants

The present invention also encompasses variants (e.g., allelic variants,

30 orthologs, etc.) of the polynucleotide sequence disclosed herein in SEQ ID NO:X, the complementary strand thereto, and/or the cDNA sequence contained in the deposited clone.

The present invention also encompasses variants of the polypeptide sequence, and/or fragments therein, disclosed in SEQ ID NO:Y and/or encoded by a deposited

35 clone.

"Variant" refers to a polynucleotide or polypeptide differing from the polynucleotide or polypeptide of the present invention, but retaining essential

5 properties thereof. Generally, variants are overall closely similar, and, in many regions, identical to the polynucleotide or polypeptide of the present invention.

The present invention is also directed to polynucleotide sequences which comprise, or alternatively consist of, a polynucleotide sequence which is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to, the
10 following non-limiting examples, the polynucleotide sequence of the coding region of the sequence in SEQ ID NO:X, the complementary strand of said coding region, the polynucleotide sequence provided in a cDNA of the deposited clone, the complementary strand of said deposited cDNA, a polynucleotide sequence encoding the polypeptide identified as SEQ ID NO:Y, a polynucleotide sequence encoding the
15 polypeptide of a cDNA provided in the deposited clone. The invention also encompasses polynucleotide fragments of any of the polynucleotide sequences provided herein.

The present invention also encompasses polynucleotide sequences which hybridize to any of the provided polynucleotide sequences of the present invention,
20 including the aforementioned polynucleotide fragments, under stringent or lower stringent hybridization conditions. Polypeptides encoded by the polynucleotide sequences which hybridize to the polynucleotide sequences of the present invention are also encompassed by the present invention.

The present invention encompasses polypeptide sequences which comprise, or
25 alternatively consist of, an amino acid sequence which is at least 80%, 98%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to, the following non-limited examples, the polypeptide sequence identified as SEQ ID NO:Y, the polypeptide sequence encoded by a cDNA provided in the deposited clone, and/or polypeptide fragments of any of the polypeptides provided herein.

30 By a nucleic acid having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the nucleic acid is identical to the reference sequence except that the nucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the polypeptide. In
35 other words, to obtain a nucleic acid having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted or substituted with another nucleotide, or a number of nucleotides up to 5% of the total nucleotides in the reference sequence may

5 be inserted into the reference sequence. The query sequence may be an entire sequence referenced in Table 1, the ORF (open reading frame), or any fragment specified as described herein.

As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%,
10 or 99% identical to a nucleotide sequence of the present invention can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the CLUSTALW computer program (Thompson, J.D., et al., *Nucleic*
15 *Acids Research*, 2(22):4673-4680, (1994)), which is based on the algorithm of Higgins, D.G., et al., *Computer Applications in the Biosciences (CABIOS)*, 8(2):189-191, (1992). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result of said global sequence alignment is in percent identity. Preferred parameters used in
20 a CLUSTALW alignment of DNA sequences to calculate percent identity are: Matrix=BLOSUM, k-tuple=1, Number of Top Diagonals=5, Gap Penalty=3, Gap Open Penalty 10, Gap Extension Penalty=0, Scoring Method=Percent, Window Size=5 or the length of the subject nucleotide sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence because of 5' or 3'
25 deletions, not because of internal deletions, a manual correction must be made to the results. This is because the CLUSTALW program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are
30 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the CLUSTALW sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above CLUSTALW program using the specified parameters, to arrive at a final percent identity score.
35 This corrected score is what may be used for the purposes of the present invention. Only bases outside the 5' and 3' bases of the subject sequence, as displayed by the CLUSTALW alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

5 For example, a 90 base subject sequence is aligned to a 100 base query
sequence to determine percent identity. The deletions occur at the 5' end of the
subject sequence and therefore, the CLUSTALW alignment does not show a
matched/alignment of the first 10 bases at 5' end. The 10 unpaired bases represent
10% of the sequence (number of bases at the 5' and 3' ends not matched/total number
10 of bases in the query sequence) so 10% is subtracted from the percent identity score
calculated by the CLUSTALW program. If the remaining 90 bases were perfectly
matched the final percent identity would be 90%. In another example, a 90 base
subject sequence is compared with a 100 base query sequence. This time the
deletions are internal deletions so that there are no bases on the 5' or 3' of the subject
15 sequence which are not matched/aligned with the query. In this case the percent
identity calculated by CLUSTALW is not manually corrected. Once again, only
bases 5' and 3' of the subject sequence which are not matched/aligned with the query
sequence are manually corrected for. No other manual corrections are required for the
purposes of the present invention.

20 By a polypeptide having an amino acid sequence at least, for example, 95%
"identical" to a query amino acid sequence of the present invention, it is intended that
the amino acid sequence of the subject polypeptide is identical to the query sequence
except that the subject polypeptide sequence may include up to five amino acid
alterations per each 100 amino acids of the query amino acid sequence. In other
25 words, to obtain a polypeptide having an amino acid sequence at least 95% identical
to a query amino acid sequence, up to 5% of the amino acid residues in the subject
sequence may be inserted, deleted, or substituted with another amino acid. These
alterations of the reference sequence may occur at the amino or carboxy terminal
positions of the reference amino acid sequence or anywhere between those terminal
30 positions, interspersed either individually among residues in the reference sequence or
in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 80%, 85%,
91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to, for instance, an
amino acid sequence referenced in Table 1 (SEQ ID NO:Y) or to the amino acid
35 sequence encoded by cDNA contained in a deposited clone, can be determined
conventionally using known computer programs. A preferred method for determining
the best overall match between a query sequence (a sequence of the present invention)
and a subject sequence, also referred to as a global sequence alignment, can be

5 determined using the CLUSTALW computer program (Thompson, J.D., et al., *Nucleic Acids Research*, 2(22):4673-4680, (1994)), which is based on the algorithm of Higgins, D.G., et al., *Computer Applications in the Biosciences (CABIOS)*, 8(2):189-191, (1992). In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by converting U's to T's. The result
10 of said global sequence alignment is in percent identity. Preferred parameters used in a CLUSTALW alignment of DNA sequences to calculate percent identity are: Matrix=BLOSUM, k-tuple=1, Number of Top Diagonals=5, Gap Penalty=3, Gap Open Penalty 10, Gap Extension Penalty=0, Scoring Method=Percent, Window Size=5 or the length of the subject nucleotide sequence, whichever is shorter.

15 If the subject sequence is shorter than the query sequence due to N- or C-terminal deletions, not because of internal deletions, a manual correction must be made to the results. This is because the CLUSTALW program does not account for N- and C-terminal truncations of the subject sequence when calculating global percent identity. For subject sequences truncated at the N- and C-termini, relative to the
20 query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the CLUSTALW sequence alignment. This percentage is then subtracted
25 from the percent identity, calculated by the above CLUSTALW program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what may be used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually
30 adjusting the percent identity score. That is, only query residue positions outside the farthest N- and C-terminal residues of the subject sequence.

For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the N-terminus of the subject sequence and therefore, the CLUSTALW alignment does not
35 show a matching/alignment of the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C-termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the CLUSTALW program. If

5 the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal deletions so there are no residues at the N- or C-termini of the subject sequence, which are not matched/aligned with the query. In this case the percent identity calculated by CLUSTALW is not
10 manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the CLUSTALW alignment, which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are required for the purposes of the present invention.

The variants may contain alterations in the coding regions, non-coding
15 regions, or both. Especially preferred are polynucleotide variants containing alterations which produce silent substitutions, additions, or deletions, but do not alter the properties or activities of the encoded polypeptide. Nucleotide variants produced by silent substitutions due to the degeneracy of the genetic code are preferred. Moreover, variants in which 5-10, 1-5, or 1-2 amino acids are substituted, deleted, or
20 added in any combination are also preferred. Polynucleotide variants can be produced for a variety of reasons, e.g., to optimize codon expression for a particular host (change codons in the mRNA to those preferred by a bacterial host such as *E. coli*).

Naturally occurring variants are called "allelic variants," and refer to one of several alternate forms of a gene occupying a given locus on a chromosome of an
25 organism. (Genes II, Lewin, B., ed., John Wiley & Sons, New York (1985).) These allelic variants can vary at either the polynucleotide and/or polypeptide level and are included in the present invention. Alternatively, non-naturally occurring variants may be produced by mutagenesis techniques or by direct synthesis.

Using known methods of protein engineering and recombinant DNA
30 technology, variants may be generated to improve or alter the characteristics of the polypeptides of the present invention. For instance, one or more amino acids can be deleted from the N-terminus or C-terminus of the secreted protein without substantial loss of biological function. The authors of Ron et al., *J. Biol. Chem.* 268: 2984-2988 (1993), reported variant KGF proteins having heparin binding activity even after
35 deleting 3, 8, or 27 amino-terminal amino acid residues. Similarly, Interferon gamma exhibited up to ten times higher activity after deleting 8-10 amino acid residues from the carboxy terminus of this protein (Dobeli et al., *J. Biotechnology* 7:199-216 (1988)).

5 Moreover, ample evidence demonstrates that variants often retain a biological activity similar to that of the naturally occurring protein. For example, Gayle and coworkers (J. Biol. Chem 268:22105-22111 (1993)) conducted extensive mutational analysis of human cytokine IL-1a. They used random mutagenesis to generate over
10 the entire length of the molecule. Multiple mutations were examined at every possible amino acid position. The investigators found that "[m]ost of the molecule could be altered with little effect on either [binding or biological activity]." In fact, only 23 unique amino acid sequences, out of more than 3,500 nucleotide sequences examined, produced a protein that significantly differed in activity from wild-type.

15 Furthermore, even if deleting one or more amino acids from the N-terminus or C-terminus of a polypeptide results in modification or loss of one or more biological functions, other biological activities may still be retained. For example, the ability of a deletion variant to induce and/or to bind antibodies which recognize the protein will likely be retained when less than the majority of the residues of the protein are
20 removed from the N-terminus or C-terminus. Whether a particular polypeptide lacking N- or C-terminal residues of a protein retains such immunogenic activities can readily be determined by routine methods described herein and otherwise known in the art.

 Thus, the invention further includes polypeptide variants which show
25 substantial biological activity. Such variants include deletions, insertions, inversions, repeats, and substitutions selected according to general rules known in the art so as have little effect on activity. For example, guidance concerning how to make phenotypically silent amino acid substitutions is provided in Bowie et al., Science 247:1306-1310 (1990), wherein the authors indicate that there are two main strategies
30 for studying the tolerance of an amino acid sequence to change.

 The first strategy exploits the tolerance of amino acid substitutions by natural selection during the process of evolution. By comparing amino acid sequences in different species, conserved amino acids can be identified. These conserved amino acids are likely important for protein function. In contrast, the amino acid positions
35 where substitutions have been tolerated by natural selection indicates that these positions are not critical for protein function. Thus, positions tolerating amino acid substitution could be modified while still maintaining biological activity of the protein.

5 The second strategy uses genetic engineering to introduce amino acid changes at specific positions of a cloned gene to identify regions critical for protein function. For example, site directed mutagenesis or alanine-scanning mutagenesis (introduction of single alanine mutations at every residue in the molecule) can be used. (Cunningham and Wells, Science 244:1081-1085 (1989).) The resulting mutant
10 molecules can then be tested for biological activity.

 As the authors state, these two strategies have revealed that proteins are surprisingly tolerant of amino acid substitutions. The authors further indicate which amino acid changes are likely to be permissive at certain amino acid positions in the protein. For example, most buried (within the tertiary structure of the protein) amino
15 acid residues require nonpolar side chains, whereas few features of surface side chains are generally conserved. Moreover, tolerated conservative amino acid substitutions involve replacement of the aliphatic or hydrophobic amino acids Ala, Val, Leu and Ile; replacement of the hydroxyl residues Ser and Thr; replacement of the acidic residues Asp and Glu; replacement of the amide residues Asn and Gln, replacement of
20 the basic residues Lys, Arg, and His; replacement of the aromatic residues Phe, Tyr, and Trp, and replacement of the small-sized amino acids Ala, Ser, Thr, Met, and Gly.

 Besides conservative amino acid substitution, variants of the present invention include, but are not limited to, the following: (i) substitutions with one or more of the non-conserved amino acid residues, where the substituted amino acid residues may or
25 may not be one encoded by the genetic code, or (ii) substitution with one or more of amino acid residues having a substituent group, or (iii) fusion of the mature polypeptide with another compound, such as a compound to increase the stability and/or solubility of the polypeptide (for example, polyethylene glycol), or (iv) fusion of the polypeptide with additional amino acids, such as, for example, an IgG Fc
30 fusion region peptide, or leader or secretory sequence, or a sequence facilitating purification. Such variant polypeptides are deemed to be within the scope of those skilled in the art from the teachings herein.

 For example, polypeptide variants containing amino acid substitutions of charged amino acids with other charged or neutral amino acids may produce proteins
35 with improved characteristics, such as less aggregation. Aggregation of pharmaceutical formulations both reduces activity and increases clearance due to the aggregate's immunogenic activity. (Pinckard et al., Clin. Exp. Immunol. 2:331-340

5 (1967); Robbins et al., Diabetes 36: 838-845 (1987); Cleland et al., Crit. Rev. Therapeutic Drug Carrier Systems 10:307-377 (1993).

Moreover, the invention further includes polypeptide variants created through the application of molecular evolution ("DNA Shuffling") methodology to the polynucleotide disclosed as SEQ ID NO:X, the sequence of the clone submitted in a
10 deposit, and/or the cDNA encoding the polypeptide disclosed as SEQ ID NO:Y. Such DNA Shuffling technology is known in the art and more particularly described elsewhere herein (e.g., WPC, Stemmer, PNAS, 91:10747, (1994)), and example 39).

A further embodiment of the invention relates to a polypeptide which comprises the amino acid sequence of the present invention having an amino acid
15 sequence which contains at least one amino acid substitution, but not more than 50 amino acid substitutions, even more preferably, not more than 40 amino acid substitutions, still more preferably, not more than 30 amino acid substitutions, and still even more preferably, not more than 20 amino acid substitutions. Of course, in order of ever-increasing preference, it is highly preferable for a peptide or polypeptide
20 to have an amino acid sequence which comprises the amino acid sequence of the present invention, which contains at least one, but not more than 10, 9, 8, 7, 6, 5, 4, 3, 2 or 1 amino acid substitutions. In specific embodiments, the number of additions, substitutions, and/or deletions in the amino acid sequence of the present invention or fragments thereof (e.g., the mature form and/or other fragments described herein), is
25 1-5, 5-10, 5-25, 5-50, 10-50 or 50-150, conservative amino acid substitutions are preferable.

Polynucleotide and Polypeptide Fragments

The present invention is directed to polynucleotide fragments of the
30 polynucleotides of the invention, in addition to polypeptides encoded therein by said polynucleotides and/or fragments.

In the present invention, a "polynucleotide fragment" refers to a short polynucleotide having a nucleic acid sequence which: is a portion of that contained in a deposited clone, or encoding the polypeptide encoded by the cDNA in a deposited
35 clone; is a portion of that shown in SEQ ID NO:X or the complementary strand thereto, or is a portion of a polynucleotide sequence encoding the polypeptide of SEQ ID NO:Y. The nucleotide fragments of the invention are preferably at least about 15 nt, and more preferably at least about 20 nt, still more preferably at least about 30 nt,

5 and even more preferably, at least about 40 nt, at least about 50 nt, at least about 75
nt, or at least about 150 nt in length. A fragment "at least 20 nt in length," for
example, is intended to include 20 or more contiguous bases from the cDNA sequence
contained in a deposited clone or the nucleotide sequence shown in SEQ ID NO:X. In
this context "about" includes the particularly recited value, a value larger or smaller
10 by several (5, 4, 3, 2, or 1) nucleotides, at either terminus, or at both termini. These
nucleotide fragments have uses that include, but are not limited to, as diagnostic
probes and primers as discussed herein. Of course, larger fragments (e.g., 50, 150,
500, 600, 2000 nucleotides) are preferred.

Moreover, representative examples of polynucleotide fragments of the
15 invention, include, for example, fragments comprising, or alternatively consisting of,
a sequence from about nucleotide number 1-50, 51-100, 101-150, 151-200, 201-250,
251-300, 301-350, 351-400, 401-450, 451-500, 501-550, 551-600, 651-700, 701-750,
751-800, 800-850, 851-900, 901-950, 951-1000, 1001-1050, 1051-1100, 1101-1150,
1151-1200, 1201-1250, 1251-1300, 1301-1350, 1351-1400, 1401-1450, 1451-1500,
20 1501-1550, 1551-1600, 1601-1650, 1651-1700, 1701-1750, 1751-1800, 1801-1850,
1851-1900, 1901-1950, 1951-2000, or 2001 to the end of SEQ ID NO:X, or the
complementary strand thereto, or the cDNA contained in a deposited clone. In this
context "about" includes the particularly recited ranges, and ranges larger or smaller
by several (5, 4, 3, 2, or 1) nucleotides, at either terminus or at both termini.
25 Preferably, these fragments encode a polypeptide which has biological activity. More
preferably, these polynucleotides can be used as probes or primers as discussed
herein. Also encompassed by the present invention are polynucleotides which
hybridize to these nucleic acid molecules under stringent hybridization conditions or
lower stringency conditions, as are the polypeptides encoded by these
30 polynucleotides.

In the present invention, a "polypeptide fragment" refers to an amino acid
sequence which is a portion of that contained in SEQ ID NO:Y or encoded by the
cDNA contained in a deposited clone. Protein (polypeptide) fragments may be "free-
standing," or comprised within a larger polypeptide of which the fragment forms a
35 part or region, most preferably as a single continuous region. Representative
examples of polypeptide fragments of the invention, include, for example, fragments
comprising, or alternatively consisting of, from about amino acid number 1-20, 21-40,
41-60, 61-80, 81-100, 102-120, 121-140, 141-160, or 161 to the end of the coding

5 region. Moreover, polypeptide fragments can be about 20, 30, 40, 50, 60, 70, 80, 90, 100, 110, 120, 130, 140, or 150 amino acids in length. In this context "about" includes the particularly recited ranges or values, and ranges or values larger or smaller by several (5, 4, 3, 2, or 1) amino acids, at either extreme or at both extremes. Polynucleotides encoding these polypeptides are also encompassed by the invention.

10 Preferred polypeptide fragments include the full-length protein. Further preferred polypeptide fragments include the full-length protein having a continuous series of deleted residues from the amino or the carboxy terminus, or both. For example, any number of amino acids, ranging from 1-60, can be deleted from the amino terminus of the full-length polypeptide. Similarly, any number of amino acids,
15 ranging from 1-30, can be deleted from the carboxy terminus of the full-length protein. Furthermore, any combination of the above amino and carboxy terminus deletions are preferred. Similarly, polynucleotides encoding these polypeptide fragments are also preferred.

Also preferred are polypeptide and polynucleotide fragments characterized by
20 structural or functional domains, such as fragments that comprise alpha-helix and alpha-helix forming regions, beta-sheet and beta-sheet-forming regions, turn and turn-forming regions, coil and coil-forming regions, hydrophilic regions, hydrophobic regions, alpha amphipathic regions, beta amphipathic regions, flexible regions, surface-forming regions, substrate binding region, and high antigenic index regions.
25 Polypeptide fragments of SEQ ID NO:Y falling within conserved domains are specifically contemplated by the present invention. Moreover, polynucleotides encoding these domains are also contemplated.

Other preferred polypeptide fragments are biologically active fragments. Biologically active fragments are those exhibiting activity similar, but not necessarily
30 identical, to an activity of the polypeptide of the present invention. The biological activity of the fragments may include an improved desired activity, or a decreased undesirable activity. Polynucleotides encoding these polypeptide fragments are also encompassed by the invention.

In a preferred embodiment, the functional activity displayed by a polypeptide
35 encoded by a polynucleotide fragment of the invention may be one or more biological activities typically associated with the full-length polypeptide of the invention. Illustrative of these biological activities includes the fragments ability to bind to at least one of the same antibodies which bind to the full-length protein, the fragments

5 ability to interact with at least one of the same proteins which bind to the full-length,
the fragments ability to elicit at least one of the same immune responses as the full-
length protein (i.e., to cause the immune system to create antibodies specific to the
same epitope, etc.), the fragments ability to bind to at least one of the same
10 polynucleotides as the full-length protein, the fragments ability to bind to a receptor of
the full-length protein, the fragments ability to bind to a ligand of the full-length
protein, and the fragments ability to multimerize with the full-length protein.
However, the skilled artisan would appreciate that some fragments may have
biological activities which are desirable and directly inapposite to the biological
activity of the full-length protein. The functional activity of polypeptides of the
15 invention, including fragments, variants, derivatives, and analogs thereof can be
determined by numerous methods available to the skilled artisan, some of which are
described elsewhere herein.

Epitopes and Antibodies

20

In the present invention, "epitopes" refer to polypeptide fragments having
antigenic and/or immunogenic activity in an organism, preferably an animal or a
plant. A preferred embodiment of the present invention relates to a polypeptide
fragment comprising an epitope, as well as the polynucleotide encoding this fragment.
25 A region of a protein molecule to which an antibody can bind is defined as an
"antigenic epitope". In contrast, an "immunogenic epitope" is defined as a part of a
protein that elicits an antibody response (i.e., in animals), or a defensive response (i.e.,
in plants). (See, for instance, Geyson et al., Proc. Natl. Acad. Sci. USA 81:3998-4002
(1983)).

30 Epitope fragments may be produced by any conventional means known in the
art. Epitopes that are immunogenic may be useful for inducing antibodies according
to methods well known in the art. Such epitopes may be presented together with a
carrier protein such as an albumin, or may be presented without a carrier if the epitope
is of sufficient length (greater than or equal to 25 amino acids). Nonetheless, epitopes
35 have been found to elicit an immune response that comprise as few as 8 to 10 amino
acids. Epitopes of the present invention may be linear epitopes derived from a
denatured polypeptide (e.g., as found in Western blots).

5 In the present invention, antigenic epitopes preferably contain a sequence of at least seven, more preferably at least nine, and most preferably between about 15 to about 30 amino acids. Antigenic epitopes are useful to raise antibodies, including monoclonal antibodies, that specifically bind the epitope. (See, for instance, Wilson et al., *Cell* 37:767-778 (1984); Sutcliffe, J.G., et al., *Science* 219:660-666 (1983)).

10 As used herein, the term “antibody” (Ab) or “monoclonal antibody” (Mab) is meant to include intact molecules, as well as, antibody fragments (such as, for example, Fab and F(ab')₂ fragments) which are capable of specifically binding to protein. Fab and F(ab')₂ fragments lack the Fc fragment of intact antibody, clear more rapidly from the circulation of the animal or plant, and may have less non-specific tissue binding than an intact antibody (Wahl et al., *J. Nucl. Med* 24:316-325
15 (1983)). Thus, these fragments are preferred, as well as the products of a FAB or other immunoglobulin expression library. Moreover, antibodies of the present invention include chimeric, single chain, and humanized antibodies.

The present invention is further directed to antibodies specific to polypeptides,
20 polypeptide fragments, and polypeptide variants of the present invention. The antibodies of the present invention may include, but are not limited to, polyclonal, monoclonal, humanized, fully-human, monospecific, bispecific, trispecific, heteroconjugate, chimeric, single chain, variable light chain, variable heavy chain, one or more complementarity-determining regions (CDRs), phage-display derived
25 antibodies (those derived from a Fab expression library, etc.), anti-idiotypic antibodies, and those antibodies having enzymatic activity (i.e., catalytic antibodies). The antibodies of the present invention may be comprised of any of the currently known antibody isotypes (e.g., IgD, IgM, IgE, IgG, IgY, and IgA, etc.), the human IgA and IgG subclasses (e.g., IgG1, IgG2, IgG3, IgG4, IgA1, or IgA2), or the mouse IgA and
30 IgG subclasses (e.g., IgG1, IgG2a, IgG2b, IgG3, IgA1, or IgA2).

The antibodies of the present invention may comprise polyclonal antibodies. Methods of preparing polyclonal antibodies are known to the skilled artisan (Harlow, et al., *Antibodies: A Laboratory Manual*, (Cold spring Harbor Laboratory Press, 2nd
ed. (1988), which is hereby incorporated herein by reference in its entirety).
35 Polyclonal antibodies can be raised in a mammal or bird, for example, by one or more injections of an immunizing agent and, if desired, an adjuvant. For the purposes of the invention, “immunizing agent” may be defined as a polypeptide of the invention,

5 including fragments, variants, and/or derivatives thereof, in addition to fusions with heterologous polypeptides and other forms of the polypeptides described herein.

Typically, the immunizing agent and/or adjuvant will be injected in the mammal by multiple subcutaneous or intraperitoneal injections, though they may also be given intramuscularly, and/or through IV). The immunizing agent may include
10 polypeptides of the present invention or a fusion protein or variants thereof. Depending upon the nature of the polypeptides (i.e., percent hydrophobicity, percent hydrophilicity, stability, net charge, isoelectric point etc.), it may be useful to conjugate the immunizing agent to a protein known to be immunogenic in the mammal being immunized. Such conjugation includes either chemical conjugation
15 by derivitizing active chemical functional groups to both the polypeptide of the present invention and the immunogenic protein such that a covalent bond is formed, or through fusion-protein based methodology, or other methods known to the skilled artisan. Examples of such immunogenic proteins include, but are not limited to keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean
20 trypsin inhibitor. Various adjuvants may be used to increase the immunological response, depending on the host species, including but not limited to Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, dinitrophenol, and potentially useful human adjuvants
25 such as BCG (bacille Calmette-Guerin) and Corynebacterium parvum. Additional examples of adjuvants which may be employed includes the MPL-TDM adjuvant (monophosphoryl lipid A, synthetic trehalose dicorynomycolate). The immunization protocol may be selected by one skilled in the art without undue experimentation.

The antibodies of the present invention may comprise monoclonal antibodies.
30 Monoclonal antibodies may be prepared using hybridoma methods, such as those described by Kohler and Milstein, Nature, 256:495 (1975) and U.S. Pat. No. 4,376,110, by Harlow, et al., Antibodies: A Laboratory Manual, (Cold spring Harbor Laboratory Press, 2nd ed. (1988), by Hammerling, et al., Monoclonal Antibodies and T-Cell Hybridomas (Elsevier, N.Y., (1981)), or other methods known to the artisan.
35 Other examples of methods which may be employed for producing monoclonal antibodies includes, but are not limited to, the human B-cell hybridoma technique (Kosbor et al., 1983, Immunology Today 4:72; Cole et al., 1983, Proc. Natl. Acad. Sci. USA 80:2026-2030), and the EBV-hybridoma technique (Cole et al., 1985,

5 Monoclonal Antibodies And Cancer Therapy, Alan R. Liss, Inc., pp. 77-96). Such antibodies may be of any immunoglobulin class including IgG, IgM, IgE, IgA, IgD and any subclass thereof. The hybridoma producing the mAb of this invention may be cultivated in vitro or in vivo. Production of high titers of mAbs in vivo makes this the presently preferred method of production.

10 In a hybridoma method, a mouse, a humanized mouse, a mouse with a human immune system, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes may be immunized in vitro.

15 The immunizing agent will typically include polypeptides of the present invention or a fusion protein thereof. Generally, either peripheral blood lymphocytes ("PBLs") are used if cells of human origin are desired, or spleen cells or lymph node cells are used if non-human mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as
20 polyethylene glycol, to form a hybridoma cell (Goding, Monoclonal Antibodies: Principles and Practice, Academic Press, (1986), pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovine and human origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells may be cultured in a suitable culture medium that preferably
25 contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

30 Preferred immortalized cell lines are those that fuse efficiently, support stable high level expression of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium. More preferred immortalized cell lines are murine myeloma lines, which can be obtained, for instance, from the Salk Institute Cell Distribution Center, San Diego, California and the American Type Culture
35 Collection, Manassas, Virginia. As inferred throughout the specification, human myeloma and mouse-human heteromyeloma cell lines also have been described for the production of human monoclonal antibodies (Kozbor, J. Immunol., 133:3001

5 (1984); Brodeur et al., *Monoclonal Antibody Production Techniques and Applications*, Marcel Dekker, Inc., New York, (1987) pp. 51-63).

The culture medium in which the hybridoma cells are cultured can then be assayed for the presence of monoclonal antibodies directed against the polypeptides of the present invention. Preferably, the binding specificity of monoclonal antibodies
10 produced by the hybridoma cells is determined by immunoprecipitation or by an *in vitro* binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunoabsorbant assay (ELISA). Such techniques are known in the art and within the skill of the artisan. The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollart, *Anal.*
15 *Biochem.*, 107:220 (1980).

After the desired hybridoma cells are identified, the clones may be subcloned by limiting dilution procedures and grown by standard methods (Goding, *supra*). Suitable culture media for this purpose include, for example, Dulbecco's Modified Eagle's Medium and RPMI-1640. Alternatively, the hybridoma cells may be grown
20 *in vivo* as ascites in a mammal.

The monoclonal antibodies secreted by the subclones may be isolated or purified from the culture medium or ascites fluid by conventional immunoglobulin purification procedures such as, for example, protein A-sepharose, hydroxyapatite chromatography, gel exclusion chromatography, gel electrophoresis, dialysis, or
25 affinity chromatography.

The skilled artisan would acknowledge that a variety of methods exist in the art for the production of monoclonal antibodies and thus, the invention is not limited to their sole production in hybridomas. For example, the monoclonal antibodies may be made by recombinant DNA methods, such as those described in US patent No. 4, 816, 567. In this context, the term "monoclonal antibody" refers to an antibody
30 derived from a single eukaryotic, phage, or prokaryotic clone. The DNA encoding the monoclonal antibodies of the invention can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine
35 antibodies, or such chains from human, humanized, or other sources). The hybridoma cells of the invention serve as a preferred source of such DNA. Once isolated, the DNA may be placed into expression vectors, which are then transformed into host cells such as Simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells

5 that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. The DNA also may be modified, for example, by substituting the coding sequence for human heavy and light chain constant domains in place of the homologous murine sequences (US Patent No. 4, 816, 567; Morrison et al, supra) or by covalently joining to the immunoglobulin
10 coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. Such a non-immunoglobulin polypeptide can be substituted for the constant domains of an antibody of the invention, or can be substituted for the variable domains of one antigen-combining site of an antibody of the invention to create a chimeric bivalent antibody.

15 The antibodies may be monovalent antibodies. Methods for preparing monovalent antibodies are well known in the art. For example, one method involves recombinant expression of immunoglobulin light chain and modified heavy chain. The heavy chain is truncated generally at any point in the Fc region so as to prevent heavy chain crosslinking. Alternatively, the relevant cysteine residues are substituted
20 with another amino acid residue or are deleted so as to prevent crosslinking.

In vitro methods are also suitable for preparing monovalent antibodies. Digestion of antibodies to produce fragments thereof, particularly, Fab fragments, can be accomplished using routine techniques known in the art.

The antibodies of the present invention may further comprise humanized
25 antibodies or fully-human (human) antibodies. Humanized forms of non-human (e.g., murine) antibodies are chimeric immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. Humanized antibodies include human immunoglobulins
30 (recipient antibody) in which residues from a complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat, or rabbit having the desired specificity, affinity and capacity. In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Humanized
35 antibodies may also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to

5 those of a non-human immunoglobulin and all or substantially all of the FR regions are those of a human immunoglobulin consensus sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones et al., *Nature*, 321:522-525 (1986); Riechmann et al., *Nature* 332:323-329 (1988)] and Presta, *Curr. Op. Struct. Biol.*, 2:593-596 (1992).

10 Methods for humanizing non-human antibodies are well known in the art. Generally, a humanized antibody has one or more amino acid residues introduced into it from a source that is non-human. These non-human amino acid residues are often referred to as "import" residues, which are typically taken from an "import" variable domain. Humanization can be essentially performed following the methods of Winter and co-workers (Jones et al., *Nature*, 321:522-525 (1986); Reichmann et al., *Nature*, 332:323-327 (1988); Verhoeyen et al., *Science*, 239:1534-1536 (1988), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. Accordingly, such "humanized" antibodies are chimeric antibodies 15 (US Patent No. 4, 816, 567), wherein substantially less than an intact human variable domain has been substituted by the corresponding sequence from a non-human species. In practice, humanized antibodies are typically human antibodies in which some CDR residues and possible some FR residues are substituted from analogous sites in rodent antibodies.

25 Human antibodies can also be produced using various techniques known in the art, including phage display libraries (Hoogenboom and Winter, *J. Mol. Biol.*, 227:381 (1991); Marks et al., *J. Mol. Biol.*, 222:581 (1991)). The techniques of Cole et al., and Boerder et al., are also available for the preparation of human monoclonal antibodies (Cole et al., *Monoclonal Antibodies and Cancer Therapy*, Alan R. Riss, 30 (1985); and Boerner et al., *J. Immunol.*, 147(1):86-95, (1991)). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in all respects, including gene 35 rearrangement, assembly, and creation of an antibody repertoire. This approach is described, for example, in US patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5,633,425; 5,661,106, and in the following scientific publications: Marks et al., *Biotechnol.*, 10:779-783 (1992); Lonberg et al., *Nature* 368:856-859 (1994);

- 5 Fishwild et al., *Nature Biotechnol.*, 14:845-51 (1996); Neuberger, *Nature Biotechnol.*, 14:826 (1996); Lonberg and Huszer, *Intern. Rev. Immunol.*, 13:65-93 (1995).

In addition, techniques developed for the production of "chimeric antibodies" (Morrison et al., 1984, *Proc. Natl. Acad. Sci.*, 81:6851-6855; Neuberger et al., 1984, *Nature*, 312:604-608; Takeda et al., 1985, *Nature*, 314:452-454) by splicing the genes
10 from a mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity can be used. A chimeric antibody is a molecule in which different portions are derived from different animal species, such as those having a variable region derived from a murine mAb and a human immunoglobulin constant region. (See, e.g., Cabilly et al., U.S. Pat.
15 No. 4,816,567; and Boss et al., U.S. Pat. No. 4,816,397, both of which are incorporated herein by reference in their entirety.)

The antibodies of the present invention may be bispecific antibodies. Bispecific antibodies are monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens. In the present
20 invention, one of the binding specificities may be directed towards a polypeptide of the present invention, the other may be for any other antigen, and preferably for a cell-surface protein, receptor, receptor subunit, tissue-specific antigen, virally derived protein, virally encoded envelope protein, bacterially derived protein, or bacterial surface protein, etc.

25 Methods for making bispecific antibodies are known in the art. Traditionally, the recombinant production of bispecific antibodies is based on the co-expression of two immunoglobulin heavy-chain/light-chain pairs, where the two heavy chains have different specificities (Milstein and Cuello, *Nature*, 305:537-539 (1983)). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas
30 (quadromas) produce a potential mixture of ten different antibody molecules, of which only one has the correct bispecific structure. The purification of the correct molecule is usually accomplished by affinity chromatography steps. Similar procedures are disclosed in WO 93/08829, published 13 May 1993, and in Traunecker et al., *EMBO J.*, 10:3655-3659 (1991).

35 Antibody variable domains with the desired binding specificities (antibody-antigen combining sites) can be fused to immunoglobulin constant domain sequences. The fusion preferably is with an immunoglobulin heavy-chain constant domain, comprising at least part of the hinge, CH2, and CH3 regions. It is preferred to have

5 the first heavy-chain constant region (CH1) containing the site necessary for light-chain binding present in at least one of the fusions. DNAs encoding the immunoglobulin heavy-chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transformed into a suitable host organism. For further details of generating bispecific antibodies see, for example
10 Suresh et al., *Meth. In Enzym*, 121:210 (1986).

Heteroconjugate antibodies are also contemplated by the present invention. Heteroconjugate antibodies are composed of two covalently joined antibodies. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells (US Patent No. 4, 676, 980), and for the treatment of HIV infection
15 (WO 91/00360; WO 92/20373; and EP03089). It is contemplated that the antibodies may be prepared in vitro using known methods in synthetic protein chemistry, including those involving crosslinking agents. For example, immunotoxins may be constructed using a disulfide exchange reaction or by forming a thioester bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-
20 mercaptobutyrimidate and those disclosed, for example, in US Patent No. 4,676,980.

Uses for Antibodies directed against polypeptides of the invention

The antibodies of the present invention have various utilities. For example, such antibodies may be used in diagnostic assays to detect the presence or
25 quantification of the polypeptides of the invention in a sample. Such a diagnostic assay may be comprised of at least two steps. The first, subjecting a sample with the antibody, wherein the sample is a tissue (e.g., animal, plant, etc.), biological fluid (e.g., blood, urine, phloem fluid, xylem fluid, plant secretion, etc.), biological extract (e.g., tissue or cellular homogenate, etc.), a protein microchip (e.g., See Arenkov P, et
30 al., *Anal Biochem.*, 278(2):123-131 (2000)), or a chromatography column, etc. And a second step involving the quantification of antibody bound to the substrate. Alternatively, the method may additionally involve a first step of attaching the antibody, either covalently, electrostatically, or reversably, to a solid support, and a second step of subjecting the bound antibody to the sample, as defined above and
35 elsewhere herein.

Various diagnostic assay techniques are known in the art, such as competitive binding assays, direct or indirect sandwich assays and immunoprecipitation assays conducted in either heterogenous or homogenous phases (Zola, *Monoclonal*

5 Antibodies: A Manual of Techniques, CRC Press, Inc., (1987), pp147-158). The
antibodies used in the diagnostic assays can be labeled with a detectable moiety. The
detectable moiety should be capable of producing, either directly or indirectly, a
detectable signal. For example, the detectable moiety may be a radioisotope, such as
2H, 14C, 32P, or 125I, a florescent or chemiluminescent compound, such as
10 fluorescein isothiocyanate, rhodamine, or luciferin, or an enzyme, such as alkaline
phoshatase, beta-galactosidase, green flourescent protein, or horseradish peroxidase.
Any method known in the art for conjugating the antibody to the detectable moiety
may be employed, including those methods described by Hunter et al., Nature,
144:945 (1962); Dafvid et al., Biochem., 13:1014 (1974); Pain et al., J. Immunol.
15 Metho., 40:219(1981); and Nygren, J. Histochem. And Cytochem., 30:407 (1982).

Antibodies directed against the polypeptides of the present invention are
useful for the affinity purification of such polypeptides from recombinant cell culture
or natural sources. In this process, the antibodies against a particular polypeptide are
immobilized on a suitable support, such as a Sephadex resin or filter paper, using
20 methods well known in the art. The immobilized antibody then is contacted with a
sample containing the polypeptides to be purified, and thereafter the support is
washed with a suitable solvent that will remove substantially all the material in the
sample except for the desired polypeptides, which are bound to the immobilized
antibody. Finally, the support is washed with another suitable solvent that will release
25 the desired polypeptide from the antibody.

Antibodies directed against polypeptides of the present invention are useful for
inhibiting allergic reactions in animals. For example, by administering a
therapeutically acceptable dose of an antibody, or antibodies, of the present invention,
or a cocktail of the present antibodies, or in combination with other antibodies of
30 varying sources, the animal may not elicit an allergic response upon ingestion of
certain antigens, particularly plant antigens. In an exemplary example, the tomato
LEA lectin, from *Lycopersicum esculentum* has been shown to elicit a strong
immune response when administered both orally and i.n. in mice. Thus, the
administration to the mouse of antibodies directed against the LEA lectin may
35 decrease or altogether eliminate an immune response to the LEA lectin as the LEA
lectin could effectively be eliminated from circulation prior to eliciting an immune
response (for example, See Lavelle EC, et al., Immunology. 99(1):30-7, (2000)).
Likewise, one could envision cloning the gene encoding an antibody directed against

5 a polypeptide of the present invention, said polypeptide having the potential to elicit an allergic and/or immune response in an organism, and transforming the organism with said antibody gene such that it is expressed (e.g., constitutively, inducibly, etc.) in the organism. Thus, the organism would effectively become resistant to an allergic response resulting from the ingestion or presence of such an immune/allergic reactive
10 polypeptide. Detailed descriptions of therapeutic and/or gene therapy applications of the present invention are provided elsewhere herein.

Alternatively, antibodies of the present invention could be produced in a plant (e.g., cloning the gene of the antibody directed against a polypeptide of the present invention, and transforming a plant with a suitable vector comprising said gene for
15 constitutive expression of the antibody within the plant), and the plant subsequently ingested by an animal, thereby conferring temporary immunity to the animal for the specific antigen the antibody is directed towards (See, for example, US Patent Nos. 5,914,123 and 6,034,298).

In another embodiment, antibodies of the present invention, preferably
20 polyclonal antibodies, more preferably monoclonal antibodies, and most preferably single-chain antibodies, can be used as a means of inhibiting gene expression of a particular gene, or genes, in a plant or organism. See, for example, International Publication Number WO 00/05391, published 2/3/00, to Dow Agrosiences LLC. The application of such methods for the antibodies of the present invention are known
25 in the art, and are more particularly described elsewhere herein (See, Examples 14 and 15).

In yet another embodiment, antibodies of the present invention may be useful for multimerizing the polypeptides of the present invention. For example, certain proteins may confer enhanced biological activity when present in a multimeric state
30 (i.e., such enhanced activity may be due to the increased effective concentration of such proteins whereby more protein is available in a localized location).

Fusion Proteins

Any polypeptide of the present invention can be used to generate fusion
35 proteins. For example, the polypeptide of the present invention, when fused to a second protein, can be used as an antigenic tag. Antibodies raised against the polypeptide of the present invention can be used to indirectly detect the second protein by binding to the polypeptide. Moreover, because certain proteins target

5 cellular locations based on trafficking signals, the polypeptides of the present invention can be used as targeting molecules once fused to other proteins.

Examples of domains that can be fused to polypeptides of the present invention include not only heterologous signal sequences, but also other heterologous functional regions. The fusion does not necessarily need to be direct, but may occur
10 through linker sequences.

Moreover, fusion proteins may also be engineered to improve characteristics of the polypeptide of the present invention. For instance, a region of additional amino acids, particularly charged amino acids, may be added to the N-terminus of the polypeptide to improve stability and persistence during purification from the host cell
15 or subsequent handling and storage. Peptide moieties may be added to the polypeptide to facilitate purification. Such regions may be removed prior to final preparation of the polypeptide. Similarly, peptide cleavage sites can be introduced inbetween such peptide moieties, which could additionally be subjected to protease activity to remove said peptide(s) from the protein of the present invention. The addition of peptide
20 moieties, including peptide cleavage sites, to facilitate handling of polypeptides are familiar and routine techniques in the art.

Moreover, polypeptides of the present invention, including fragments, and specifically epitopes, can be combined with parts of the constant domain of immunoglobulins (IgA, IgE, IgG, IgM) or portions thereof (CH1, CH2, CH3, and any
25 combination thereof, including both entire domains and portions thereof), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life in vivo. One reported example describes chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian
30 immunoglobulins. (EP A 394,827; Traunecker et al., Nature 331:84-86 (1988).) Fusion proteins having disulfide-linked dimeric structures (due to the IgG) can also be more efficient in binding and neutralizing other molecules, than the monomeric secreted protein or protein fragment alone. (Fountoulakis et al., J. Biochem. 270:3958-3964 (1995).)

35 Similarly, EP-A-O 464 533 (Canadian counterpart 2045869) discloses fusion proteins comprising various portions of the constant region of immunoglobulin molecules together with another human protein or part thereof. In many cases, the Fc part in a fusion protein is beneficial in therapy and diagnosis, and thus can result in,

5 for example, improved pharmacokinetic properties. (EP-A 0232 262.) Alternatively, deleting the Fc part after the fusion protein has been expressed, detected, and purified, would be desired. For example, the Fc portion may hinder therapy and diagnosis if the fusion protein is used as an antigen for immunizations. In drug discovery, for example, human proteins, such as hIL-5, have been fused with Fc portions for the
10 purpose of high-throughput screening assays to identify antagonists of hIL-5. (See, D. Bennett et al., *J. Molecular Recognition* 8:52-58 (1995); K. Johanson et al., *J. Biol. Chem.* 270:9459-9471 (1995).)

Moreover, the polypeptides of the present invention can be fused to marker sequences (also referred to as "tags"). Due to the availability of antibodies specific to
15 such "tags", purification of the fused polypeptide of the invention, and/or its identification is significantly facilitated since antibodies specific to the polypeptides of the invention are not required. Such purification may be in the form of an affinity purification whereby an anti-tag antibody or another type of affinity matrix (e.g., anti-tag antibody attached to the matrix of a flow-thru column) that binds to the epitope
20 tag is present. In preferred embodiments, the marker amino acid sequence is a hexa-histidine peptide, such as the tag provided in a pQE vector (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311), among others, many of which are commercially available. As described in Gentz et al., *Proc. Natl. Acad. Sci. USA* 86:821-824 (1989), for instance, hexa-histidine provides for convenient purification of the fusion
25 protein. Another peptide tag useful for purification, the "HA" tag, corresponds to an epitope derived from the influenza hemagglutinin protein. (Wilson et al., *Cell* 37:767 (1984)).

The skilled artisan would acknowledge the existence of other "tags" which could be readily substituted for the tags referred to supra for purification and/or
30 identification of polypeptides of the present invention (Jones C., et al., *J Chromatogr A.* 707(1):3-22 (1995)). For example, the c-myc tag and the 8F9, 3C7, 6E10, G4m B7 and 9E10 antibodies thereto (Evan et al., *Molecular and Cellular Biology* 5:3610-3616 (1985)); the Herpes Simplex virus glycoprotein D (gD) tag and its antibody (Paborsky et al., *Protein Engineering*, 3(6):547-553 (1990), the Flag-peptide – i.e., the
35 octapeptide sequence DYKDDDDK (SEQ ID NO:8), (Hopp et al., *Biotech.* 6:1204-1210 (1988); the KT3 epitope peptide (Martin et al., *Science*, 255:192-194 (1992)); a-tubulin epitope peptide (Skinner et al., *J. Biol. Chem.*, 266:15136-15166, (1991)); the T7 gene 10 protein peptide tag (Lutz-Freyermuth et al., *Proc. Natl. Sci. USA*,

5 87:6363-6397 (1990)), the FITC epitope (Zymed, Inc.), the GFP epitope (Zymed, Inc.), and the Rhodamine epitope (Zymed, Inc.).

The present invention also encompasses the attachment of the coding region of a repeating series of up to nine arginine amino acids to a polynucleotide of the present invention. The invention also encompasses chemically derivitizing a
10 polypeptide of the present invention with a repeating series of up to nine arginine amino acids. Such a tag, when attached to a polypeptide, has recently been shown to serve as a universal pass, allowing compounds access to the interior of cells without additional derivitization or manipulation (Wender, P., et al., unpublished data).

Protein fusions involving polypeptides of the present invention, including
15 fragments and/or variants thereof, can be used for the following, non-limiting examples, subcellular localization of proteins, determination of protein-protein interactions via immunoprecipitation, purification of proteins via affinity chromatography, functional and/or structural characterization of protein. The present invention also encompasses the application of hapten specific antibodies for any of
20 the uses referenced above for epitope fusion proteins. For example, the polypeptides of the present invention could be chemically derivitized to attach hapten molecules (e.g., DNP, (Zymed, Inc.)). Due to the availability of monoclonal antibodies specific to such haptens, the protein could be readily purified using immunoprecipitation, for example.

25 Polypeptides of the present invention, including fragments and/or variants thereof, in addition to, antibodies directed against such polypeptides, fragments, and/or variants, may be fused to any of a number of known, and yet to be determined, toxins, such as ricin, saporin (Mashiba H, et al., Ann N Y Acad Sci. 1999;886:233-5), HC toxin (Tonukari NJ, et al., Plant Cell. 2000 Feb;12(2):237-248), BT endotoxin, or
30 pseudomonias endotoxin. Such fusions could be used to deliver the toxins to desired tissues for which a ligand or a protein capable of binding to the polypeptides of the invention exists.

The invention encompasses the fusion of antibodies directed against polypeptides of the present invention, including variants and fragments thereof, to
35 said toxins for delivering the toxin to specific locations in a cell, to specific plant tissues, and/or to specific plant species. Such bifunctional antibodies are known in the art, though a review describing additional advantageous fusions, including citations for methods of production, can be found in P.J. Hudson, Curr. Opp. In Imm.

5 11:548-557, (1999); this publication, in addition to the references cited therein, are hereby incorporated by reference in their entirety herein. In this context, the term “toxin” may be expanded to include any heterologous protein, a small molecule, radionucleotides, cytotoxic drugs, liposomes, adhesion molecules, glycoproteins, ligands, cell or tissue-specific ligands, enzymes, of bioactive agents, biological
10 response modifiers, anti-fungal agents, hormones, steroids, vitamins, peptides, peptide analogs, anti-allergenic agents, anti-tubercular agents, anti-viral agents, antibiotics, anti-protozoan agents, chelates, radioactive particles, radioactive ions, X-ray contrast agents, monoclonal antibodies, polyclonal antibodies and genetic material. In view of the present disclosure, one skilled in the art could determine whether any particular
15 “toxin could be used in the compounds of the present invention. Examples of suitable “toxins” listed above are exemplary only and are not intended to limit the “toxins” that may be used in the present invention.

Thus, any of these above fusions can be engineered using the polynucleotides or the polypeptides of the present invention.

20

Vectors, Host Cells, and Protein Production

The present invention also relates to vectors containing the polynucleotide of the present invention, host cells, and the production of polypeptides by recombinant techniques. The vector may be, for example, a phage, plasmid, viral, or retroviral
25 vector. Retroviral vectors may be replication competent or replication defective. In the latter case, viral propagation generally will occur only in complementing host cells.

The polynucleotides may be joined to a vector containing a selectable marker for propagation in a host. Generally, a plasmid vector is introduced in a precipitate,
30 such as a calcium phosphate precipitate, or in a complex with a charged lipid. If the vector is a virus, it may be packaged in vitro using an appropriate packaging cell line and then transduced into host cells.

The polynucleotide insert of the present invention should be operatively linked to an appropriate promoter, such as the 35S promoter, the 34S promoter, CMV
35 promoter, phage lambda PL promoter, the E. coli lac, trp, phoA and tac promoters, the SV40 early and late promoters and promoters of retroviral LTRs, to name a few. In addition, it may be desirable, or required, in some instances to have tissue-specific or cell type-specific promoters operably linked to a polynucleotide of the present

5 invention. Examples of suitable plant-expressible promoters selectively expressed in particular tissues or cell types are well known in the art and include, but are not limited to, seed specific promoters (e.g., WO 89/03887), organ-primordia specific promoters (An et al., *Plant Cell*, 8:15-30, (1996)), stem-specific promoters (Keller et al., *EMBO J.*, 7:3625-3633, (1988)), leaf specific promoters (Hudspeth et al., *Plant*
10 *Mol. Biol.*, 12:579-589, (1989)), mesophyl-specific promoters (such as the light inducible Rubisco promoters), root-specific promoters (Keller et al., *Genes Devel.*, 3:1639-1646, (1989)), tuber-specific promoters (Keil et al., *EMBO J.*, 8:1323-1330, (1989)), vascular tissue specific promoters (Peleman et al., *Gene*, 84:359-369, (1989)), meristem specific promoters (such as the promoter of the
15 SHOOTMERISTEMLESS (STM) gene, Long, et al., *Nature*, 379:66-69, (1996)), primordia specific promoter (such as the *Antirrhinum CycD3a* gene promoter, Doonan et al., in "Plant Cell Division" (Francis, Duditz, and Inze, Eds), Portland Press, London, (1998)), anther specific promoters (WO 89/10396, WO 92/13956, and WO 92/13957), stigma-specific promoters (WO 91/02068), degiscence-zone specific
20 promoters (WO 97/13865), seed-specific promoters (WO 89/03887), etc.

Additional promoters that may be operably linked to a polynucleotide of the present invention may be found in McElroy and Brettel, *Tibtech*, Vol. 12, February, 1994. Moreover, a number of promoters are currently being used for transformation of dicotyledonous plants. These promoters come from a variety of different sources. One
25 group of commonly used promoters were isolated from *Agrobacterium tumefaciens*, where they function to drive the expression of opine synthase genes carried on the T-DNA segment that is integrated into the plant genome during infection. These promoters include the octopine synthase (ocs) promoter (L. Comai et al., 1985; C. Waldron et al., 1985), the mannopine synthase (mas) promoter (L. Comai et al., 1985;
30 K. E. McBride and K. R. Summerfelt, 1990) and the nopaline synthase (nos) promoter (M. W. Bevan et al., 1983; L. Herrera-Estrella et al., 1983, R. T. Fraley et al., 1983, M. De Block et al., 1984;, R. Hain et al., 1985). These promoters are active in a wide variety of plant tissue.

In addition, the promoters disclosed in the following publications may also be
35 operably linked to a polynucleotide of the present invention: US Patent Nos. 5,623,067; 5,712,112; 5,723,751; 5,723,754; 5,723,757; 5,744,334; 5,750,385; 5,750,399; 5,767,363; 5,783,393; 5,789,214; 5,792,922; 5,792,933; 5,801,027; 5,804,694; 5,814,618; 5,824,857; 5,824,863; 5,824,865; 5,824,866; 5,824,872; and

5 5,929,302; and International Publication Nos. WO 97/49727, WO 98/00533, WO 98/03655, WO 98/07846, WO 98/08961, WO 98/08962, WO 98/10734, WO 98/16634, WO 98/22593, WO 98/38295, and WO 98/44097; and European Patent Application No. EP 0 846 770.

Several viral promoters are also used to, drive heterologous gene expression in
10 dicots (J. C. Kridl and R. M. Goodman, 1986) and may be operably linked to a polynucleotide of the present invention. The Cauliflower Mosaic Virus 35S promoter is one of the promoters used most often for dicot transformation because it confers high levels of gene expression in almost all tissues (J. Odell et al., 1985; D. W. Ow et al., 1986; D. M. Shah et al., 1986). Modifications of this promoter are also used,
15 including a configuration with two tandem 35S promoters (R. Kay et al., 1987) and the mas-35S promoter (L. Comai et al., 1990), which consists of the mannopine synthase promoter in tandem with the 35S promoter. Both of these promoters drive even higher levels of gene expression than a single copy of the 35S promoter. Other viral promoters that have been used include the Cauliflower Mosaic Virus 19S promoter (J.
20 Paszkowski et al., 1984; E. Balazs et al.) and the 34S promoter from the figwort mosaic virus (M. Sanger et al., 1990).

Alternatively, the polynucleotide insert of the present invention could be operatively linked to any of a number of inducible promoters known in the art, which include, but are not limited to: tetracycline inducible promoters, small-molecule
25 inducible promoters, light inducible promoters, chemical compounds (e.g., safeners, herbicides, glucocorticoids, etc.), abiotic stress inducible promoters (e.g., wounding, heavy metals, cold-sensitive promoters, heat-sensitive promoters, salt sensitive promoters, drought sensitive promoters, hypoxia inducible (such as those disclosed in EP1012317), etc.), biotic stress promoters (e.g., pathogen or pest infection including
30 infection by fungi, viruses, bacteria, insects, nematodes, mycoplasmas, and mycoplasma-like organisms, etc.). Examples of plant-expressible inducible promoters suitable for the invention are: nematode inducible promoters (such as those disclosed in WO 92/21757 and/or EP1007709), fungus inducible promoters (WO 93/19188, WO 96/28561), chemically inducible Arabidopsis PR-1 promoter (WO 98/03536), the
35 inducible promoters disclosed in WO 98/45445, the inducible promoters disclosed in US Patent No. 5,804,693, the tomato soft fruit inducible promoter disclosed in US Patent No. 5, 821, 398, promoters inducible after application of glucocorticoids such as dexamethasone, or promoters repressed or activated after application of tetracyclin

5 (Gatz et al., PNAS USA, 85:1394-1397, (1988)). Other suitable inducible promoters will be known to the skilled artisan.

In addition, the polynucleotide insert of the present invention could be operatively linked to "artificial" or chimeric promoters and transcription factors. Specifically, the artificial promoter could comprise, or alternatively consist, of any
10 combination of cis-acting DNA sequence elements that are recognized by trans-acting transcription factors. Preferably, the cis acting DNA sequence elements and trans-acting transcription factors are operable in plants. Further, the trans-acting transcription factors of such "artificial" promoters could also be "artificial" or chimeric in design themselves and could act as activators or repressors to said
15 "artificial" promoter. For example, a chimeric promoter of the invention could comprise one or more, upstream activating sequences from the Octopine Synthase gene (OCS), matrix attachment regions (MAR), etc.

The expression constructs will further contain sites for transcription initiation, termination, and, in the transcribed region, a ribosome binding site for translation.
20 The coding portion of the transcripts expressed by the constructs will preferably include a translation initiating codon at the beginning and a termination codon (UAA, UGA or UAG) appropriately positioned at the end of the polypeptide to be translated.

The expression constructs may additionally comprise 5' leader sequences in the expression constructs. Such leader sequences can act to enhance translation.
25 Translation leaders are known in the art and include: picornavirus leaders, for example, EMCV leader (Encephalomyocarditis 5' noncoding region) (Elroy-Stein, O., Fuerst, T.R., and Moss, B. (1989) PNAS USA, 86:6126-6130); potyvirus leaders, for example, TEV leader (Tobacco Etch Virus) (Allison et al. (1986)); MDMV leader (Maize Dwarf Mosaic Virus) Virology, 154:9-20); and human immunoglobulin
30 heavy-chain binding protein (BiP), (Macejak, D.G., and Sarnow, P. (1991) Nature, 353:90-94; untranslated leader from the coat protein mRNA of alfalfa mosaic virus (AMV RNA 4), (Jobling, S.A., and Gehrke, L., (1987) Nature, 325:622-625); tobacco mosaic virus leader (TW), (Gallie, D.R. et al. (1989) Molecular Biology of RNA, pages 237-256); and maize chlorotic mottle virus leader (MCNW) (Lommel, S.A. et
35 al. (1991) Virology, 81:382-385). See also, Della-Cioppa et al. (1987) Plant Physiology, 81:965-966. Other methods known to enhance translation can also be utilized, for example, introns, and the like.

5 As indicated, the expression vectors will preferably include at least one
selectable marker. Such markers include, but are not limited to, dihydrofolate
reductase, G418 or neomycin resistance, kanamycin resistance, hygromycin
resistance, bialaphos resistance, sulfonoamide resistance, streptomycin resistance,
10 spectinomycin resistance, chlorosulfuron resistance, glyphosphate resistance, and
methotrexate resistance, for eukaryotic cell culture and tetracycline, kanamycin or
ampicillin resistance genes for culturing in *E. coli* and other bacteria. Representative
examples of appropriate hosts include, but are not limited to, bacterial cells, such as *E.*
coli, *Streptomyces* and *Salmonella typhimurium* cells; fungal cells, such as yeast cells
(e.g., *Saccharomyces cerevisiae* or *Pichia pastoris* (ATCC Accession No. 201178));
15 insect cells such as *Drosophila* S2 and *Spodoptera Sf9* cells; animal cells such as
CHO, COS, 293, and Bowes melanoma cells; plant cells, and specifically plant cells
and/or tissues derived from any of the plants listed in Table 3. Appropriate culture
mediums and conditions for the above-described host cells are known in the art.

The polynucleotides and polypeptides of the present invention can be targeted
20 to the chloroplast or amyloplast for expression. In this manner, the expression
construct will additionally contain a polynucleotide sequence encoding a transit
peptide operably linked to a polynucleotide of the present invention to direct the
polynucleotide of the present invention to the chloroplasts. Such transit peptides are
known in the art. See, for example, Von Heijne et al. (1991) *Plant Mol. Biol. Rep.*
25 9:104-126; Clark et al. (1989) *J Biol. Chem.* 264:17544-17550; della-Cioppa et al.
(1987) *Plant Physiol.* 84065060 R.omer et al. (1993) *Biochem. Biophys. Res*
Commun. 196:1414-1421; and Shah et al. (1986) *Science* 233:478-481.

The expression construct may also comprise any other necessary regulators
such as nuclear localization signals (Kalderon et al. (1984) *Cell* 39:499-509; and
30 Lassner et al. (1991) *Plant Molecular Biology* 17:229-234); plant translational
consensus sequences (Joshi, C.P. (1987) *Nucleic Acids Research* 15:6643 6653),
introns (Luehrsen and Walbot (1991) *Mol. Gen. Genet.* 225:81-93) and the like,
operably linked to a polynucleotide of the present invention.

The polynucleotide sequences encoding the proteins or polypeptides of the
35 present invention may be particularly useful in the genetic manipulation of plants. In
this manner, the polynucleotides of the invention are provided in expression cassettes
for expression in the plant of interest. Where appropriate, the gene(s) may be
optimized for increased expression in the transformed plant. That is, the

5 polynucleotides can be synthesized using plant preferred codons for improved expression specific to a particular species. Methods are available in the art for synthesizing plant preferred genes. See, for example, U.S. Patent Nos. 5,380,831, 5,436, 391, and Murray et al. (1989) *Nucleic Acids Res.* 17:477-498, herein incorporated by reference.

10 Depending upon the species in which the DNA sequence of interest is to be expressed, it may be desirable to synthesize the sequence with plant preferred codons, or alternatively with chloroplast preferred codons. The plant preferred codons may be determined from the codons of highest frequency in the proteins expressed in the largest amount in the particular plant species of interest. See, EPA 0359472; EPA
15 0385962; WO 91/16432; Perlak et al. (1991) *Proc. Natl. Acad. Sci. USA* 88:3324-3328; and Murray et al. (1989) *Nucleic Acids Research*. In this manner, the polynucleotide sequences can be optimized for expression in any plant. It is recognized that all or any part of the gene sequence may be optimized or synthetic. That is, synthetic or partially optimized sequences may also be used.

20 Additionally, it may be desirable to selectively express a polypeptide of the present invention in a specific target cell or tissue of a plant by synthesizing the encoding polynucleotide sequence to contain codons optimized for high translational efficiency within the particular target cell or tissue. Such methods are known in the art and are specifically provided in PCT International Publication No. WO 00/42190
25 (which is hereby incorporated herein by reference).

Additional sequence modifications are known to enhance gene expression in a cellular host. These include elimination of sequences encoding spurious polyadenylation signals, exon-intron splice site signals, transposon like repeats, and other such well-characterized sequences that may be deleterious to gene expression.
30 The G-C content of the sequence may be adjusted to levels average for a given cellular host, as calculated by reference to known genes expressed in the host cell. When possible, the sequence may be modified to avoid predicted hairpin secondary mRNA structures.

Among vectors preferred for use in bacteria include pQE70, pQE60 and pQE-
35 9, available from QIAGEN, Inc.; pBluescript vectors, Phagescript vectors, pNH8A, pNH16a, pNH18A, pNH46A, available from Stratagene Cloning Systems, Inc.; and ptc99a, pKK223-3, pKK233-3, pDR540, pRIT5 available from Pharmacia Biotech, Inc. Among preferred eukaryotic vectors are pWLNEO, pSV2CAT, pOG44, pXT1

5 and pSG available from Stratagene; and pSVK3, pBPV, pMSG and pSVL available from Pharmacia. Preferred expression vectors for use in yeast systems include, but are not limited to pYES2, pYD1, pTEF1/Zeo, pYES2/GS, pPICZ,pGAPZ, pGAPZalph, pPIC9, pPIC3.5, pHIL-D2, pHIL-S1, pPIC3.5K, pPIC9K, and PAO815 (all available from Invitrogen, Carlsbad, CA).

10 Preferred expression vectors in plant systems include, but are not limited to, Bin 19 (ATCC Deposit No: 37327), GA437 (ATCC Deposit No: 37350), pAK1003 (ATCC Deposit No: 37425), pAS2022 (ATCC Deposit No: 37426), pAS2023 (ATCC Deposit No: 37427), pAP2034 (ATCC Deposit No: 37428), pC22 (ATCC Deposit No: 37493), pHS24 (ATCC Deposit No: 37841), pHS85 (ATCC Deposit No: 37842),
15 pPM1 (ATCC Deposit No: 40172), pGV3111SE (ATCC Deposit No: 53213), pCGN978 (ATCC Deposit No: 67064), pFL61 (ATCC Deposit No: 77215), pGPTV-KAN (ATCC Deposit No: 77388), pGPTV-HPT (ATCC Deposit No: 77389), pGPTV-DHFR (ATCC Deposit No: 77390), pGPTV-BAR (ATCC Deposit No: 77391), pGPTV-BLEO (ATCC Deposit No: 77392), and/or pPE1000 (ATCC Deposit
20 No: 87573). The skilled artisan would appreciate that any of the above vectors could easily be modified to either include or delete specific elements as may be required for operability. Other suitable vectors will be readily apparent to the skilled artisan.

Introduction of the construct into the host cell can be effected by biolistic transformation (Klein et al., *Nature*, 327:70-73 (1987)), PEG-mediated transfection
25 (Paskowski, et al., *EMBO J.*, 3:2717, (1984)), calcium phosphate transfection, DEAE-dextran mediated transfection, cationic lipid-mediated transfection, electroporation (Fromm, et al., *PNAS*, USA, 82:5824 (1985)), transduction, infection, *Agrobacterium tumefaciens* -directed infection, or other methods. Such methods are described in many standard laboratory manuals, such as Davis et al., *Basic Methods in Molecular*
30 *Biology* (1986).

A polypeptide of this invention can be recovered and purified from recombinant cell cultures by well-known methods including ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, affinity
35 chromatography, hydroxylapatite chromatography and lectin chromatography. Most preferably, high performance liquid chromatography ("HPLC") is employed for purification.

5 Polypeptides of the present invention can also be recovered from: products purified from natural sources, including bodily fluids, tissues and cells, whether directly isolated or cultured; products of chemical synthetic procedures; and products produced by recombinant techniques from a prokaryotic or eukaryotic host, including, for example, bacterial, yeast, higher plant, insect, and mammalian cells.

10 Depending upon the host employed in a recombinant production procedure, the polypeptides of the present invention may be glycosylated or may be non-glycosylated. In addition, polypeptides of the invention may also include an initial modified methionine residue, in some cases as a result of host-mediated processes. Thus, it is well known in the art that the N-terminal methionine encoded by the translation initiation codon generally is removed with high efficiency from any
15 protein after translation in all eukaryotic cells. While the N-terminal methionine on most proteins also is efficiently removed in most prokaryotes, for some proteins, this prokaryotic removal process is inefficient, depending on the nature of the amino acid to which the N-terminal methionine is covalently linked.

20 In addition to encompassing host cells containing the vector constructs discussed herein, the invention also encompasses primary, secondary, and immortalized host cells that have been engineered to delete or replace endogenous genetic material (e.g., coding sequence), and/or to include genetic material (e.g., heterologous polynucleotide sequences) that is operably associated with the polynucleotides of the invention, and which activates, alters, and/or amplifies
25 endogenous polynucleotides. For example, techniques known in the art may be used to operably associate heterologous control regions (e.g., promoter and/or enhancer) and endogenous polynucleotide sequences via homologous recombination, resulting in the formation of a new transcription unit (see, e.g., U.S. Patent No. 5,641,670, issued June 24, 1997; U.S. Patent No. 5,733,761, issued March 31, 1998; International
30 Publication No. WO 96/29411, published September 26, 1996; International Publication No. WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA 86:8932-8935 (1989); and Zijlstra et al., Nature 342:435-438 (1989), the disclosures of each of which are incorporated by reference in their entireties).

35 In addition, polypeptides, analogs, derivatives, and/or fragments of the invention can be chemically synthesized. (See, e.g., Merrifield, 1963, J. Amer. Chem. Soc. 85:2149-2156). For example, polypeptides can be synthesized by solid phase techniques, cleaved from the resin, and purified by preparative high performance

5 liquid chromatography (e.g., see Creighton, 1983, *Proteins, Structures and Molecular Principles*, W.H. Freeman and Co., N.Y., pp. 50-60). The polypeptides can also be synthesized by use of a peptide synthesizer. The composition of the synthetic peptides may be confirmed by amino acid analysis or sequencing (e.g., the Edman degradation procedure; see Creighton, 1983, *Proteins, Structures and Molecular Principles*, W.H. Freeman and Co., N.Y., pp. 34-49; Mass Spectroscopy peptide sequencing, etc).
10 Furthermore, it desired, nonclassical amino acids or chemical amino acid analogs can be introduced as a substitution or addition into the polypeptide of the invention. Non-classical amino acids include but are not limited to the D-isomers of the common amino acids, 2,4-diaminobutyric acid, alpha.-amino isobutyric acid, 4-aminobutyric acid, Abu, 2-amino butyric acid, .gamma.-Abu, .epsilon.-Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid, 3-amino propionic acid, ornithine, norleucine, norvaline, hydroxyproline, sarcosine, citrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, beta.-alanine, fluoro-amino acids, designer amino acids such as .beta.-methyl amino acids, C.alpha.-methyl amino acids,
15 N.alpha.-methyl amino acids, and amino acid analogs in general. Furthermore, the amino acid can be D (dextrorotary) or L (levorotary).

Manipulations of the polypeptide sequences of the invention may be made at the protein level. Included within the scope of the invention are polypeptides of the invention, fragments thereof, derivatives or analogs which are differentially modified
25 during or after translation, e.g., by glycosylation, acetylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to an antibody molecule or other cellular ligand, etc. Any of numerous chemical modifications may be carried out by known techniques, including, but not limited to, specific chemical cleavage by cyanogen bromide, trypsin, chymotrypsin, papain, V8 protease, NaBH₄, acetylation, formulation, oxidation, reduction,
30 metabolic synthesis in the presence of tunicamycin, etc.

Moreover, the invention encompasses additional post-translational modifications which include, for example, the addition of N-linked or O-linked carbohydrate moieties or chains, the addition of detectable labels which may be
35 fluorescent, radioisotopic, enzymatic, in nature, the addition of epitope tagged peptide fragments (e.g., FLAG, HA, GST, thioredoxin, maltose binding protein, etc.), attachment of affinity tags such as biotin and/or streptavidin, the covalent attachment of chemical moieties to the amino acid backbone, N- or C-terminal processing of the

5 polypeptides ends (e.g., proteolytic processing), deletion of the N-terminal methioine residue, etc.

Furthermore, the invention encompasses chemical derivitization of the polypeptides of the present invention, preferably where the chemical is a hydrophilic polymer residue. Exemplary hydrophilic polymers, including derivitives, may be
10 those that include polymers in which the repeating units contain one or more hydroxy groups (polyhydroxy polymers), including, for example, poly(vinyl alcohol); polymers in which the repeating units contain one or more amino groups (polyamine polymers), including, for example, peptides, polypeptides, proteins and lipoproteins, such as albumin and natural lipoproteins; polymers in which the repeating units
15 contain one or more carboxy groups (polycarboxy polymers), including, for example, carboxymethylcellulose, alginic acid and salts thereof, such as sodium and calcium alginate, glycosaminoglycans and salts thereof, including salts of hyaluronic acid, phosphorylated and sulfonated derivatives of carbohydrates, genetic material, such as interleukin-2 and interferon, and phosphorothioate oligomers; and polymers in which
20 the repeating units contain one or more saccharide moieties (polysaccharide polymers), including, for example, carbohydrates.

The molecular weight of the hydrophilic polymers may vary, and is generally about 50 to about 5,000,000, with polymers having a molecular weight of about 100 to about 50,000 being preferred. More preferred polymers have a molecular weight of
25 about 150 to about 10,000, with molecular weights of 200 to about 8,000 being even more preferred.

Additional preferred polymers which may be used to derivitize polypeptides of the invention, include, for example, poly(ethylene glycol) (PEG), poly(vinylpyrrolidone), polyoxomers, polysorbate and poly(vinyl alcohol), with PEG
30 polymers being particularly preferred. Preferred among the PEG polymers are PEG polymers having a molecular weight of from about 100 to about 10,000. More preferably, the PEG polymers have a molecular weight of from about 200 to about 8,000, with PEG 2,000, PEG 5,000 and PEG 8,000, which have molecular weights of 2,000, 5,000 and 8,000, respectively, being even more preferred. Other suitable
35 hydrophilic polymers, in addition to those exemplified above, will be readily apparent to one skilled in the art based on the present disclosure. Generally, the polymers used may include polymers that can be attached to the polypeptides of the invention via alkylation or acylation reactions.

5 As with the various polymers exemplified above, it is contemplated that the polymeric residues may contain functional groups in addition, for example, to those typically involved in linking the polymeric residues to the polypeptides of the present invention. Such functionalities include, for example, carboxyl, amine, hydroxy and thiol groups. These functional groups on the polymeric residues can be further
10 reacted, if desired, with materials that are generally reactive with such functional groups and which can assist in targeting specific tissues in the body including, for example, diseased tissue. Exemplary materials which can be reacted with the additional functional groups include, for example, proteins, including antibodies, carbohydrates, peptides, glycopeptides, glycolipids, lectins, and nucleosides.

15 In addition to residues of hydrophilic polymers, the chemical used to derivitize the polypeptides of the present invention can be a saccharide residue. Exemplary saccharides which can be derived include, for example, monosaccharides or sugar alcohols, such as erythrose, threose, ribose, arabinose, xylose, lyxose, fructose, sorbitol, mannitol and sedoheptulose, with preferred monosaccharides being fructose,
20 mannose, xylose, arabinose, mannitol and sorbitol; and disaccharides, such as lactose, sucrose, maltose and cellobiose. Other saccharides include, for example, inositol and ganglioside head groups. Other suitable saccharides, in addition to those exemplified above, will be readily apparent to one skilled in the art based on the present disclosure. Generally, saccharides which may be used for derivitization include
25 saccharides that can be attached to the polypeptides of the invention via alkylation or acylation reactions.

 Moreover, the invention also encompasses derivitization of the polypeptides of the present invention, for example, with lipids (including cationic, anionic, polymerized, charged, synthetic, saturated, unsaturated, and any combination of the
30 above, etc.). stabilizing agents.

 The invention encompasses derivitization of the polypeptides of the present invention, for example, with compounds that may serve a stabilizing function (e.g., to increase the polypeptides half-life in solution, to make the polypeptides more water soluble, to increase the polypeptides hydrophilic or hydrophobic character, etc.).
35 Polymers useful as stabilizing materials may be of natural, semi-synthetic (modified natural) or synthetic origin. Exemplary natural polymers include naturally occurring polysaccharides, such as, for example, arabinans, fructans, fucans, galactans, galacturonans, glucans, mannans, xylans (such as, for example, inulin), levan,

5 fucoidan, carrageenan, galatocarolose, pectic acid, pectins, including amylose, pullulan, glycogen, amylopectin, cellulose, dextran, dextrin, dextrose, glucose, polyglucose, polydextrose, pustulan, chitin, agarose, keratin, chondroitin, dermatan, hyaluronic acid, alginic acid, xanthin gum, starch and various other natural homopolymer or heteropolymers, such as those containing one or more of the
10 following aldoses, ketoses, acids or amines: erythrose, threose, ribose, arabinose, xylose, lyxose, allose, altrose, glucose, dextrose, mannose, gulose, idose, galactose, talose, erythrulose, ribulose, xylulose, psicose, fructose, sorbose, tagatose, mannitol, sorbitol, lactose, sucrose, trehalose, maltose, cellobiose, glycine, serine, threonine, cysteine, tyrosine, asparagine, glutamine, aspartic acid, glutamic acid, lysine, arginine, histidine, glucuronic acid, gluconic acid, glucaric acid, galacturonic acid,
15 mannuronic acid, glucosamine, galactosamine, and neuraminic acid, and naturally occurring derivatives thereof Accordingly, suitable polymers include, for example, proteins, such as albumin, polyalginates, and polylactide-coglycolide polymers. Exemplary semi-synthetic polymers include carboxymethylcellulose,
20 hydroxymethylcellulose, hydroxypropylmethylcellulose, methylcellulose, and methoxycellulose. Exemplary synthetic polymers include polyphosphazenes, hydroxyapatites, fluoroapatite polymers, polyethylenes (such as, for example, polyethylene glycol (including for example, the class of compounds referred to as Pluronic.RTM., commercially available from BASF, Parsippany, N.J.),
25 polyoxyethylene, and polyethylene terephthalate), polypropylenes (such as, for example, polypropylene glycol), polyurethanes (such as, for example, polyvinyl alcohol (PVA), polyvinyl chloride and polyvinylpyrrolidone), polyamides including nylon, polystyrene, polylactic acids, fluorinated hydrocarbon polymers, fluorinated carbon polymers (such as, for example, polytetrafluoroethylene), acrylate,
30 methacrylate, and polymethylmethacrylate, and derivatives thereof. Methods for the preparation of derivitized polypeptides of the invention which employ polymers as stabilizing compounds will be readily apparent to one skilled in the art, in view of the present disclosure, when coupled with information known in the art, such as that described and referred to in Unger, U.S. Pat. No. 5,205,290, the disclosure of which is
35 hereby incorporated by reference herein in its entirety.

Moreover, the invention encompasses additional modifications of the polypeptides of the present invention. Such additional modifications are known in the

5 art, and are specifically provided, in addition to methods of derivitization, etc., in US Patent No. 6,028,066, which is hereby incorporated in its entirety herein.

Uses of the Polynucleotides

Each of the polynucleotides identified herein can be used in numerous ways as
10 reagents. The following description should be considered exemplary and utilizes known techniques.

The polynucleotides of the present invention are useful for chromosome identification. There exists an ongoing need to identify new chromosome markers, since few chromosome marking reagents, based on actual sequence data (repeat
15 polymorphisms), are presently available. Each polynucleotide of the present invention can be used as a chromosome marker.

Briefly, sequences can be mapped to chromosomes by preparing PCR primers (preferably 15-25 bp) from the sequences shown in SEQ ID NO:X. Primers can be selected using computer analysis so that primers do not span more than one predicted
20 exon in the genomic DNA. These primers are then used for PCR screening of somatic cell hybrids containing individual human chromosomes. Only those hybrids containing the human gene corresponding to the SEQ ID NO:X will yield an amplified fragment.

Similarly, somatic hybrids provide a rapid method of PCR mapping the
25 polynucleotides to particular chromosomes. Three or more clones can be assigned per day using a single thermal cycler. Moreover, sublocalization of the polynucleotides can be achieved with panels of specific chromosome fragments. Other gene mapping strategies that can be used include in situ hybridization, prescreening with labeled flow-sorted chromosomes, and preselection by hybridization to construct
30 chromosome specific-cDNA libraries.

Precise chromosomal location of the polynucleotides can also be achieved using fluorescence in situ hybridization (FISH) of a metaphase chromosomal spread. This technique uses polynucleotides as short as 500 or 600 bases; however, polynucleotides 2,000-4,000 bp are preferred. For a review of this technique, see
35 Verma et al., "Human Chromosomes: a Manual of Basic Techniques," Pergamon Press, New York (1988).

For chromosome mapping, the polynucleotides can be used individually (to mark a single chromosome or a single site on that chromosome) or in panels (for

5 marking multiple sites and/or multiple chromosomes). Preferred polynucleotides correspond to the noncoding regions of the cDNAs because the coding sequences are more likely conserved within gene families, thus increasing the chance of cross hybridization during chromosomal mapping.

10 Once a polynucleotide has been mapped to a precise chromosomal location, the physical position of the polynucleotide can be used in linkage analysis. Linkage analysis establishes coinheritance between a chromosomal location and presentation of a particular disease. Disease mapping data are known in the art. Assuming 1 megabase mapping resolution and one gene per 20 kb, a cDNA precisely localized to a chromosomal region associated with the disease could be one of 50-500 potential
15 causative genes.

Thus, once coinheritance is established, differences in the polynucleotide and the corresponding gene between affected and unaffected organisms can be examined. First, visible structural alterations in the chromosomes, such as deletions or translocations, are examined in chromosome spreads or by PCR. If no structural
20 alterations exist, the presence of point mutations are ascertained. Mutations observed in some or all affected organisms, but not in normal organisms, indicates that the mutation may cause the disease. However, complete sequencing of the polypeptide and the corresponding gene from several normal organisms is required to distinguish the mutation from a polymorphism. If a new polymorphism is identified, this
25 polymorphic polypeptide can be used for further linkage analysis.

Furthermore, increased or decreased expression of the gene in affected organisms as compared to unaffected organisms can be assessed using polynucleotides of the present invention. Any of these alterations (altered expression, chromosomal rearrangement, or mutation) can be used as a diagnostic or prognostic
30 marker.

Thus, the invention also provides a diagnostic method useful during diagnosis of a disorder, involving measuring the expression level of polynucleotides of the present invention in cells or body fluid from an organism and comparing the measured gene expression level with a standard level of polynucleotide expression
35 level, whereby an increase or decrease in the gene expression level compared to the standard is indicative of a disorder.

By "measuring the expression level of a polynucleotide of the present invention" is intended qualitatively or quantitatively measuring or estimating the level

5 of the polypeptide of the present invention or the level of the mRNA encoding the polypeptide in a first biological sample either directly (e.g., by determining or estimating absolute protein level or mRNA level) or relatively (e.g., by comparing to the polypeptide level or mRNA level in a second biological sample). Preferably, the polypeptide level or mRNA level in the first biological sample is measured or
10 estimated and compared to a standard polypeptide level or mRNA level, the standard being taken from a second biological sample obtained from an individual not having the disorder or being determined by averaging levels from a population of organisms not having a disorder. As will be appreciated in the art, once a standard polypeptide level or mRNA level is known, it can be used repeatedly as a standard for
15 comparison.

By "biological sample" is intended any biological sample obtained from an organism, body fluids, cell line, tissue culture, or other source which contains the polypeptide of the present invention or mRNA. As indicated, biological samples include body fluids (such as the following non-limiting examples, phloem, xylem,
20 secreted fluids, nectar, nodule fluid, appressorium fluid, canker fluid, gall fluid, fruit juice, trichome fluid, vacuole fluid, plastid fluid, cytosolic fluid, root exudates, interstitial fluid, etc.) which contain the polypeptide of the present invention, and other tissue sources found to express the polypeptide of the present invention (such as the following non-limiting examples root, stem, apical meristem, leaves, flowers,
25 pedals, etc.). Methods for obtaining tissue biopsies and body fluids from plants are well known in the art. For example, methods for isolating interstitial fluid of plant cells and tissues can be found, for example, in International Publication No. WO 00/09725. Where the biological sample is to include mRNA, a tissue biopsy is the preferred source.

30 The method(s) provided above may preferably be applied in a diagnostic method and/or kits in which polynucleotides and/or polypeptides are attached to a solid support. In one exemplary method, the support may be a "gene chip" or a "biological chip" as described in US Patents 5,837,832, 5,874,219, and 5,856,174. Further, such a gene chip with polynucleotides of the present invention attached may
35 be used to identify polymorphisms between the polynucleotide sequences, with polynucleotides isolated from a test subject. The knowledge of such polymorphisms (i.e. their location, as well as, their existence) would be beneficial in identifying disease loci for many disorders, including proliferative diseases and conditions. Such

5 a method is described in US Patents 5,858,659 and 5,856,104. The US Patents referenced supra are hereby incorporated by reference in their entirety herein.

The present invention encompasses polynucleotides of the present invention that are chemically synthesized, or reproduced as peptide nucleic acids (PNA), or according to other methods known in the art. The use of PNAs would serve as the preferred form if the polynucleotides are incorporated onto a solid support, or gene
10 chip. For the purposes of the present invention, a peptide nucleic acid (PNA) is a polyamide type of DNA analog and the monomeric units for adenine, guanine, thymine and cytosine are available commercially (Perceptive Biosystems). Certain components of DNA, such as phosphorus, phosphorus oxides, or deoxyribose
15 derivatives, are not present in PNAs. As disclosed by P. E. Nielsen, M. Egholm, R. H. Berg and O. Buchardt, *Science* 254, 1497 (1991); and M. Egholm, O. Buchardt, L.Christensen, C. Behrens, S. M. Freier, D. A. Driver, R. H. Berg, S. K. Kim, B. Norden, and P. E. Nielsen, *Nature* 365, 666 (1993), PNAs bind specifically and tightly to complementary DNA strands and are not degraded by nucleases. In fact,
20 PNA binds more strongly to DNA than DNA itself does. This is probably because there is no electrostatic repulsion between the two strands, and also the polyamide backbone is more flexible. Because of this, PNA/DNA duplexes bind under a wider range of stringency conditions than DNA/DNA duplexes, making it easier to perform multiplex hybridization. Smaller probes can be used than with DNA due to the
25 stronger binding characteristics of PNA:DNA hybrids. In addition, it is more likely that single base mismatches can be determined with PNA/DNA hybridization because a single mismatch in a PNA/DNA 15-mer lowers the melting point ($T_{sub.m}$) by 8°-20° C, vs. 4°-16° C for the DNA/DNA 15-mer duplex. Also, the absence of charge groups in PNA means that hybridization can be done at low ionic strengths and reduce
30 possible interference by salt during the analysis.

In addition to the foregoing, a polynucleotide can be used to control gene expression through triple helix formation or antisense DNA or RNA. Antisense techniques are discussed, for example, in Okano, *J. Neurochem.* 56: 560 (1991); "Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca
35 Raton, FL (1988). Triple helix formation is discussed in, for instance Lee et al., *Nucleic Acids Research* 6: 3073 (1979); Cooney et al., *Science* 241: 456 (1988); and Dervan et al., *Science* 251: 1360 (1991). Both methods rely on binding of the polynucleotide to a complementary DNA or RNA. For these techniques, preferred

5 polynucleotides are usually oligonucleotides 20 to 40 bases in length and complementary to either the region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxy-nucleotides as Antisense Inhibitors
10 of Gene Expression, CRC Press, Boca Raton, FL (1988).) Triple helix formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques are effective in model systems, and the information disclosed herein can be used to design antisense or triple helix polynucleotides in an effort to treat or
15 prevent disease.

The present invention encompasses the addition of a nuclear localization signal, operably linked to the 5' end, 3' end, or any location therein, to any of the oligonucleotides, antisense oligonucleotides, triple helix oligonucleotides, ribozymes, PNA oligonucleotides, and/or polynucleotides, of the present invention. See, for
20 example, G. Cutrona, et al., Nat. Biotech., 18:300-303, (2000); which is hereby incorporated herein by reference.

Polynucleotides of the present invention are also useful in gene therapy. One goal of gene therapy is to insert a normal gene into an organism having a defective gene, in an effort to correct the genetic defect. The polynucleotides disclosed in the
25 present invention offer a means of targeting such genetic defects in a highly accurate manner. Another goal is to insert a new gene that was not present in the host genome, thereby producing a new trait in the host cell. In one example, polynucleotide sequences of the present invention may be used to construct chimeric RNA/DNA oligonucleotides corresponding to said sequences, specifically designed to induce host
30 cell mismatch repair mechanisms in a plant upon systemic injection, for example (Bartlett, R.J., et al., Nat. Biotech, 18:615-622 (2000), which is hereby incorporated by reference herein in its entirety). Such RNA/DNA oligonucleotides could be designed to correct genetic defects in certain host strains, and/or to introduce desired traits in the plant host (e.g., introduction of a specific polymorphism within an
35 endogenous gene corresponding to a polynucleotide of the present invention that may confer resistance to certain herbicides, pesticides, fungicides, etc.). Alternatively, the polynucleotide sequence of the present invention may be used to construct duplex oligonucleotides corresponding to said sequence, specifically designed to correct

5 genetic defects in certain host strains, and/or to introduce desired traits in the plant
host (e.g., introduction of a specific polymorphism within an endogenous gene
corresponding to a polynucleotide of the present invention that may confer resistance
to certain herbicides, pesticides, fungicides, etc.). Such methods of using duplex
oligonucleotides are known in the art and are encompassed by the present invention
10 (see EP1007712, which is hereby incorporated by reference herein in its entirety).

The polynucleotides are also useful for identifying organisms from minute
biological samples. The United States military, for example, is considering the use of
restriction fragment length polymorphism (RFLP) for identification of its personnel.
In this technique, an individual's genomic DNA is digested with one or more
15 restriction enzymes, and probed on a Southern blot to yield unique bands for
identifying personnel. This method does not suffer from the current limitations of
"Dog Tags" which can be lost, switched, or stolen, making positive identification
difficult. The polynucleotides of the present invention can be used as additional DNA
markers for RFLP.

20 The polynucleotides of the present invention can also be used as an alternative
to RFLP, by determining the actual base-by-base DNA sequence of selected portions
of an organisms genome. These sequences can be used to prepare PCR primers for
amplifying and isolating such selected DNA, which can then be sequenced. Using
this technique, organisms can be identified because each organism will have a unique
25 set of DNA sequences. Once an unique ID database is established for an organism,
positive identification of that organism, living or dead, can be made from extremely
small tissue samples. Similarly, polynucleotides of the present invention can be used
as polymorphic markers, in addition to, the identification of transformed or non-
transformed plant cells and/or tissues.

30 There is also a need for reagents capable of identifying the source of a
particular tissue. Such need arises, for example, when presented with tissue of
unknown origin. Appropriate reagents can comprise, for example, DNA probes or
primers specific to particular tissue prepared from the sequences of the present
invention. Panels of such reagents can identify tissue by species and/or by organ type.
35 In a similar fashion, these reagents can be used to screen tissue cultures for
contamination. Moreover, as mentioned above, such reagents can be used to screen
and/or identify transformed and non-transformed plant cells and/or tissues.

5 Moreover, the polynucleotides of the present invention, the polynucleotides encoding the polypeptides of the present invention, the cDNA contained in the deposit, including variants and/or fragments therein, may be useful for modulating, inhibiting, increasing, decreasing, or introducing the following, non-limiting traits in a plant: drought tolerance, UV tolerance, flower development, terpene synthesis, abiotic
10 stresse tolerance, heat stress tolerance, cold stress tolerance, nutritional stress tolerance, xenobiotic stress tolerance, protein storage capability, oil storage capability, amino acid content, amino acid composition, carbohydrate storage capability, oil content, oil composition, carbohydrate content, carbohydrate composition, fiber content, fiber composition, metabolite content, metaboliter composition, vitamin
15 content, and/or vitamin composition. The polynucleotides of the invention, may also be useful in modulating plant yield, plant development, plant differentiation, root growth, root morphology, plant color, plant aroma, plant flavor, palatability of plant tissue, plant organoleptic properties, may be useful in phytoremediation, and/or plant defense. Moreover, the polypeptides of the invention may also be useful in
20 modulating the plants ability to serve as a plant nutraceutical, pharmaceutical, or phytoceutical. Alternatively, polypeptides of the invention may also be useful in modulating the plants ability to produce plant nutraceuticals, pharmaceuticals, or phytoceuticals of either endogenous or exogenous origin (e.g., from another plant species, a human, a mammal, an animal, or other organism). In these contexts, the
25 term "plant" may be applied to mean any plant cell, plant tissue, plant fluid, or plant feature, and includes plant infection structures, which may include, but are not limited to an appressorium, a gall, a canker, and/or nodules. In these contexts, the term "modulate" may be applied to mean the qualitative or quantitative increase, decrease, introduction of, inhibition of, complete loss of, or over-expression of a specific trait or
30 characteristic as described above and elsewhere herein.

 In the very least, the polynucleotides of the present invention can be used as molecular weight markers on Southern gels, as diagnostic probes for the presence of a specific mRNA in a particular cell type, as a probe to "subtract-out" known sequences in the process of discovering novel polynucleotides, for selecting and making
35 oligomers for attachment to a "gene chip" or other support, to raise anti-DNA antibodies using DNA immunization techniques, and as an antigen to elicit an immune response.

5 Uses of the Polypeptides

Each of the polypeptides identified herein can be used in numerous ways. The following description should be considered exemplary and utilizes known techniques.

A polypeptide of the present invention can be used to assay protein levels in a biological sample using antibody-based techniques. For example, protein expression
10 in tissues can be studied with classical immunohistological methods. (Jalkanen, M., et al., J. Cell. Biol. 101:976-985 (1985); Jalkanen, M., et al., J. Cell . Biol. 105:3087-3096 (1987)). Other antibody-based methods useful for detecting protein gene expression include immunoassays, such as the enzyme linked immunosorbent assay (ELISA) and the radioimmunoassay (RIA). Suitable antibody assay labels are known
15 in the art and include enzyme labels, such as, glucose oxidase, and radioisotopes, such as iodine (125I, 121I), carbon (14C), sulfur (35S), tritium (3H), indium (112In), and technecium (99mTc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

In addition to assaying protein levels in a biological sample, proteins can also
20 be detected in vivo by imaging. Antibody labels or markers for in vivo imaging of protein include those detectable by X-radiography, NMR or ESR. For X-radiography, suitable labels include radioisotopes such as barium or cesium, which emit detectable radiation but are not overtly harmful to the subject. Suitable markers for NMR and ESR include those with a detectable characteristic spin, such as deuterium, which may
25 be incorporated into the antibody by labeling of nutrients for the relevant hybridoma.

A protein-specific antibody or antibody fragment which has been labeled with an appropriate detectable imaging moiety, such as a radioisotope (for example, 131I, 112In, 99mTc), a radio-opaque substance, or a material detectable by nuclear magnetic resonance, is introduced (for example, parenterally, subcutaneously, or
30 intraperitoneally) into the organism. It will be understood in the art that the size of the subject and the imaging system used will determine the quantity of imaging moiety needed to produce diagnostic images. In the case of a radioisotope moiety, the quantity of radioactivity injected will normally range from about 5 to 20 millicuries of 99mTc. The labeled antibody or antibody fragment will then preferentially
35 accumulate at the location of cells which contain the specific protein. In vivo tumor imaging is described in S.W. Burchiel et al., "Immunopharmacokinetics of Radiolabeled Antibodies and Their Fragments." (Chapter 13 in Tumor Imaging: The

5 Radiochemical Detection of Cancer, S.W. Burchiel and B. A. Rhodes, eds., Masson Publishing Inc. (1982)). Such methods are equally applicable to plants.

Thus, the invention provides a diagnostic method of a disorder, which involves (a) assaying the expression of a polypeptide of the present invention in cells or body fluid of an organism; (b) comparing the level of gene expression with a
10 standard gene expression level, whereby an increase or decrease in the assayed polypeptide gene expression level compared to the standard expression level is indicative of a disorder.

Moreover, polypeptides of the present invention can be used for the treatment, detection, and/or prevention of a disease, or disease state. For example, an organism
15 can be administered a polypeptide of the present invention in an effort to replace absent or decreased levels of the polypeptide, to supplement absent or decreased levels of a different polypeptide, to inhibit the activity of a polypeptide, either directly or indirectly, to activate the activity of a polypeptide, either directly or indirectly, (e.g., by binding to a receptor), to reduce the activity of a membrane bound receptor
20 by competing with it for free ligand, or to bring about a desired response (e.g., induction of differentiation, growth, senescence, germination, etc.).

Similarly, antibodies directed to a polypeptide of the present invention can also be used to treat, prevent, and/or diagnose disease. For example, administration of an antibody directed to a polypeptide of the present invention can bind and reduce
25 overproduction of the polypeptide. Similarly, administration of an antibody can activate the polypeptide, such as by binding to a polypeptide bound to a membrane (receptor).

At the very least, the polypeptides of the present invention can be used as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration
30 columns using methods well known to those of skill in the art. Polypeptides can also be used to raise antibodies, which in turn are used to measure protein expression from a recombinant cell, as a way of assessing transformation of the host cell. Moreover, the polypeptides of the present invention can be used to test the following biological activities.

35

Transgenic Methods

Another aspect of the present invention is to gene therapy methods for treating or preventing disorders, diseases and conditions. The gene therapy methods relate to the

5 introduction of nucleic acid (DNA, RNA, and antisense DNA or RNA) sequences into
an organism, preferably a plant, to achieve expression of a polypeptide of the present
invention. This method requires a polynucleotide which codes for a polypeptide of the
invention operatively linked to a promoter and any other genetic elements necessary
for the expression of the polypeptide by the target tissue. Such transgenic and delivery
10 techniques are known in the art, see, for example, WO90/11092, which is herein
incorporated by reference.

Thus, for example, cells from a plant may be engineered with a polynucleotide
(DNA or RNA) comprising a promoter operably linked to a polynucleotide of the
invention *ex vivo*, with the engineered cells then being introduced back into the plant
15 to "treat" the deficiency. Such methods are well-known in the art and are equally
applicable to plants. For example, see Belldegrun et al., *J. Natl. Cancer Inst.*, 85:207-
216 (1993); Ferrantini et al., *Cancer Research*, 53:107-1112 (1993); Ferrantini et al.,
J. Immunology 153: 4604-4615 (1994); Kaido, T., et al., *Int. J. Cancer* 60: 221-229
(1995); Ogura et al., *Cancer Research* 50: 5102-5106 (1990); Santodonato, et al.,
20 *Human Gene Therapy* 7:1-10 (1996); Santodonato, et al., *Gene Therapy* 4:1246-1255
(1997); and Zhang, et al., *Cancer Gene Therapy* 3: 31-38 (1996)), which are herein
incorporated by reference.

As discussed in more detail below, the polynucleotide constructs can be
delivered by any method that delivers materials to the cells of an organism, such as,
25 biolistic injection into the plant tissues (apical meristem, root, flower, stem, and the
like). The polynucleotide constructs may be delivered in an acceptable liquid or
aqueous carrier.

In one embodiment, the polynucleotide of the invention is delivered as a naked
polynucleotide. The term "naked" polynucleotide, DNA or RNA refers to sequences
30 that are free from any delivery vehicle that acts to assist, promote or facilitate entry
into the cell, including viral sequences, viral particles, liposome formulations,
lipofectin or precipitating agents and the like. However, the polynucleotides of the
invention can also be delivered in liposome formulations and lipofectin formulations
and the like can be prepared by methods well known to those skilled in the art. Such
35 methods are described, for example, in U.S. Patent Nos. 5,593,972, 5,589,466, and
5,580,859, which are herein incorporated by reference.

The polynucleotide vector constructs of the invention used in the gene therapy
method are preferably constructs that will not integrate into the host genome nor will

5 they contain sequences that allow for replication. Alternatively, the polynucleotide vector constructs of the invention may integrate into the host genome and may replicate. Appropriate vectors include pWLNEO, pSV2CAT, pOG44, pXT1 and pSG available from Stratagene; pSVK3, pBPV, pMSG and pSVL available from Pharmacia; and pEF1/V5, pcDNA3.1, and pRc/CMV2 available from Invitrogen.
10 Other suitable vectors will be readily apparent to the skilled artisan.

Any strong promoter known to those skilled in the art can be used for driving the expression of polynucleotide sequence of the invention. Suitable promoters include the 35S, 34S, and actin promoters, in addition to any other promoter known in the art and/or described elsewhere herein. The promoter also may be the native
15 promoter for the polynucleotides of the invention.

Unlike other gene therapy techniques, one major advantage of introducing naked nucleic acid sequences into target cells is the transitory nature of the polynucleotide synthesis in the cells. Studies have shown that non-replicating DNA sequences can be introduced into cells to provide production of the desired
20 polypeptide for periods of up to six months.

The preferred route of administration is by the parenteral route of injection into the interstitial space of tissues. However, other parenteral routes may also be used. In addition, naked DNA constructs can be delivered to a plants circulatory system by direct injection.

25 The naked polynucleotides are delivered by any method known in the art, including, but not limited to, direct needle injection at the delivery site, topical administration, and so-called "gene guns". These delivery methods are known in the art.

The constructs may also be delivered with delivery vehicles such as viral
30 sequences, viral particles, liposome formulations, lipofectin, precipitating agents, etc. Such methods of delivery are known in the art.

In certain embodiments, the polynucleotide constructs of the invention are complexed in a liposome preparation. Liposomal preparations for use in the instant invention include cationic (positively charged), anionic (negatively charged) and
35 neutral preparations. However, cationic liposomes are particularly preferred because a tight charge complex can be formed between the cationic liposome and the polyanionic nucleic acid. Cationic liposomes have been shown to mediate intracellular delivery of plasmid DNA (Felgner et al., Proc. Natl. Acad. Sci. USA ,

5 84:7413-7416 (1987), which is herein incorporated by reference); mRNA (Malone et al., Proc. Natl. Acad. Sci. USA , 86:6077-6081 (1989), which is herein incorporated by reference); and purified transcription factors (Debs et al., J. Biol. Chem., 265:10189-10192 (1990), which is herein incorporated by reference), in functional form.

10 Cationic liposomes are readily available. For example, N[1-2,3-dioleoyloxy)propyl]-N,N,N-triethylammonium (DOTMA) liposomes are particularly useful and are available under the trademark Lipofectin, from GIBCO BRL, Grand Island, N.Y. (See, also, Felgner et al., Proc. Natl Acad. Sci. USA , 84:7413-7416 (1987), which is herein incorporated by reference). Other commercially available
15 liposomes include transfectace (DDAB/DOPE) and DOTAP/DOPE (Boehringer).

Other cationic liposomes can be prepared from readily available materials using techniques well known in the art. See, e.g. PCT Publication NO: WO 90/11092 (which is herein incorporated by reference) for a description of the synthesis of DOTAP (1,2-bis(oleoyloxy)-3-(trimethylammonio)propane) liposomes. Preparation
20 of DOTMA liposomes is explained in the literature, see, e.g., Felgner et al., Proc. Natl. Acad. Sci. USA, 84:7413-7417, which is herein incorporated by reference. Similar methods can be used to prepare liposomes from other cationic lipid materials.

Similarly, anionic and neutral liposomes are readily available, such as from Avanti Polar Lipids (Birmingham, Ala.), or can be easily prepared using readily
25 available materials. Such materials include phosphatidyl, choline, cholesterol, phosphatidyl ethanolamine, dioleoylphosphatidyl choline (DOPC), dioleoylphosphatidyl glycerol (DOPG), dioleoylphosphatidyl ethanolamine (DOPE), among others. These materials can also be mixed with the DOTMA and DOTAP starting materials in appropriate ratios.

30 Methods for making liposomes using these materials are well known in the art. For example, commercially dioleoylphosphatidyl choline (DOPC), dioleoylphosphatidyl glycerol (DOPG), and dioleoylphosphatidyl ethanolamine (DOPE) can be used in various combinations to make conventional liposomes, with or without the addition of cholesterol. Thus, for example, DOPG/DOPC vesicles can be
35 prepared by drying 50 mg each of DOPG and DOPC under a stream of nitrogen gas into a sonication vial. The sample is placed under a vacuum pump overnight and is hydrated the following day with deionized water. The sample is then sonicated for 2 hours in a capped vial, using a Heat Systems model 350 sonicator equipped with an

5 inverted cup (bath type) probe at the maximum setting while the bath is circulated at 15EC. Alternatively, negatively charged vesicles can be prepared without sonication to produce multilamellar vesicles or by extrusion through nucleopore membranes to produce unilamellar vesicles of discrete size. Other methods are known and available to those of skill in the art.

10 The liposomes can comprise multilamellar vesicles (MLVs), small unilamellar vesicles (SUVs), or large unilamellar vesicles (LUVs), with SUVs being preferred. The various liposome-nucleic acid complexes are prepared using methods well known in the art. See, e.g., Straubinger et al., *Methods of Immunology*, 101:512-527 (1983), which is herein incorporated by reference. For example, MLVs containing nucleic acid can be prepared by depositing a thin film of phospholipid on the walls of a glass
15 tube and subsequently hydrating with a solution of the material to be encapsulated. SUVs are prepared by extended sonication of MLVs to produce a homogeneous population of unilamellar liposomes. The material to be entrapped is added to a suspension of preformed MLVs and then sonicated. When using liposomes containing
20 cationic lipids, the dried lipid film is resuspended in an appropriate solution such as sterile water or an isotonic buffer solution such as 10 mM Tris/NaCl, sonicated, and then the preformed liposomes are mixed directly with the DNA. The liposome and DNA form a very stable complex due to binding of the positively charged liposomes to the cationic DNA. SUVs find use with small nucleic acid fragments. LUVs are
25 prepared by a number of methods, well known in the art. Commonly used methods include Ca²⁺-EDTA chelation (Papahadjopoulos et al., *Biochim. Biophys. Acta*, 394:483 (1975); Wilson et al., *Cell*, 17:77 (1979)); ether injection (Deamer et al., *Biochim. Biophys. Acta*, 443:629 (1976); Ostro et al., *Biochem. Biophys. Res. Commun.*, 76:836 (1977); Fraley et al., *Proc. Natl. Acad. Sci. USA*, 76:3348 (1979));
30 detergent dialysis (Enoch et al., *Proc. Natl. Acad. Sci. USA*, 76:145 (1979)); and reverse-phase evaporation (REV) (Fraley et al., *J. Biol. Chem.*, 255:10431 (1980); Szoka et al., *Proc. Natl. Acad. Sci. USA*, 75:145 (1978); Schaefer-Ridder et al., *Science*, 215:166 (1982)), which are herein incorporated by reference.

Generally, the ratio of DNA to liposomes will be from about 10:1 to about
35 1:10. Preferably, the ration will be from about 5:1 to about 1:5. More preferably, the ration will be about 3:1 to about 1:3. Still more preferably, the ratio will be about 1:1. U.S. Patent NO: 5,676,954 (which is herein incorporated by reference) reports on the injection of genetic material, complexed with cationic liposomes carriers, into mice.

5 U.S. Patent Nos. 4,897,355, 4,946,787, 5,049,386, 5,459,127, 5,589,466, 5,693,622,
5,580,859, 5,703,055, and international publication NO: WO 94/9469 (which are
herein incorporated by reference) provide cationic lipids for use in transfecting DNA
into cells and mammals. U.S. Patent Nos. 5,589,466, 5,693,622, 5,580,859,
5,703,055, and international publication NO: WO 94/9469 (which are herein
10 incorporated by reference) provide methods for delivering DNA-cationic lipid
complexes to mammals. Such methods are equally applicable to plants and is within
the skill of the artisan.

In certain embodiments, cells are engineered, *ex vivo* or *in vivo*, using a
retroviral particle containing RNA that comprises a sequence encoding polypeptides
15 of the invention. Retroviral integration has been detected to occur in plants based
upon the identification of a pararetrovirus sequences within the tobacco genome.
Since such integration was determined to occur at very limited integration sites, such
a pararetrovirus may represent a desirable genetic transformation vehicle for the
polynucleotides of the present invention (Jakowitsch, J., et al., PNAS 96(23):13241-6
20 (1999).

The retroviral plasmid vector is employed to transduce packaging cell lines to
form producer cell lines. Examples of packaging cells which may be transfected
include, but are not limited to, the PE501, PA317, R-2, R-AM, PA12, T19-14X, VT-
19-17-H2, RCRE, RCRIP, GP+E-86, GP+envAm12, and DAN cell lines as described
25 in Miller, Human Gene Therapy , 1:5-14 (1990), which is incorporated herein by
reference in its entirety. The vector may transduce the packaging cells through any
means known in the art. Such means include, but are not limited to, electroporation,
the use of liposomes, and CaPO₄ precipitation. In one alternative, the retroviral
plasmid vector may be encapsulated into a liposome, or coupled to a lipid, and then
30 administered to a host.

The producer cell line generates infectious retroviral vector particles that
include polynucleotides encoding polypeptides of the invention. Such retroviral
vector particles then may be employed, to transduce eukaryotic cells, either *in vitro* or
in vivo. The transduced eukaryotic cells will express polypeptides of the invention.

35 The present invention also encompasses the application of retrotransposons to
the genetic transformation of plants. The retrotransposons would preferably represent
retrotransposons with known plant host range and would comprise polynucleotides
encoding polypeptides of the present invention. Many retrotransposons are known in

5 the art, some of which are described by Bennetzen JL, Trends Microbiol., 4(9):347-53 (1996) which is hereby incorporated herein by reference.

Another method of gene therapy involves operably associating heterologous control regions and endogenous polynucleotide sequences (e.g. encoding the polypeptide sequence of interest) via homologous recombination (see, e.g., U.S. Patent NO: 5,641,670, issued June 24, 1997; International Publication NO: WO 96/29411, published September 26, 1996; International Publication NO: WO 94/12650, published August 4, 1994; Koller et al., Proc. Natl. Acad. Sci. USA, 86:8932-8935 (1989); and Zijlstra et al., Nature, 342:435-438 (1989). This method involves the activation of a gene which is present in the target cells, but which is not normally expressed in the cells, or is expressed at a lower level than desired.

Polynucleotide constructs are made, using standard techniques known in the art, which contain the promoter with targeting sequences flanking the promoter. Suitable promoters are described herein. The targeting sequence is sufficiently complementary to an endogenous sequence to permit homologous recombination of the promoter-targeting sequence with the endogenous sequence. The targeting sequence will be sufficiently near the 5' end of the desired endogenous polynucleotide sequence so the promoter will be operably linked to the endogenous sequence upon homologous recombination.

The promoter and the targeting sequences can be amplified using PCR. Preferably, the amplified promoter contains distinct restriction enzyme sites on the 5' and 3' ends. Preferably, the 3' end of the first targeting sequence contains the same restriction enzyme site as the 5' end of the amplified promoter and the 5' end of the second targeting sequence contains the same restriction site as the 3' end of the amplified promoter. The amplified promoter and targeting sequences are digested and ligated together.

The promoter-targeting sequence construct is delivered to the cells, either as naked polynucleotide, or in conjunction with transformation-facilitating agents, such as liposomes, viral sequences, viral particles, whole viruses, lipofection, precipitating agents, etc., described in more detail above. The P promoter-targeting sequence can be delivered by any method, included direct needle injection, intravenous injection, topical administration, infusion, particle accelerators, etc. The methods are described in more detail below.

5 The promoter-targeting sequence construct is taken up by cells. Homologous recombination between the construct and the endogenous sequence takes place, such that an endogenous sequence is placed under the control of the promoter. The promoter then drives the expression of the endogenous sequence.

 Preferably, the polynucleotide encoding a polypeptide of the invention
10 contains a secretory signal sequence that facilitates secretion of the protein. Typically, the signal sequence is positioned in the coding region of the polynucleotide to be expressed towards or at the 5' end of the coding region. The signal sequence may be homologous or heterologous to the polynucleotide of interest and may be homologous or heterologous to the cells to be transformed. Additionally, the signal
15 sequence may be chemically synthesized using methods known in the art.

 Any mode of administration of any of the above-described polynucleotides constructs can be used so long as the mode results in the expression of one or more molecules in an amount sufficient to provide a therapeutic effect. This includes direct needle injection, systemic injection, infusion, biolistic injectors, particle accelerators
20 (i.e., "gene guns"), gelfoam sponge depots, other commercially available depot materials, osmotic pumps (e.g., Alza minipumps), and decanting or topical application. For example, direct injection of naked calcium phosphate-precipitated plasmid into rat liver and rat spleen or a protein-coated plasmid into the portal vein has resulted in gene expression of the foreign gene in the rat livers. (Kaneda et al.,
25 Science, 243:375 (1989)). Moreover, direct injection of naked DNA has been reported in plants and is encompassed by the present invention (Davey MR, et al., Plant Mol Biol, 13(3):273-85 (1989), and Potrykus I, Ciba Found Symp, 154:198-212 (1990)).

 A preferred method of local administration is by direct injection. Preferably, a
30 recombinant molecule of the present invention complexed with a delivery vehicle is administered by direct injection into or locally within the area of the organisms circulatory system (e.g., phloem, xylem, etc). Administration of a composition locally within the area of the organisms circulatory system refers to injecting the composition centimeters and preferably, millimeters within the organisms circulatory system.

35 Another method of local administration is to contact a polynucleotide construct of the present invention in or around a surgical wound or grafting. For example, the polynucleotide construct can be coated on the surface of tissue inside the wound or the construct can be injected into areas of tissue inside the wound.

5 Therapeutic compositions useful in systemic administration, include recombinant molecules of the present invention complexed to a targeted delivery vehicle of the present invention. Suitable delivery vehicles for use with systemic administration comprise liposomes comprising ligands for targeting the vehicle to a particular site.

10 Preferred methods of systemic administration, include injection, aerosol, percutaneous (topical) delivery. Injections can be performed using methods standard in the art. Aerosol delivery can also be performed using methods standard in the art (see, for example, Stribling et al., Proc. Natl. Acad. Sci. USA , 189:11277-11281 (1992), which is incorporated herein by reference). Topical delivery can be performed
15 by mixing a polynucleotide construct of the present invention with a lipophilic reagent (e.g., DMSO) that is capable of passing into the skin.

 Determining an effective amount of substance to be delivered can depend upon a number of factors including, for example, the chemical structure and biological activity of the substance, the age and weight of the plant or animal,
20 the precise condition requiring treatment and its severity, and the route of administration. The frequency of treatments depends upon a number of factors, such as the amount of polynucleotide constructs administered per application, as well as the half-life of the polynucleotide and polypeptides (i.e., the effective period of application). The precise amount, number of applications and timing of applications will be determined per
25 desired application.

 Therapeutic compositions of the present invention can be administered to any organism, preferably to plants. Preferred plants, may include the following non-limiting examples, including barley, oats, rye, sorghum, pea, sunflower, tobacco, cotton, petunia, tomato, broccoli, lettuce, apple, plum, orange, and lemon, and more
30 preferably rice, maize, conola, wheat, sugarbeet, sugarcane, and soybean, in addition to other plants known in the art and referenced more particularly elsewhere herein (e.g., Table 3).

 Moreover, the present invention encompasses transgenic cells, including, but not limited to seeds, organisms, and plants into which genes encoding polypeptides of the
35 present invention have been introduced. Non-limiting examples of suitable recipient plants for introducing polynucleotides of the invention, polynucleotides encoding the polypeptides of the invention, the cDNA contained in a deposit, and/or fragments, and variants therein, are listed in Table 3 below:

TABLE 3**RECIPIENT PLANTS**

<u>{PRIVATE }COMMON NAME</u>	<u>FAMILY</u>	<u>LATIN NAME</u>
Maize	Gramineae	<i>Zea mays</i>
Maize, Dent	Gramineae	<i>Zea mays dentiformis</i>
Maize, Flint	Gramineae	<i>Zea mays vulgaris</i>
Maize, Pop	Gramineae	<i>Zea mays microsperma</i>
Maize, Soft	Gramineae	<i>Zea mays amyloacea</i>
Maize, Sweet	Gramineae	<i>Zea mays amylodisaccharata</i>
Maize, Sweet	Gramineae	<i>Zea mays saccharata</i>
Maize, Waxy	Gramineae	<i>Zea mays ceratina</i>
Wheat, Dinkel	Pooideae	<i>Triticum spelta</i>
Wheat, Durum	Pooideae	<i>Triticum durum</i>
Wheat, English	Pooideae	<i>Triticum turgidum</i>
Wheat, Large Spelt	Pooideae	<i>Triticum spelta</i>
Wheat, Polish	Pooideae	<i>Triticum polonium</i>
Wheat, Poulard	Pooideae	<i>Triticum turgidum</i>
Wheat, Singlegrained	Pooideae	<i>Triticum monococcum</i>
Wheat, Small Spelt	Pooideae	<i>Triticum monococcum</i>
Wheat, Soft	Pooideae	<i>Triticum aestivum</i>
Rice	Gramineae	<i>Oryza sativa</i>
Rice, American Wild	Gramineae	<i>Zizania aquatica</i>
Rice, Australian	Gramineae	<i>Oryza australiensis</i>
Rice, Indian	Gramineae	<i>Zizania aquatica</i>
Rice, Red	Gramineae	<i>Oryza glaberrima</i>
Rice, Tuscarora	Gramineae	<i>Zizania aquatica</i>
Rice, West African	Gramineae	<i>Oryza glaberrima</i>
Barley	Pooideae	<i>Hordeum vulgare</i>
Barley, Abyssinian Intermediate, also Irregular	Pooideae	<i>Hordeum irregulare</i>
Barley, Ancestral Two-row	Pooideae	<i>Hordeum spontaneum</i>
Barley, Beardless	Pooideae	<i>Hordeum trifurcatum</i>
Barley, Egyptian	Pooideae	<i>Hordeum trifurcatum</i>
Barley, four-rowed	Pooideae	<i>Hordeum vulgare polystichon</i>
Barley, six-rowed	Pooideae	<i>Hordeum vulgare hexastichon</i>
Barley, Two-rowed	Pooideae	<i>Hordeum distichon</i>
Cotton, Abroma	Dicotyledoneae	<i>Abroma augusta</i>
Cotton, American Upland	Malvaceae	<i>Gossypium hirsutum</i>
Cotton, Asiatic Tree, also Indian Tree	Malvaceae	<i>Gossypium arboreum</i>
Cotton, Brazilian, also, Kidney, and,	Malvaceae	<i>Gossypium barbadense brasiliense</i>

<u>{PRIVATE }COMMON NAME</u>	<u>FAMILY</u>	<u>LATIN NAME</u>
Pernambuco		
Cotton, Levant	Malvaceae	Gossypium herbaceum
Cotton, Long Silk, also Long Staple, Sea Island	Malvaceae	Gossypium barbadense
Cotton, Mexican, also Short Staple	Malvaceae	Gossypium hirsutum
Soybean, Soya	Leguminosae	Glycine max
Sugar beet	Chenopodiaceae	Beta vulgaris altissima
Sugar cane	Woody-plant	Arenga pinnata
Tomato	Solanaceae	Lycopersicon esculentum
Tomato, Cherry	Solanaceae	Lycopersicon esculentum cerasiforme
Tomato, Common	Solanaceae	Lycopersicon esculentum commune
Tomato, Currant	Solanaceae	Lycopersicon pimpinellifolium
Tomato, Husk	Solanaceae	Physalis ixocarpa
Tomato, Hyenas	Solanaceae	Solanum incanum
Tomato, Pear	Solanaceae	Lycopersicon esculentum pyriforme
Tomato, Tree	Solanaceae	Cyphomandra betacea
Potato	Solanaceae	Solanum tuberosum
Potato, Spanish, Sweet potato	Convolvulaceae	Ipomoea batatas
Rye, Common	Pooideae	Secale cereale
Rye, Mountain	Pooideae	Secale montanum
Pepper, Bell	Solanaceae	Capsicum annuum grossum
Pepper, Bird, also Cayenne, Guinea	Solanaceae	Capsicum annuum minimum
Pepper, Bonnet	Solanaceae	Capsicum sinense
Pepper, Bullnose, also Sweet	Solanaceae	Capsicum annuum grossum
Pepper, Cherry	Solanaceae	Capsicum annuum cerasiforme
Pepper, Cluster, also Red Cluster	Solanaceae	Capsicum annuum fasciculatum
Pepper, Cone	Solanaceae	Capsicum annuum conoides
Pepper, Goat, also Spur	Solanaceae	Capsicum frutescens
Pepper, Long	Solanaceae	Capsicum frutescens longum
Pepper, Oranamental Red, also Wrinkled	Solanaceae	Capsicum annuum abbreviatum
Pepper, Tabasco Red	Solanaceae	Capsicum annuum conoides
Lettuce, Garden	Compositae	Lactuca sativa
Lettuce, Asparagus, also Celery	Compositae	Lactuca sativa asparagina
Lettuce, Blue	Compositae	Lactuca perennis
Lettuce, Blue, also Chicory	Compositae	Lactuca pulchella

<u>{PRIVATE }COMMON NAME</u>	<u>FAMILY</u>	<u>LATIN NAME</u>
Lettuce, Cabbage, also Head	Compositae	Lactuca sativa capitata
Lettuce, Cos, also Longleaf, Romaine	Compositae	Lactuca sativa longifolia
Lettuce, Crinkle, also Curled, Cutting, Leaf	Compositae	Lactuca sativa crispa
Celery	Umbelliferae	Apium graveolens dulce
Celery, Blanching, also Garden	Umbelliferae	Apium graveolens dulce
Celery, Root, also Turniprooted	Umbelliferae	Apium graveolens rapaceum
Eggplant, Garden	Solanaceae	Solanum melongena
Sorghum	Sorghum	All crop species
Alfalfa	Leguminosae	Medicago sativum
Carrot	Umbelliferae	Daucus carota sativa
Bean, Climbing	Leguminosae	Phaseolus vulgaris vulgaris
Bean, Sprouts	Leguminosae	Phaseolus aureus
Bean, Brazilian Broad	Leguminosae	Canavalia ensiformis
Bean, Broad	Leguminosae	Vicia faba
Bean, Common, also French, White, Kidney	Leguminosae	Phaseolus vulgaris
Bean, Egyptian	Leguminosae	Dolichos lablab
Bean, Long, also Yardlong	Leguminosae	Vigna sesquipedalis
Bean, Winged	Leguminosae	Psophocarpus tetragonolobus
Oat, also Common, Side, Tree	Avena	Sativa
Oat, Black, also Bristle, Lopsided	Avena	Strigosa
Oat, Bristle	Avena	
Pea, also Garden, Green, Shelling	Leguminosae	Pisum, sativum sativum
Pea, Blackeyed	Leguminosae	Vigna sinensis
Pea, Edible Podded	Leguminosae	Pisum sativum axiphium
Pea, Grey	Leguminosae	Pisum sativum speciosum
Pea, Winged	Leguminosae	Tetragonolobus purpureus
Pea, Wrinkled	Leguminosae	Pisum sativum medullare
Sunflower	Compositae	Helianthus annuus
Squash, Autumn, Winter	Dicotyledoneae	Cucurbita maxima
Squash, Bush, also Summer	Dicotyledoneae	Cucurbita pepo melopepo
Squash, Turban	Dicotyledoneae	Cucurbita maxima turbaniformis
Cucumber	Dicotyledoneae	Cucumis sativus
Cucumber, African, also Bitter		Momordica charantia

<u>{PRIVATE }COMMON NAME</u>	<u>FAMILY</u>	<u>LATIN NAME</u>
Cucumber, Squirting, also Wild		Ecballium elaterium
Cucumber, Wild		Cucumis anguria
Poplar, California	Woody-Plant	Populus trichocarpa
Poplar, European Black		Populus nigra
Poplar, Gray		Populus canescens
Poplar, Lombardy		Populus italica
Poplar, Silverleaf, also White		Populus alba
Poplar, Western Balsam		Populus trichocarpa
Tobacco	Solanaceae	Nicotiana
Arabidopsis Thaliana	Cruciferae	Arabidopsis thaliana
Turfgrass	Lolium	
Turfgrass	Agrostis	
	Other families of turfgrass	
Clover	Leguminosae	

5

Biological Activities

The polynucleotides or polypeptides, or agonists or antagonists of the present invention can be used in assays to test for one or more biological activities. If these
 10 polynucleotides and polypeptides do exhibit activity in a particular assay, it is likely that these molecules may be involved in the diseases associated with the biological activity. Thus, the polynucleotides or polypeptides, or agonists or antagonists could be used to treat the associated disease.

15

Hyperproliferative Disorders

Polynucleotides or polypeptides, or agonists or antagonists of the invention can be used to detect, prevent, and/or confer resistance to hyperproliferative diseases, disorders, and/or conditions, which include, but are not limited to, cankers, galls,
 20 tumors, appressorium, etc. Polynucleotides or polypeptides, or agonists or antagonists of the present invention may inhibit the proliferation of the disorder through direct or indirect interactions. Alternatively, polynucleotides or polypeptides, or agonists or

5 antagonists of the present invention may proliferate other cells which can inhibit the hyperproliferative disorder.

One preferred embodiment utilizes polynucleotides of the present invention to inhibit aberrant cellular division, by gene therapy using the present invention, and/or protein fusions or fragments thereof.

10 Thus, the present invention provides a method for treating or preventing cell proliferative diseases, disorders, and/or conditions by inserting into an abnormally proliferating cell a polynucleotide of the present invention, wherein said polynucleotide represses said expression.

Another embodiment of the present invention provides a method of treating or
15 preventing cell-proliferative diseases, disorders, and/or conditions in organisms comprising administration of one or more active gene copies of the present invention to an abnormally proliferating cell or cells. In a preferred embodiment, a polynucleotide of the present invention is a DNA construct comprising a recombinant expression vector effective in expressing a DNA sequence encoding said
20 polynucleotides. In another preferred embodiment of the present invention, the DNA construct encoding the polynucleotides of the present invention is inserted into cells to be treated utilizing a retrovirus vector, or more preferably a plant retrotransposon-based vector. In a most preferred embodiment, the viral vector is defective and will not transform non-proliferating cells, only proliferating cells. Moreover, in a
25 preferred embodiment, the polynucleotides of the present invention inserted into proliferating cells either alone, or in combination with or fused to other polynucleotides, can then be modulated via an external stimulus (i.e. magnetic, specific small molecule, chemical, or drug administration, etc.), which acts upon the promoter upstream of said polynucleotides to induce expression of the encoded
30 protein product. As such, a beneficial therapeutic affect of the present invention may be expressly modulated (i.e. to increase, decrease, or inhibit expression of the present invention) based upon said external stimulus.

Polynucleotides of the present invention may be useful in repressing expression of proliferating genes or antigens. By "repressing expression of the
35 proliferating genes" is intended the suppression of the transcription of the gene, the degradation of the gene transcript (pre-message RNA), the inhibition of splicing, the destruction of the messenger RNA, the prevention of the post-translational

5 modifications of the protein, the destruction of the protein, or the inhibition of the normal function of the protein.

For local administration to abnormally proliferating cells, polynucleotides of the present invention may be administered by any method known to those of skill in the art including, but not limited to transformation, electroporation, microinjection of
10 cells, or in vehicles such as liposomes, lipofectin, or as naked polynucleotides, or any other method described throughout the specification. The polynucleotide of the present invention may be delivered by known gene delivery systems such as, but not limited to, retroviral vectors (Gilboa, J. Virology 44:845 (1982); Hocke, Nature 320:275 (1986); Wilson, et al., Proc. Natl. Acad. Sci. U.S.A. 85:3014), vaccinia virus
15 system (Chakrabarty et al., Mol. Cell Biol. 5:3403 (1985) or other efficient DNA delivery systems (Yates et al., Nature 313:812 (1985)) known to those skilled in the art. These references are exemplary only and are hereby incorporated by reference. In order to specifically deliver or transform cells which are abnormally proliferating and spare non-dividing cells, it is preferable to utilize a retrovirus, or a plant
20 retrotransposon-based vector (as described in the art and elsewhere herein) delivery system known to those of skill in the art. Since host DNA replication is required for retroviral DNA to integrate and the retrovirus will be unable to self replicate due to the lack of the retrovirus genes needed for its life cycle. Utilizing such a retroviral delivery system for polynucleotides of the present invention will target said gene and
25 constructs to abnormally proliferating cells and will spare the non-dividing normal cells.

The polynucleotides of the present invention may be delivered directly to cell proliferative disorder/disease sites in the plants body by use of imaging devices used to guide an injecting needle directly to the disease site. The polynucleotides of the
30 present invention may also be administered to disease sites at the time of grafting, etc.

By "cell proliferative disease" is meant any disease or disorder, affecting any one or any combination of organs, cavities, or body parts, which is characterized by single or multiple local abnormal proliferations of cells, groups of cells, or tissues.

Any amount of the polynucleotides of the present invention may be
35 administered as long as it has a biologically inhibiting effect on the proliferation of the treated cells. Moreover, it is possible to administer more than one of the polynucleotide of the present invention simultaneously to the same site. By "biologically inhibiting" is meant partial or total growth inhibition as well as

5 decreases in the rate of proliferation or growth of the cells. The biologically inhibitory dose may be determined by assessing the effects of the polynucleotides of the present invention on target or abnormally proliferating cell growth in tissue culture, tumor growth in plants and cell cultures, or any other method known to one of ordinary skill in the art.

10 The present invention is further directed to antibody-based therapies which involve administering anti-polypeptides and anti-polynucleotide antibodies to a plant, for detecting, preventing, and/or conferring resistance to one or more of the described diseases, disorders, and/or conditions. Methods for producing anti-polypeptides and anti-polynucleotide antibodies polyclonal and monoclonal antibodies are described in
15 detail elsewhere herein. Such antibodies may be provided in pharmaceutically acceptable compositions as known in the art or as described herein.

A summary of the ways in which the antibodies of the present invention may be used therapeutically includes binding polynucleotides or polypeptides of the present invention locally or systemically in the body or by direct cytotoxicity of the
20 antibody, e.g. as mediated by complement (CDC) or by effector cells (ADCC). Some of these approaches are described in more detail below. Armed with the teachings provided herein, one of ordinary skill in the art will know how to use the antibodies of the present invention for diagnostic, monitoring or therapeutic purposes without undue experimentation.

25 In particular, the antibodies, fragments and derivatives of the present invention are useful for detecting, preventing, and/or conferring resistance to a plant having or developing cell proliferative and/or differentiation diseases, disorders, and/or conditions as described herein. Such treatment comprises administering a single or multiple doses of the antibody, or a fragment, derivative, or a conjugate thereof.

30 The antibodies of this invention may be advantageously utilized in combination with other monoclonal or chimeric antibodies, or with cytokinins or plant growth factors, for example, which serve to increase the number or activity of effector cells which interact with the antibodies.

It is preferred to use high affinity and/or potent in vivo inhibiting and/or
35 neutralizing antibodies against polypeptides or polynucleotides of the present invention, fragments or regions thereof, for both immunoassays directed to and therapy of diseases, disorders, and/or conditions related to polynucleotides or polypeptides, including fragments thereof, of the present invention. Such antibodies,

5 fragments, or regions, will preferably have an affinity for polynucleotides or polypeptides, including fragments thereof. Preferred binding affinities include those with a dissociation constant or K_d less than $5 \times 10^{-6}M$, $10^{-6}M$, $5 \times 10^{-7}M$, $10^{-7}M$, $5 \times 10^{-8}M$, $10^{-8}M$, $5 \times 10^{-9}M$, $10^{-9}M$, $5 \times 10^{-10}M$, $10^{-10}M$, $5 \times 10^{-11}M$, $10^{-11}M$, $5 \times 10^{-12}M$, $10^{-12}M$, $5 \times 10^{-13}M$, $10^{-13}M$, $5 \times 10^{-14}M$, $10^{-14}M$, $5 \times 10^{-15}M$, and $10^{-15}M$.

Polypeptides, including protein fusions, of the present invention, or fragments thereof may be useful in inhibiting proliferative cells or tissues through the induction of apoptosis. Said polypeptides may act either directly, or indirectly to induce apoptosis of proliferative cells and tissues. Apoptosis has been described in plants and is thought to be regulated via mechanisms analogous to apoptosis induction in animals (see, for example,

15 LoSchiavo F, et al., *Eur J Cell Biol.*, 79(4):294-8 (2000); and Tian R, et al., *FEBS Lett.*, 474(1):11-15 (2000)).

Moreover, in another preferred embodiment of the present invention, said polypeptides may induce apoptosis through other mechanisms, such as in the activation of other proteins which will activate apoptosis (e.g., caspase 3, poly(ADP-ribose) polymerase (PARP), etc), or through stimulating the expression of said proteins, either alone or in combination with small molecule drugs or adjuvants, such as apoptonin, galectins, thioredoxins, antiinflammatory proteins (See for example, 25 *Mutat Res* 400(1-2):447-55 (1998), *Med Hypotheses*.50(5):423-33 (1998), *Chem Biol Interact.* Apr 24;111-112:23-34 (1998), *J Mol Med*.76(6):402-12 (1998), *Int J Tissue React*;20(1):3-15 (1998), which are all hereby incorporated by reference).

In another embodiment, the invention provides a method of delivering compositions containing the polypeptides of the invention (e.g., compositions containing polypeptides or polypeptide antibodies associated with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs) to targeted cells expressing the polypeptide of the present invention. Polypeptides or polypeptide antibodies of the invention may be associated with with heterologous polypeptides, heterologous nucleic acids, toxins, or prodrugs via hydrophobic, hydrophilic, ionic and/or covalent interactions.

35 Polypeptides, protein fusions to, or fragments thereof, of the present invention are useful in enhancing the immunogenicity and/or antigenicity of proliferating cells or tissues, either directly, or indirectly, to said antigens and immunogens.

5

Infectious Diseases

Infectious agents may inhibit the plants overall ability to maintain plant homeostasis and/or cellular homeostasis. For example, the infectious agent may inhibit the plants ability to control cellular division, differentiation, and development; absorption of water and minerals from the soil and the translocation of these substances throughout the plant; photosynthesis and translocation of the photosynthetic products to areas of use or storage; metabolism of synthesized compounds; reproduction; and storage of plant food-stuffs for overwintering or reproduction, for example.

Affected cells and tissues of diseased plants are usually weakened or destroyed by disease-causing agents. The ability of such infected cells and tissues to perform normal physiological functions is thus reduced or completely inhibited, causing cell or plant death. The type of tissue affected determines the physiological function affected. For example, infection of the root (e.g., root rot), interferes with absorption of water and nutrients from the soil; infection of the xylem vessels (e.g., vascular wilts, cankers, etc.) interferes with translocation of water and minerals to the crown of the plant; infection of the foliage (e.g., leaf spots, blights, mosaics, etc.) interferes with photosynthesis; infection of the cortex (e.g., cortical canker, viral and mycoplasmal infections of phloem, etc.) interferes with the downward translocation of photosynthetic products; flower infections (e.g., bacterial and fungal blights, viral, mycoplasmal, and fungal infections of flowers, etc.) interfere with reproduction; and infections of fruit (e.g., fruit rot, etc.) interfere with reproduction or storage of reserve food stuffs for the new plant. The above list of infectious traits and/or symptoms is exemplary and should not be construed as limiting the present invention. Additional infectious traits are known in the art, some of which are described elsewhere herein (see for example, Agrios, G.N., in "Plant Pathology", 3rd Ed., Academic Press, Inc., (1988); which is hereby incorporated in its entirety herein).

Viruses are one example of an infectious agent that can cause disease or traits that can be detected, prevented, and/or resistance conferred to by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention. Examples of viruses, include, but are not limited to the following DNA and RNA viruses and viral families: Arbovirus, Adenoviridae, Arenaviridae, Arterivirus, Birnaviridae, Bunyaviridae, Caliciviridae, Circoviridae, Coronaviridae, Dengue, EBV, HIV,

5 Flaviviridae, Hepadnaviridae (Hepatitis), Herpesviridae (such as, Cytomegalovirus, Herpes Simplex, Herpes Zoster), Mononegavirus (e.g., Paramyxoviridae, Morbillivirus, Rhabdoviridae), Orthomyxoviridae (e.g., Influenza A, Influenza B, and parainfluenza), Papiloma virus, Papovaviridae, Parvoviridae, Picornaviridae, Poxviridae (such as Smallpox or Vaccinia), Reoviridae (e.g., Rotavirus), Retroviridae
10 (HTLV-I, HTLV-II, Lentivirus), and Togaviridae (e.g., Rubivirus). . Additional examples of viruses, include, but are not limited to to the following Tobamovirus group (e.g., Tobacco Mosaic), Tobravirus group (e.g., Tobacco rattle virus), Hordeivirus group (e.g., Barlet strip mosaic), Potexvirus group (e.g., Potato virus X), Carlavirus group (e.g., Carnation latent virus), Potyvirus group (e.g., potato virus Y),
15 Closterovirus group (e.g., beet yellows virus), Maize chlorotic dwarf virus, Tobacco necrosis virus, Tymovirus group (e.g., Turnip yellow mosiac virus), Tombusvirus group (e.g., Tomato bushy stunt virus), Sobemovirus group (e.g., Southern bean mosaic virus), Luteovirus group (e.g., Barley yellow dwarf virus), Comovirus group (e.g., Cowpea mosaic virus), Nepovirus group (e.g., Tobacco ringspot virus), Pea
20 enation mosaic virus, Dianthovirus group (e.g., Carnation ringspot virus), Cucumovirus group (e.g., Cucumber mosaic virus), Bromovirus group (e.g., Brome mosaic virus), Ilavirus group (e.g., Tobacco streak virus), Alfalfa mosaic virus, Tomatto spotter wilt virus, Rhabdoviridae (e.g., Lettuce necrotic yellows virus), Rioviridae (e.g., Wound tumor virus), Geminivirus group (e.g., Maize streak virus),
25 and Caulimovirus (e.g., Cauliflower mosaic virus). Additional viruses capable of infecting a plant or animal are known in the art (see, for example, G.N., Agrios, supra, and Jones, T.C., in "Veterinary Pathology", 4th Edition, Lea and Febiger, Philadelphia, (1972).

Viruses falling within these families can cause a variety of diseases or
30 symptoms, generally including, but not limited to: mosaics, ring spots, stunting, dwarfing, leaf roll, yellowing, streaking, pox, enation, tumors, pitting of stem, aspermy, sterility, pitting of fruit, flattening and distortion of the stem; and specifically include, but not limited to tobacco mosaic, bean mosaic, apple mosaic, pear ring pattern mosaic, maize dwarf mosaic, tulip breaking, tobacco ringspot,
35 prunus necrtic ringspot, elm ringspot, chysanthemum ringspot, lilac ringspot, blueberry ringspot, beet yellows, wheat streak mosaic, tobacco etch, vein enation, vein clearing, vein banding, vein necrosis, potato leaf roll, grape fan leaf, tomato shoestring, stunting, banana bunchy top, citrus tristeza, cacao swollen shoot, stem

5 pitting, apple flat limb, pear rough bark, stem necrosis, graft brown line, cherry black
canker, elm zonate canker, citrus woody gall, clover wound tumor, apple russet ring,
apple scar skin, pear stony pit, spotted wilt, etc. Viruses may also lead to decreased
photosynthesis, decreased chlorophyll per leaf, decreased photosynthesis, decreased
chlorophyll per leaf, decreased chlorophyll efficiency, decrease in plant hormone
10 production, decreased growth rate, decreased soluble nitrogen, decreased
carbohydrate levels, either an increase or decrease in respiration, aberrant plant
metabolism, decrease water translocation, decreased nutrient retention, increased
transpiration, reduced yields, modulate transcription of the plant, modulate translation
of the plant, and aberrant cellular metabolism. Additional symptoms caused by viral
15 infections are known in the art (see, for example, G.N., Agrios, supra; Jones, T.C., in
"Veterinary Pathology", 4th Edition, Lea and Febiger, Philadelphia, (1972); and in
"Viral and Rickettsial Infections of Animals", eds, Betts, A.O., and York, C.J.,
Academic Press, NY, (1967)). Polynucleotides or polypeptides, or agonists or
antagonists of the invention, can be used to either directly or indirectly, detect,
20 prevent, and/or confer resistance to any of these traits or diseases. For example, a
polynucleotide or polypeptide of the present invention may directly inhibit a disorder
or infection when transgenically overexpressed in a plant. Alternatively, for example,
a polynucleotide or polypeptide of the present invention may indirectly inhibit a
disorder or infection by inhibiting the ability of the virus to transmit the infection
25 from one plant to another.

As inferred to above, plant viral infections may be transmitted through a
number of mechanisms, which include, but are not limited to the following:
transmission through vegetative propagation, mechanical transmission through sap,
seed transmission, pollen transmission, insect transmission, mite transmission,
30 nematode transmission, fungal transmission, and dodder transmission.
Polynucleotides or polypeptides, or agonists or antagonists of the invention, can be
used to either directly or indirectly, detect, prevent, inhibit, and/or confer resistance to
any of these mechanisms of viral transmission.

Similarly, bacterial agents that can cause disease or symptoms in plants or
35 animals can be detected, prevented, and/or resistance conferred to the plant by a
polynucleotide or polypeptide and/or agonist or antagonist of the present invention.
Examples of such bacterial agents include, but not limited to, the following: Gram-
Negative and Gram-positive bacteria and bacterial families and fungi: Bacillaceae

5 (e.g., Anthrax, Clostridium), Bacteroidaceae, Blastomycosis, Bordetella, Borrelia
 (e.g., Borrelia burgdorferi), Brucellosis, Candidiasis, Campylobacter,
 Coccidioidomycosis, Cryptococcosis, Dermatocycoses, E. coli (e.g., Enterotoxigenic
 E. coli and Enterohemorrhagic E. coli), Enterobacteriaceae (Klebsiella, Salmonella
 (e.g., Salmonella typhi, and Salmonella paratyphi), Serratia, Yersinia), Erysipelothrix,
 10 Helicobacter, Legionellosis, Leptospirosis, Listeria, Mycoplasmatales,
 Mycobacterium leprae, Vibrio cholerae, Neisseriaceae (e.g., Acinetobacter,
 Gonorrhea, Meningococcal), Meisseria meningitidis, Pasteurellacea Infections (e.g.,
 Actinobacillus, Heamophilus (e.g., Heamophilus influenza type B), Pasteurella),
 Pseudomonas, Rickettsiaceae, Chlamydiaceae, Syphilis, Shigella spp.,
 15 Staphylococcal, Meningiococcal, Pneumococcal and Streptococcal (e.g.,
 Streptococcus pneumoniae and Group B Streptococcus). Additional examples of
 bacterial agents include, for example, Agrobacterium, Clavibacter, Erwinia,
 Pseudomonas, Xanthomonas, Streptomyces, Xylella, Mycoplasm, Acholeplasma, and
 Spiroplasmas. Additional bacterial agents capable of infecting a plant or animal are
 20 known in the art (see, for example, G.N., Agrios, supra, and Jones, T.C., in
 "Veterinary Pathology", 4th Edition, Lea and Febiger, Philadelphia, (1972)

Bacterial agents falling within any of the aforementioned families can cause a
 variety of diseases or symptoms, generally including, but not limited to: leaf spots,
 leaf blights, soft rots (e.g., of fruit, root, and storage organs, etc.), wilts, overgrowths,
 25 scabs, cankers, nodules, galls, yellowing, phloem necrosis, X-disease, fasciation, and
 hairy root; and specifically, for example, crown gall, twig gall, cane gall, potato ring
 rot, tomato canker and wilt, banana wilt bud blast, cutting rot, black venation, bulb
 rot, citrus canker, walnut blight, pototo scab, sweet potato soil rot, tobacco wildfire,
 bean blight, cucumber angular leaf spot, cotton angular leaf spot, cereal blight, grass
 30 blight, tomato bacterial spot, pepper bacterial spot, stone fruit bacterial spot, bacterial
 vascular wilts, cucurbit bacterial wilt, pear fire blight, apple fire blight, pototo ring
 rot, southern bacterial wilt of solanaceous plants, Moko disease of banana, gummosis
 of stone fruit trees, Pierce's Disease of grape, almond leaf scorch, alfalfa dwarf,
 phony peach, plum leaf scald, ratoon stunting, clover club leaf, aster yellows, big bud,
 35 apple proliferation, peach yellows, apple rubbery wood, pear decline, elm phloem
 necrosis, coconut lethal yellowing, citrus tubborn, decreased photosynthesis,
 decreased chorophyll per leaf, decreased chlorophyll efficiency, decrease in plant
 hormone production, decreased growth rate, decreased soluble nitrogen, decreased

5 carbohydrate levels, either an increase or decrease in respiration, aberrant plant
metabolism, decrease water translocation, decreased nutrient retention, increased
transpiration, reduced yields, modulate transcription of the plant, modulate translation
of the plant, aberrant cellular metabolism, and corn stunt. Additional symptoms
and diseases caused by bacterial agents are known in the art (see, for example, G.N.,
10 Agrios, supra, and Jones, T.C., in "Veterinary Pathology", 4th Edition, Lea and
Febiger, Philadelphia, (1972). Polynucleotides or polypeptides, agonists or
antagonists of the invention, can be used, either directly or indirectly, to detect,
prevent, and/or confer resistance to any of these symptoms or diseases.

Similarly, fungal agents that can cause disease or symptoms in plants
15 or animals can be detected, prevented, and/or resistance conferred to the plant by a
polynucleotide or polypeptide and/or agonist or antagonist of the present invention.
Examples of such fungal agents include, but not limited to, the following:
Actinomycetales (e.g., *Corynebacterium*, *Mycobacterium*, *Nocardia*), *Cryptococcus*
neoformans, *Aspergillus*, *Myxomycota* (e.g., *Myxomycetes* (*Fuligo*, *Muciliago*,
20 *Physarum*, *Physarales*, etc), and *Plasmodiophoromycetes* (*Plasmodiophora* (e.g., *P.*
brassicae), *Polymyxa* (e.g., *P. gramminis*, etc.), *Spongospora* (e.g., *S. subteranea*,
etc))), *Eumycota* (e.g., *Mastigomycotina*, *Chytridiomycetes* (e.g., *Olpidium*
brassicae, *Physoderma maydia*, *Synchytrium endobioticum*, *Urophlyctis alfalfae*,
etc.), *Oomycetes*, *Saprolegniales* (e.g., *Aphanomyces*, etc.), *Peronosporales*,
25 *Pythiaceae*, *Pythium* (e.g., *Phytophthora infestans*, etc.), *Albuginaceae* (e.g., *Albugo*
candida, etc.), *Peronosporaceae* (e.g., *Plasmopara viticola*, *Peronospora nicotianae*,
Bremia lactucae, *Sclerospora graminicola*, and *Pseudoperonospora cubensis*, etc.),
Zygomycotina, *Zygomycetes*, *Mucorales*, *Rhizopus* (e.g., *Choanephora cucurbitarum*,
etc.), *Endogonales*, *Endogone*, *Ascomycotina*, *Hemiascomycetes*, *Endomycetales*
30 (e.g., *Saccharomyces cerevisiae*, etc.), *Taphrina*, *Pytenomycetes*, *Erysiphales* (e.g.,
Erysiphe, *Microsphaera*, *Podosphaera leucotricha*, *Spaerotheca pannosa*, *Uncinula*
necator etc.), *Sphaeriales* (e.g., *Botryosphaeria obtusa*, *Ceratocystis*, *Diaporthe*,
Endothia parasitica, *Eutypa armeniacae*, *Glomerella cingulata*, *Gnomonia*, *Hypoxyton*
mammatum, *Rosellinia*, *Valsa*, *Xylaria*, etc.), *Hypocreales* (e.g., *Claviceps purpurea*,
35 *Gibberella*, *Nectria*, etc.), *Loculoascomycetes*, *Myriangiales* (e.g., *Elsinoe*, etc.),
Dothideales (e.g., *Capnodium*, *Didymella*, *Guignardia bidwellii*, *Microcyclus elei*,
Plowrightia morbosum, etc.) *Pleosporales* (e.g., *Cochliobolus sativus*,
Gaeumannomyces graminis, *Pyrenophora*, *Venturia inaequalis*, etc.), *Discomycetes*,

- 5 Phacidiales (e.g., *Rhytisma acerinum*), Helotiales (e.g., *Diplocarpos rosae*, *Higginsia hiemalis*, *Lophodermium*, *Monilinia fructicola*, *Pseudopeziza trifolii*, *Sclerotinia sclerotiorum*, etc.), Deuteromycotina, Coelomycetes, Sphaeropsidales (e.g., *Ascochyta pisi*, *Coniothyrium*, *Cytospora*, *Diplodia maydis*, *Phoma lingam*, *Phomopsis*, *Phyllosticta*, *Septoria apii*, etc.), Melanconiales (e.g., *Celletotrichum*,
10 *Coryneium beijerincki*, *Cylindrosporium*, *Gloeosporium*, *Marssonina*, *Melanconium fuligenum*, *Sphaceloma*, etc.), Hyphomycetes, Hyphales (e.g., *Alternaria*, *Asperigillus*, *Bipolaris*, *dreschslerea*, *Excerophilum*, *Botrytis cinerea*, *Cercospora*, *Fulvia fulva*, *Fusarium*, *Geotrichum candidum*, *Graphium ulmi*, *Penicium*, *Phymatotrichum omnivorum*, *Pyricularia*, *Spilocaea*, *Theilaviopsis basicola*,
15 *Trichoderma*, *Verticillum*, etc.), Agonomycetes, Agonomycetales (e.g., *Rhizoctonia*, *Sclerotium*, etc.), Basidiomycotina, Hemibasidiomycetes, Ustilaginales (e.g., *Sphacelthea*, *Tilletia*, *Urocystis cepulae*, *Ustilago*, etc.), Uredinales (e.g., *Cronartium*, *Gymnosporangium juniperi-virginianae*, *Melampsora lini*, *Phragmidium*, *Puccinia*, *Uromyces appendiculatus*, etc.), Hymenomycetes, Exobasidiales (e.g.,
20 *Exobasidium*, etc.), Aphyllchorales (e.g., *Aethalia*, *Corticium*, *Heterobasidium*, *Lenzites*, *Peniophora*, *Polyporus*, *Poria*, *Schizophyllum*, *Stereum*, etc.), Tulasnellales (e.g., *Thanatephorus*, *Typhula*, etc.), Agaricales (e.g., *Armillaria mellea*, *Marasmius*, *Pholiota*, *Pleurotus*, etc.), and particularly Ascomycota, Basidiomycota, Zygomycota, Oomycota, Plasmodiophoromycota, *Puccinia recondita*, *Rhizo puschinensis*, *Plasmo*
25 *paraviticola*, *Plasmodiophora brassicae*, *Erwinia amylovora*, *Elsinoe fawcettii*, *Phaeosphaeria nodorum*, *Mycosphaerella arachidis*, *Mycosphaerella berkeleyi*, *Mycosphaerella fijiensis*, *Mycosphaerella graminicola*, *Pyrenophora teres*, *Pyremophora tritici-repentis*, *Venturia carpophila*, *Alternaria brassicae*, *Alternaria kikuchiana*, *Alternaria mali*, *Alternaria solani*, *Cercospora beticola*, *Cladosporium*
30 *cucumerinum*, *Septoria lycopersici*, *Blumaria graminis*, *Erysiphe cichoracearum*, *Podosphaera leucotricha*, *Sphaerotheca fuliginea*, *Uncinila necator*, *Emericella nidulans*, *Penicillium digitatum*, *Penicillium italicum*, *Gibberella fujikuroi*, *Nectria haematococca*, *Fusarium culmorum*, *Fusarium oxysporum*, *Fusarium roseum*, *Gliocladium virens*, *Botryotinia fuckeliana*, *Monilinia fructigena*, *Sclerotinia*
35 *homoeocarpa*, *Sclerotinia sclerotiorum*, *Mollisia yallundae*, *Glomerella lagenaria*, *Saccaromyces cerevisiae*, *Neurospora crassa*, *Gaemannomyces graminis*, *Magneporthe grisea*, *Monographella nivalis*, *Rhyncosporium secalis*, *Athelia rolfsii*, *Typhula incarnata*, *Thanatephorus cucumeris*, *Rhizoctonia solani*, *Hemileia vastatrix*,

5 Puccinia hordei, and Uromyces appendiculatus,. Additional fungal agents are known in the art (see, for example, G.N., Agrios, supra, G.C. Ainsworth, in "Fungal Diseases of Animals", Commonwealth Agricultural Bureaux, Farmham Royal Bucks, England, (1959), and Jones, T.C., in "Veterinary Pathology", 4th Edition, Lea and Febiger, Philadelphia, (1972)).

10 Fungal agents falling within any of the aforementioned divisions, subdivisions, classes, orders, genus's, or species can cause a variety of diseases or symptoms, generally including, but not limited to: necrosis, plant death, cell death, hypotrophy, hypoplasia, stunting, hyperplasia (e.g., clubroot, galls, warts, witches brooms, leaf curls, etc.), tumors, leaf spots, blight, cankers, dieback, root rot, damping
15 off, basal stem rot, soft rots, dry rots, anthracnose, scab, decline, wilt, rust, mildew, and smut; and specifically, fructifications, powdery scab of potato, clubroot of crucifers, black wort of potato, crown wart of alfalfa, brown spot of corn, seed rot, seedling damping off, root and stem rot, blight, tuber rot, white rust, upper side, downy mildews, oospores on soybean seed, rhizopus soft rots, rhizopus fruit rot,
20 choanephora squash rot, bread mold, bud rot, stem rot, collar rot, crown rot, trunk rot, black pod disease, late blight of potatoes, Anthracnose diseases, Colletotrichum diseases, onion smudge, ergot, Botrytis diseases, vascular wilts, Dutch Elm disease, Gibberella diseases, Sclerotinia Diseases, Rhizoctonia Diseases, Sclerotium Diseases, postharvest decay of fruits and vegetables, decreased photosynthesis, decreased
25 chorophyll per leaf, decreased chlorophyll efficiency, decrease in plant hormone production, decreased growth rate, decreased soluble nitrogen, decreased carbohydrate levels, either an increase or decrease in respiration, aberant plant metabolism, decrease water translocation, decreased nutrient retention, increased transpiration, reduced yields, modulate transcription of the plant, modulate translation
30 of the plant, and aberrant cellular metababolism; and particularly citrus scab, grape black rot, wheat septoria nodorum blotch, early leaf spot of peanut, banana black sigatoka, wheat septoria tritici blotch, barley net blotch, cereal tan spot, apple scab, peach feckle, cabbage blackspot, apple leaf blotch, tomato and potato early blight, cercospora leaf spot of beet, cucumber scab, tomato seporial leaf spot, wheat powdery
35 mildew, barlet powdery mildew, cucumber powdery mildew, apple powdery mildew, cucumber powdery mildew, grape powdery mildew, saporphyte, green mold of citrus, blue mold of citrus, rice bakanae disease, cereal headscab, cucumber fusarium wilt, tomato fusarium wilt, radish yellow fusarium, rice damping, pepper botrytis blight,

5 botrytis blights of bean, botrytis blights of cuke, botrytis blights of grape, botrytis
blights of marigold, brown rot of stone fruits, turf dollar spot, wheat eye spot,
cucumber anthracnose, pestalotial leaf spot, rice blast, turf snow mold, barley scald,
typhula blight, rhizoctonia damping, coffee rust, wheat lead rust, barley leaf rust, and
10 bean rust. Polynucleotides or polypeptides, agonists or antagonists of the invention,
can be used, either directly or indirectly, to detect, prevent, and/or confer resistance to
any of these symptoms or diseases. Additional fungal agents capable of infecting a
plant or animal are known in the art (see, for example, G.N., Agrios, supra, G.C.
Ainsworth, in "Fungal Diseases of Animals", Commonwealth Agricultural Bureaux,
Farmham Royal Bucks, England, (1959), and Jones, T.C., in "Veterinary Pathology",
15 4th Edition, Lea and Febiger, Philadelphia, (1972)).

Moreover, parasitic agents causing disease or symptoms that can be
detected, prevented, and/or conferred resistance to by a polynucleotide or polypeptide
and/or agonist or antagonist of the present invention include, but not limited to, the
following families or class: Amebiasis, Babesiosis, Coccidiosis, Cryptosporidiosis,
20 Dientamoebiasis, Dourine, Ectoparasitic, Giardiasis, Helminthiasis, Leishmaniasis,
Theileriasis, Toxoplasmosis, Trypanosomiasis, and Trichomonas and Sporozoans
(e.g., Plasmodium virax, Plasmodium falciparum, Plasmodium malariae, and
Plasmodium ovale). Additional examples of parasitic agents include, for example,
Cuscutaceae (e.g., Cuscuta, dodder, etc.), Viscaceae (e.g., Arceuthobium (dwarf
25 mistletoe of conifers), Phoradendron (American true mistletoes of broadleaved trees),
and Viscum (European true mistletoes)), Orobanchaceae (e.g., Orobanche
(broomrapes of tobacco)), and Scrophulariaceae (e.g., Striga (witchweeds of monocot
plants)). These parasites can cause a variety of diseases or symptoms, including, but
not limited to: decreased water storage, decreased mineral availability, decreased
30 carbohydrate storage, decreased plant defense, and increased susceptibility to fungal,
bacterial, or viral infections. Additional parasitic diseases are known in the art (see,
for example, G.N., Agrios, supra, and Jones, T.C., in "Veterinary Pathology", 4th
Edition, Lea and Febiger, Philadelphia, (1972)). Polynucleotides or polypeptides, or
agonists or antagonists of the invention, can be used to detect, prevent, and/or confer
35 resistance to any of these symptoms or diseases.

Methods of treatment or prevention using a polypeptide or polynucleotide
and/or agonist or antagonist of the present invention are well known in the art, though
may include administering an effective amount of a polypeptide to the plant, seed,

5 tissue, or cells. Alternatively, treatment or prevention could be conferred by transforming the plant, seed, tissue, or cells with a polynucleotide of the present invention, or cells could be removed from the plant, transformed, and then returning the engineered cells to the plant (ex vivo therapy). Moreover, the polypeptide or polynucleotide of the present invention can be used as an antigen to create antibodies
10 that inhibit the pathogenicity of a particular infectious disease (e.g., inhibiting the expression of a plant-specific gene or protein critical for the pathogenicity of an infectious organism). In addition, any method of detecting, preventing, conferring resistance to, or inhibiting an infectious agent using a polypeptide or polynucleotide and/or agonist or antagonist of the present invention may be applied to detecting,
15 preventing, conferring resistance to, or inhibiting an infectious agent in a human, animal, mammal, or other organism, with or without additional modification.

Pest Tolerance

Nematodes are one example of pests capable of causing disease or traits that
20 can be detected, prevented, and/or resistance conferred to by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention. Examples of nematodes include, but are not limited to, the following: Tylenchida, Tylenchina, Tylenchoidia, Tylenchidea, Tylenchidae, Anguina, Ditylenchus, Tylenchorhynchidae, Tylenchorhynchus, Pratylenchidae, Pratylenchus, Radopholus, Hoplolaimidae,
25 Hoplolaimus, Rotylenchus, Helicotylenchus, Belonolaimidae, Belonolaimus, Heteroderoidea, Heteroderidae, Globodera, Heterodera, Meloidogyne, Nacobbidae, Nacobbus, Rotylenchulus, Criconematoidea, Criconematidae, Criconemella, Hemicyclophora, Paratylenchidae, Paratylenchus, Tylenchuidae, Tylenchulus, Aphelenchina, Aphelenchoidea, Aphelenchoididae, Aphlenenchoides,
30 Bursaphelenchus, Rhadinaphelenchus, Dorylaimida, Longidoridae, Longidorus, Xiphinema, Trichodoridae, Paratrachodorus, And Trichodorus. Other members of the nematode phylum, orders, suborders, superfamilies, families, genres, and species are known in the art.

Nematodes falling within any of the aforementioned, orders, suborders,
35 superfamilies, families, genres, and/or species can cause a variety of diseases or symptoms in plants, generally including, but not limited to: root knots, root galls, reduced plant growth, plant nutrient deficiencies, yellowing, wilting, reduced yields, poor product quality, plant galls, necrotic lesions, rots, twisting or distortion of leaves

5 and stems, abnormal floral development; hypertrophy, hypotrophy, cysts, and chlorosis. Polynucleotides or polypeptides, agonists or antagonists of the invention, can be used, either directly or indirectly, to detect, prevent, and/or confer resistance to any of these symptoms or diseases. Additional diseases and/or disorders caused by nematodes are known in the art.

10 Insects are another example of pests capable of causing disease or traits that can be detected, prevented, and/or resistance conferred to by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention. Examples of insects include, but are not limited to, the following: insects selected from the orders Coleoptera, Diptera, Hymenoptera, Lepidoptera, Mallophaga, Homoptera, Hemiptera,
15 Orthoptera, Thysanoptera, Dermaptera, Isoptera, Anoplura, Siphonaptera, Trichoptera, etc., particularly Coleoptera.

Polynucleotides or polypeptides, agonists or antagonists of the invention, can be used, either directly or indirectly, to detect, prevent, and/or confer resistance to any of the following, non-limiting, symptoms or diseases caused by insect pests of major
20 crops: Maize: *Ostrinia nubilalis*, European corn borer; *Agrotis ipsilon*, black cutworm; *Helicoverpa zea*, corn earworm; *Spodoptera flugiperda*, fall armyworm; *Diatraea grandiosella*, southwestern corn borer; *Elasmopalpus lignosellus*, lesser cornstalk borer; *Diatraea saccharalis*, sugarcane borer; *Diabrotica virgifera*, western corn rootworm; *Diabrotica barberi*, northern corn rootworm; *Diabrotica undecimpunctata*
25 *howardi*, spotted cucumber beetle, *Melanotus* spp., wireworms; *Cyclocephala borealis*, northern masked chafer (white grub); *Cyclocephala immaculata*, southern masked chafer (white grub); *Popillia japonica*, Japanese beetle; *Chaetocnema pulicaria*, corn flea beetle; *Sphenophorus maidis*, maize billbug; *Rhopalosiphum maidis*, corn leaf aphid; *Anuraphis maidiradicis*, corn root aphid; *Blissus leucopterus*,
30 chinch bug; *Melanoplus femurrubrum*, redlegged grasshopper; *Melanoplus sanguinipes*, migratory grasshopper; *Delia platura*, seedcorn maggot; *Agromyza parvicornis*, corn blotch leafminer; *Anaphothrips obscurus*, grass thrips; *Solenopsis milesta*, thief ant; *Tetranychus urticae*, twospotted spider mite; *Busseola fusca*, African Maize Stem Borer (AMB); *Sesamia calamistis*, African Pink Borer (AP13);
35 *Eldana saccharina*, African Sugarcane Borer (ASB); *Chilo partellus*, Sorghum Stem Borer (SSB); *Ostrinia furnacalis*, Oriental Corn Borer (OCB); *Sesamia nonagrioides*, Corn borer in Europe/N. Africa; Sorghum: *Chilo partellus*, sorghum borer; *Spodoptera flugiperda*, fall armyworm; *Helicoverpa zea*, corn earworm; *Elasmopalpus lignosellus*,

5 lesser cornstalk borer; *Agrotis subterranea*, granulate cutworm; *Phyllophaga crinita*, white grub; *Eleodes*, *Conoderus*, and *Aeolus* spp., wireworms; *Oulema melanopus*, cereal leaf beetle; *Chaetocnema pulicaria*, corn flea beetle; *Sphenophorus maidis*, maize billbug; *Rhopalosiphum maidis*; corn leaf aphid; *Siphaflava*, yellow sugarcane aphid; *Blissus leucopterus*, chinch bug; *Contarinia sorghicola*, sorghum midge;

10 *Tetranychus cinnabarinus*, carmine spider mite; *Tetranychus urticae*, twospotted spider mite; *Schizaphis graminum*, Greenbug (aphid); Wheat: *Pseudaletia unipunctata*, army worm; *Spodoptera frugiperda*, fall armyworm; *Elasmopalpus lignosellus*, lesser cornstalk borer; *Agrotis orthogonia*, plae western cutworm; *Oulema melanopus*, cereal leaf beetle; *Hypera punctata*, clover leaf weevil; *Diabrotica undecimpunctata howardi*, spotted cucumber beetle; Russian wheat aphid; *Schizaphis graminum*, greenbug; *Sitobion avenae*, English grain aphid; *Melanoplus femurrubrum*, redlegged grasshopper; *Melanoplus differentialis*, differential grasshopper; *Melanoplus sanguinipes*, migratory grasshopper; *Mayetiola destructor*, Hessian fly; *Sitodiplosis niosellana*, wheat midge; *Meromyza americana*, wheat stem

20 maggot; *Hylemya coarctata*, wheat bulb fly; *Frankliniella jusca*, tobacco thrips; *Cephus cinctus*, wheat stem sawfly; *Eriophyes tulipae*, wheat curl mite; Sunflower. *Suleima helianthana*, sunflower bud moth; *Homeosoma ellectellum*, sunflower head moth; *Zygoramma exclamationis*, sunflower beetle; *Bothyrus gibbosus*, carrot beetle; *Neolasioptera murfeldtiana*, sunflower seed midge; *Cochylis hospes*, banded

25 sunflower moth; *Rachiplusia nu*, argentina looper; *Smicronyx julvus*, red sunflower seed weevil; *Cylindrocopturus adpersus*, spotted sunflower stem weevil; Cotton: *Heliothis virescens*, tobacco budworm; *Helicoverpa zea*, bollworm; *Spodoptera exigua*, beet armyworm; *Pectinophora gossypiella*, pink bollworm; *Anthonomus grandis*, boll weevil; *Aphis gossypii*, cotton aphid; *Pseudatomoscelis seriatus*, cotton

30 fleahopper; *Trialeurodes abutilonea*, bandedwinged whitefly; *Lygus lineolaris*, tarnished plant bug; *Melanoplus femurrubrum*, redlegged grasshopper; *Melanoplus differentialis*, differential grasshopper; *Thrips tabaci*, onion thrips; *Frankliniella jusca*, tobacco thrips; *Tetranychus cinnabarinus*, carmine spider mite; *Tetranychus urticae*, twospotted spider mite; Rice: *Diatraea saccharalis*, sugarcane borer;

35 *Spodoptera frugiperda*, fall armyworm; *Helicoverpa zea*, corn earworm; *Colaspis brunnea*, grape colaspis; *Lissorhoptrus oryzophilus*, rice water weevil; *Sitophilus oryzae*, rice weevil; *Nephotettix nigropictus*, rice leafhopper; *Blissus leucopterus*, chinch bug; *Acrosternum hilare*, green stink bug; *Pseudoplusia includens*, soybean

5 looper; *Anticarsia gemmatalis*, velvetbean caterpillar; *Plathypena scabra*, green
 cloverworm; *Ostrinia nubilalis*, European corn borer; *Agrotis ipsilon*, black cutworm;
Spodoptera exigua, beet armyworm; *Heliothis virescens*, cotton boll worm;
Helicoverpa zea, cotton bollworm; *Epilachna varivestis*, Mexican bean beetle; *Myzus*
 10 *persicae*, green peach aphid; *Empoasca jabae*, potato leafhopper; *Acrosternum hilare*,
 green stink bug; *Melanoplus femurrubrum*, redlegged grasshopper; *Melanoplus*
differentialis, differential grasshopper; *Delia platura*, seedcorn maggot; *Sericothrips*
variabilis, soybean thrips; *Thrips tabaci*, onion thrips; *Tetranychus turkestanii*,
 strawberry spider mite; *Tetranychus urticae*, twospotted spider mite; Barley. *Ostrinia*
nubilalis, European corn borer; *Agrotis ipsilon*, black cutworm; *Schizaphis graminum*,
 15 greenbug; *Blissus leucopterus*, chinch bug; *Acrosternum hilare*, green stink bug;
Euschistus servus, brown stink bug; *Delia platura*, seedcorn maggot; *Mayetiola*
destructor, Hessian fly; *Petrobia latens*, brown wheat mite; Oil Seed Rgpe;
Brevicoryne brassicae, cabbage aphid; Flea beetle, *Phyllotreta* spp.; Bertha
 Armyworm; *Mamestra configurata*; Diamondback Moth; *Plutella xylostella*; Alfalfa:
 20 alfalfa looper, *Autographa californica*; alfalfa snout beetle, *Otiorhynchus ligusticii*;
 alfalfa caterpillar, *Colias eurytheme*; alfalfa blotch leafrunner, *Agronyza frontella*;
 Egyptian alfalfa weevil, *Hypera brunneipennis*; meadow spittlebug, *Philaerius*
spumarius; spotted alfalfa aphid, *Theriothis meculata*; clover leaf weevil, *Hypera*
punctata; pea aphid, *Acyrtosiphon pisum*; blue alfalfa aphid, *Acyrtosiphon kondoi*;
 25 green cloverworm, *Plathypena scabra*; clover root curculio, *Sitona hispidulus*; alfalfa
 seed chalcid, *Brachophagus roddi*; tarnished plantbug, *Lygus lineolaris*; Say stink
 bug, *Chlorochroa sayi*; velvetbean caterpillar, *Anticarsia friegiperda*, alfalfa weevil,
Hypera postica; fall annyworm, *Spodoptera*; potato leafhopper, *Empoasca jabae*;
 soybean looper, *Pseudolusia includens*; Three cornered alfalfa hopper, *Spissistilus*
 30 *jestinus*; etc. Additional insect pests are known in the art. See, for example, Many B.
 Stoetzel (1989) Common Names of Insects & Related Organisms, Entomological
 Society of America, herein incorporated by reference.

In addition, polynucleotides or polypeptides, agonists or antagonists of the
 invention may be useful in detecting, preventing, and/or conferring resistance to other
 35 plant pests – including, but not limited to, insects, herbacious species, fungi, bacteria,
 viruses, and other pests disclosed herein. For example, polynucleotides or
 polypeptides, agonists or antagonists of the invention may, either directly, or
 indirectly, modulate (preferably decrease) the biosynthesis, assimilation, and/or

5 concentration of a nutrient essential for pest survival. Alternatively, polynucleotides or polypeptides, agonists or antagonists of the invention may modulate (preferably increase) the biosynthesis of a metabolite and/or protein capable of decreasing the biosynthesis, assimilation, and/or concentration of a nutrient essential for pest survival. One example concerns the observation that plants harboring the cholesterol oxidase gene were resistant to boll weevil (*Anthonomus grandis grandis* Boheman) larvae (Purcell, JP., *Biochem Biophys Res Commun*, 196(3):1406-13 (1993)). Since cholesterol is an essential nutrient for most organisms, including insects, polynucleotides or polypeptides, agonists or antagonists of the invention capable of modulating levels and/or activity of cholesterol degrading enzymes, including cholesterol oxidase, would be useful in conferring resistance to plant pests. In another example, plants expressing certain latherogens have been shown to confer resistance to some pathogens. Such latherogens have been identified as being able to interfere with collagen synthesis and/or aggregation.

In another embodiment, the polynucleotides or polypeptides, agonists or antagonists of the invention, can be used, either directly or indirectly, to detect, prevent, and/or confer resistance to any of the pests and/or diseases described herein through direct topical application to the plants for which protection is desired. For example, several protein- and/or peptide-based commercial pesticides have recently been introduced into the marketplace that provide broad pesticide resistance to pests when applied directly to the plant – Messenger, from Eden Bioscience, and Actigard, from Novartis Crop Protection (*Nat. Biotech.*, 18:595 (2000)). Messenger is based upon the *Erwinia amylovora* harpin gene (Wei, Z., *Science*, 257:85-88 (1992)). Such protein/peptide-based pesticides have been shown to activate the systemic acquired resistance (SAR) pathway in plants. The invention encompasses additional modes of action for conferring resistance to plant pests, though direct or indirect activation of the SAR pathway by the polynucleotides or polypeptides, agonists or antagonists of the invention, is preferred. The method of topical application, including any formulation requirements, will vary based the unique characteristics of each polynucleotide or polypeptide, agonist or antagonist of the invention. In addition, polynucleotides or polypeptides, agonists or antagonists of the invention that provides plant pathogen resistance may also have other beneficial uses, such as, for example, enhancing plant growth.

5 **Plant Defense**

A polypeptide or polynucleotide and/or agonist or antagonist of the present invention can be used to increase a plants defense mechanisms against either environmental or pathogenic stresses (e.g., viral, fungal, mycoplasma, bacterial, nematode, herbicidal, insecticidal, acid rain, drought, chemical, etc.). Such defense mechanisms may be a combination of structural characteristics (i.e., to serve as a physical barrier to inhibit a pathogen, for example, from entering or spreading throughout the plant), and biochemical reactions either on the scale of the whole plant or of individual cells (e.g., producing substances that are either toxic to the pathogen, or create an environment that is non-permissive for pathogen survival, etc.).

Structurally, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for increasing the number of trichomes, increasing the thickness and/or composition of wax secretions or the waxy layer, increasing the thickness and/or composition of the cuticle, altering the structure of the epidermal cell wall, altering the size, shape, and/or location of the stomata and lenticels, inducing the plant to create or increase a layer of thick-walled cells (e.g., cork cell layer, etc.), increasing the thickness and/or composition of the outer epidermal cell wall, inducing the formation of an abscission layer, induce the formation of tyloses, induce the production and/or deposition of gums, inducing the thickening of the outer parenchyma cell layer of the cell wall, inducing the thickening of the cell wall, inducing the deposition of callose papillae in the inner layer of the cell wall, inducing a necrotic or hypersensitive defense reaction in cells and/or tissues (i.e., cell death), inducing the polymerization of oxidized phenolic compounds into lignin-like substances to structurally interfere with pathogen development, and/or inducing a cytoplasmic defense reaction.

Biochemically, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for releasing pathogenic inhibitors into the plants environment, releasing fungitoxic exudates, and/or releasing phenolic compounds (e.g., protocatechic acid, catechol, etc.). Alternatively, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for increasing the synthesis of phenolic compounds (e.g., chlorogenic acids, caffeic acids, scopoletin, oxidation products of phenolic compounds, phytoalexins (see, Bell, et al., Ann. Rev. Plant Physiol, 32, 1981, for specific examples of phytoalexins), phaseolin, rishitin, kievitone, pisatin, glyceollin, gossypol, capsidiol, etc.), tannins, and/or

5 saponins (e.g., tomatine, avenacin, etc.) within the cells and tissues of the plant. Alternatively, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for increasing the expression of plant hydrolytic enzymes (e.g., glucanases, chitinases, etc.) that may cause degradation of the pathogen cell wall, etc.

10 In another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for inhibiting the expression of recognition factors essential for host-pathogen interaction (e.g., specific oligosaccharides, carbohydrate moieties, receptors, ligands, proteins, glycoproteins, lectins, etc.). For example, a polypeptide or polynucleotide and/or agonist or
15 antagonist of the present invention, may be useful for inhibiting the expression of a protein that serves as a target for a pathogenic toxin, thus rendering the host insensitive to the toxin.

In another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful in inhibiting the ability of the plants
20 metabolic machinery to complete essential steps required for a competent pathogenic response (e.g., inhibiting the ability of plant ribosomes to recognize the pathogens nucleic acid, such as a viral nucleic acid; and/or inhibiting the ability of the plants DNA polymerase machinery to recognize and/or synthesize pathogenic DNA; or inhibiting the plants ability to catalyze a specific enzymatic step essential to eliciting a
25 pathogenic response, etc.)

In yet another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for inhibiting either the production or transport or retention of essential nutrients required for a permissive pathogenic infection (e.g., inhibiting the transport of non-essential minerals or vitamins required
30 for a pathogenic response, etc.).

In one embodiment of the invention, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for increasing the expression or activity of phenol oxidizing enzymes (e.g., polyphenoloxidases, peroxidase, etc.), increasing the expression or activity of phenylalanine ammonia
35 lyase, increasing the activity or expression of proteins capable of forming pectin salts or pectin complexes, etc.

In a further embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for either directly or indirectly

5 inhibiting the activity of a pathogenic protein essential to eliciting an infection (e.g.,
inhibiting the enzymatic activity of the protein, such as for a hydrolytic enzyme, for
example, inhibiting the proteins ability to bind to a receptor or ligand, inhibiting
protein-protein or protein-DNA interactions of the pathogenic protein, etc.).
Specifically, a polypeptide or polynucleotide and/or agonist or antagonist of the
10 present invention, may be useful for either directly or indirectly inhibiting wildfire
toxin, chlorosis-inducing toxins, tabtoxin, phaseoloyoxin, rhizobitoxine, wilt-inducing
bacterial polysaccharides, amylovorin, glycopeptide toxins, peptide toxins,
syringomycin, tagetitoxin, helminthosporoside, victorin, helminthosporium maydis T-
toxin, helminthosporium carbonum toxin, periconia circinata toxin, phyllosticta
15 maydis toxin, alternaria toxins, fusarial wilt toxins, ophiobolin, helminthosporal,
terpinoid toxins, fusicocin, pyricularin, colletotol, alternaric acid, tentoxin,
phytotoxins, zinniol, tentoxin, ascochitine, diaporthin, skyrin, Didymella applanata
toxin, Myrothecium roridum toxin, Leptosphaerulina briosiana toxin, Alternaria
tenuis phenolic toxins, Cercospora beticola toxin, Verticillium albo-atrum toxin,
20 Phytophthora nicotianae var. parasitica toxin, Phytophthora megasperma var. sojae
toxin, Ceratocystis ulmi toxins, peptidorhamnomannan, Stemphylium botryosum
toxins, stemphylin, stemphyloxin, Pyrenophora teres toxins, N-(2-amino-2-
carboxyethyl) aspartic acid, aspergillomarasmine A, and Rhynchosporosides toxins,
for example.

25 In another embodiment, a polypeptide or polynucleotide and/or agonist or
antagonist of the present invention, may be useful for either increasing or inducing the
production of cyanogenic glycosides or esters, increasing the activity or expression of
hydrolytic enzymes capable of hydrolyzing cyanogenic glycosides or esters,
increasing the activity or expression of enzymes capable of releasing cyanide into
30 plant cells and tissues, increasing the activity or expression of enzymes capable of
detoxifying cyanide (e.g., formamide hydro-lyase, etc.) and/or increasing the
expression of b-proteins, etc.

In yet a further embodiment, a polypeptide or polynucleotide and/or agonist or
antagonist of the present invention, in addition to the non-coding 5' or 3' region of
35 any of the polynucleotides of the present invention, and/or fragments thereof, may be
useful for producing viral coat proteins, bacterial or fungal proteins, lipoproteins,
polysaccharides, yeast RNA, polyanions, polyacrylic acid, salicyic acid, and/or 2-
chloroethylphosphonic acid, in the plant, for example. Such proteins and/or

5 compounds have been shown to elicit a local resistance response to plant pathogens when applied systemically or topically to the plant.

In another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for either increasing or inducing the production of secondary metabolites, which include, but are not limited to the
10 following: acetyl salicylic acid, aconitine, atropine, cytosine, germinine, cardiac glycosides (e.g., calotropin, oleandrin, etc.), linarine, quinine, atropine, taxine, cicutoxin, hyoscyamine, pyrethrin, rotenone, camphor, etc.

In another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for either increasing or inducing the
15 production of non-protein amino acids, which include, but are not limited to the following: b-cyanoalanine, azetidine 2-carboxylic acid, canavanine, 3,4-dihydroxyphenylalanine, etc.

In another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention, may be useful for either increasing or inducing the
20 production of terpenes, which include, but are not limited to the following: 1.8 cineole, camphor, a-pinene, b-pinene, camphene, thujone, etc.

In yet another embodiment, a polypeptide or polynucleotide and/or agonist or antagonist of the present invention may either directly or indirectly inhibit the infectious agent, without necessarily increasing the plants defense mechanisms.

25 The present invention encompasses the application of one, two, three, four, or more, including any combination thereof, of any of the methods of increasing plant defense mechanisms against either an environmental or infectious agent described above and elsewhere herein. Additional methods of increasing a plants defense mechanisms are known in the art. Additionally, a list of compounds and/or proteins
30 that could serve as targets for increased production or expression by the use of a polynucleotide or polypeptide of the present invention to increase a plants defense mechanisms are known in the art (see, for example, Agrious, N.C., supra; Goodman, R.N., in "The Biochemistry and Physiology of Plant Disease", University of Missouri Press, Columbia, 1986; and Lambers, H., et al., in "Plant Physiological Ecology",
35 Springer-Verlag, New York, (1998); which are hereby incorporated herein by reference in their entirety).

5 **Plant Hormones**

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention can be used to modulate the hormone levels within a plant (including any of its cells, tissues, and/or organs, etc.). Examples of hormones that may be modulated by the present invention, either directly or indirectly, generally include, but are not
10 limited to, the following: auxins, indoleacetic acid, gibberellins, cytokinins, ethylene, abscisic acid, polyamines, jasmonates, tuberonic acid, salicylic acid, systemin, brassinolides, zeatin; and specifically, indole-3-acetic acid, indole-3-butyric acid, 4-chloroindole-3-acetic acid, indole-3-acetyl-1-O-B-D-glucose, indole-3-acetyl-myoinositol, jasmonic acid, methyl jasmonate, kinetin, including any known derivatives of
15 the hormones described above, etc. In this context, modulate should be applied to mean a quantitative, or qualitative increase, decrease, induction, or termination, of the expression levels of any of the aforementioned hormones. Additional examples of plant hormones are known in the art (see, for example, Davies, P.J., in "Plant Hormones: Physiology, Biochemistry, and Molecular Biology", Kluwer Academic
20 Publishers, Boston, 1995; which is hereby incorporated by reference in its entirety herein).

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant auxin levels, would necessarily be capable of the following, non-limiting, effects on a plant: stimulating cell enlargement,
25 stimulating stem growth, stimulating cell division in the cambium, stimulating differentiation of phloem and xylem, stimulating root initiation on stem cuttings, stimulating the development of branch roots, stimulating the differentiation of roots, mediating the bending (tropistic) response of shoots and roots to gravity and light, repression of lateral buds, delay of leaf senescence, inhibition or promotion of leaf and
30 fruit abscission (via ethylene), induction of fruit setting and growth, enhancement of assimilate transport via phloem, delay of fruit ripening, promotion of flowering in Bromeliads, stimulating flower growth, promotion of femaleness in dioecious flowers, and stimulating the production of ethylene, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present
35 invention capable of modulating plant gibberellin levels, would necessarily be capable of the following, non-limiting, effects on a plant: stimulating cell division and cell elongation, inducing hyperelongation, inducing bolting, inducing stem elongation in response to long days, inducing germination, inducing germination in seeds in the

5 absence of stratification or hardening, stimulating production of α -amylase, inducing fruit setting and growth, and inducing maleness in dioecious flowers, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant cytokinin levels, would necessarily be capable of the following, non-limiting, effects on a plant: inducing cell division in the
10 presence of auxin, inducing cell division in crown gall tumors, inducing cell division in apical meristem, inducing cell division in rapidly dividing cells, promoting shoot initiation, inducing bud formation, inducing growth of lateral buds, releasing lateral bud growth from apical dominance, inducing cell enlargement, inducing leaf expansion, enhancing stomatal opening, stimulating the accumulation of chlorophyll,
15 inducing the conversion of etioplasts to chloroplasts, and delaying leaf senescence, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant ethylene levels, would necessarily be capable of the following, non-limiting, effects on a plant: releasing the plant from dormancy,
20 inducing shoot and root growth and differentiation, inducing adventitious root formation, inducing leaf and fruit abscission, inducing flowering, inducing femaleness in dioecious flowers, inducing flower opening, inducing flower and leaf senescence, and inducing fruit ripening, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant abscisic acid levels, would necessarily be
25 capable of the following, non-limiting, effects on a plant: inducing stomatal closure, inhibition of shoot growth, inducing storage protein synthesis in seeds, inhibition of α -amylase production in germinating cereal grains, induction of some aspects of dormancy, and induction of proteinase inhibitor synthesis, for example.

30 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant polamine levels, would necessarily be capable of the following, non-limiting, effects on a plant: regulation of growth and development of plant cells and tissues, modulating the synthesis of macromolecules, modulating the activity of macromolecules, stabilizing cellular plasma membrane,
35 decreasing leakage of betacyanin from wounded tissue, preservation of thylakoid structure in excised barley leaves, counteraction of hormone-induced affects on the cell membrane, binding to nucleic acids, protection of nucleic acids from alkylating agents, controlling chromosome condensation, controlling nuclear membrane

5 dissolution during pre-prophase, and modulating the structure and function of tRNA's, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant jasmonate levels, would necessarily be capable of the following, non-limiting, effects on a plant: inhibition of plant growth, inhibition
10 of seed germination, promotion of senescence, promotion of abscission, promotion of tuber formation, promotion of fruit ripening, promotion of pigment formation, promotion of tendril coiling, induction of proteinase inhibitors, and inhibit insect infestation, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present
15 invention capable of modulating plant salicylic acid levels, would necessarily be capable of the following, non-limiting, effects on a plant: induction of thermogenesis, providing resistance to pathogens via induction of pathogenesis related proteins, enhancement of flower longevity, inhibition of ethylene biosynthesis, inhibition of seed germination, inhibiting the wound response, counteracting the plants response to
20 abscisic acid, for example.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant brassinosteroid levels, would necessarily be capable of the following, non-limiting, effects on a plant: promotion of stem elongation, inhibition of root growth, inhibition of root development, promotion of
25 ethylene biosynthesis, and promotion of epinasty, for example.

The polynucleotides or polypeptide and/or agonist or antagonist of the present invention may modulate one, two, three, or more, or any combination of the above, hormones in a plant. Additional effects of hormones on a plant, including its cells, tissues, and organs are known in the art and the aforementioned plant hormone effects
30 should not be construed as limiting the utility of any of the polynucleotide or polypeptides of the present invention.

Regeneration

A polynucleotide or polypeptide and/or agonist or antagonist of the present
35 invention can be used to differentiate, proliferate, and/or attract cells, leading to the regeneration of tissues. (See, Science 276:59-87 (1997).) The regeneration of tissues could be used to repair, replace, and/or protect tissue damaged by environmental insults (e.g., herbicidal, photobleaching, acid rain, drought, nematodes, insects,

5 chemical, etc.), disease (e.g., fungal, viral, bacterial, mycoplasmal, etc.), necrosis, hypersensitive reaction, and/or cytokine damage.

Tissues that could be regenerated using polynucleotides or polypeptides of the present invention include tissues (e.g., apical meristem, lateral shoot, lateral bud, leaf, pith, vascular cambium, stem, phloem, xylem, cortex, epidermis, lateral root, root
10 meristem, cuticle, etc.), in addition to cellular organelles and constituents (vacuole, mitochondrion, chloroplast, plastid, lysosomes, peroxisomes, glyoxysomes, cytoplasm, endoplasmic reticulum, ribosomes, vacuolar membrane, nucleus, nuclear membrane, plasmodesmata, scherosomes, microbodies, primary cell wall, etc.).

15 Nutrients

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention can be used to modulate the plants nutritional status through a number of different mechanisms. For example, A polynucleotide or polypeptide and/or agonist or antagonist of the present invention can be used to modulate the plants ability to
20 retain a particular nutrient, to modulate the plants ability to synthesize a particular nutrient, to modulate the plants ability to assimilate a nutrient, to modulate the plants ability to absorb or uptake a particular nutrient, to modulate the plants ability to transport a particular nutrient, to modulate the plants ability to store a particular nutrient, to modulate the plants ability to survive under nutrient deficiencies, and to
25 prevent, detect, and/or provide resistance to nutrient deficiency symptoms and traits.

Specific examples of nutrients that may be modulated in a plant by a polynucleotide or polypeptide and/or agonist or antagonist of the present invention include the following, non-limiting, nutrients: carbon, hydrogen, oxygen, nitrogen, phosphorus, sulfur, potassium, calcium, magnesium, boron, chlorine, copper, iron,
30 manganese, zinc, molybdenum, cobalt, selenium, silicon, sodium, nickel, water, carbon dioxide, in addition to metabolic by-products, etc. Additional nutrients essential to maintaining plant homeostasis are known in the art.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant boron levels, may be capable of preventing,
35 detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of plant boron deficiency: terminal leaf necrosis, premature leaf abscission layer formation, terminal shoot internode shortening, blackening and/or death of

5 apical meristem tissue, shortening of root shoots, plant dwarfing, plant stunting, impairment of flower development, impairment of seed development, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant calcium levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting,
10 symptoms of plant calcium deficiency: chlorotic leaves, leaf curling, leaf rolling, degradation of meristematic tissues in stems and roots, meristematic tissue death, decreased root development, decreased root fiber content, decreased fruit development, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present
15 invention capable of modulating plant chlorine levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of plant chlorine deficiency: leaf tip wilting, leaf chlorosis, leaf bronzing, basipetal leaf necrosis proximal to areas of wilting, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present
20 invention capable of modulating plant copper levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of plant copper deficiency: terminal shoot wilting, terminal shoot death, fading of leaf color, reduction of carotene in plant cells and tissues, reduction of other pigments in plant cells and tissues, etc.

25 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant iron levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of iron deficiency: interveinal white chlorosis of young leaves first, chlorosis of aerial tissues, aerial tissue necrosis, bleaching of leaves, scorching of
30 leave margins and tips, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant magnesium levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of magnesium deficiency: mottling chlorosis with green veins and
35 leaf web tissue yellow or white on old leaves first, wilting of leaves, formation of leaf abscission layer in the absence of the wilting stage, necrosis of plant cells and tissues, etc.

5 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant manganese levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of manganese deficiency: mottling chlorosis with green veins and leaf web tissue yellow or white on young leaves first, then spreading to old leaves,
10 yellowish green stem, hardening and/or wooding of stem, reduction of carotene, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant molybdenum levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of molybdenum deficiency: light yellow chlorosis of leaves,
15 failure of leaf blade expansion, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant nitrogen levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of nitrogen deficiency: stunting plant growth of young plants, yellowish
20 green leaves in young plants, light green leaves in older leaves followed by yellowing and drying or shedding, increased accumulation of anthocyanins in veins, thin stem, spindely appearance of plant, reduced flowering, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant phosphorus levels, may be capable of
25 preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of phosphorus deficiency: stunting of young plants, dark blue-green leaves with purplish undertones, slender stems, increased accumulation of anthocyanin in leaves, necrosis of leaves, cessation of meristematic growth, decreased rate of fruit ripening, plant dwarfing at maturity, etc.

30 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant potassium levels, may be capable of preventing, detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of potassium deficiency: dark green leaves, pale green monocotyledon leaves, yellowing streaking of monocotyledon leaves, marginal
35 chlorosis of leaves, necrosis of leaves appearing first on old leaves, wrinkling of veins, currugating of veins, crinkling of veins, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant sulfur levels, may be capable of preventing,

5 detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of sulfur deficiency: light green to yellow leaves appearing first along veins of young leaves, slender stems, etc.

A polynucleotide or polypeptide and/or agonist or antagonist of the present invention capable of modulating plant zinc levels, may be capable of preventing,
10 detecting, alleviating, and/or conferring resistance to the following, non-limiting, symptoms of zinc deficiency: chlorosis of leaves and/or necrosis of leaves affecting young leaves first, rosetting, premature formation of abscission layer of leaves, whitish chlorotic streaks between veins in older leaves, whiting of upper leaves in monocotyledons, chlorosis of lower leaves in dicotyledons, etc.

15 Additional symptoms of plant nutrient deficiencies are known in the art (see for example, Noggle, G.R., and Fritz, G.J., in "Introductory Plant Physiology", 2nd edition, Prentice-Hall, Inc., Englewood Cliffs, 1983).

In a specific embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may be able to modulate plant nutrient levels
20 either directly or indirectly by increasing the activity, kinetics, and/or expression of transport proteins, ion channels, and/or ion carrier proteins.

In another embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may be able to modulate plant nutrient levels by
25 either directly or indirectly increasing or inducing the secretion of mineral solubilizing or mineral stabilizing compounds or chelating compounds (e.g., citric acid, malic acid, isocitric acid, etc.). Alternatively, the secreted compound may be an organic chelating compound (e.g., phytochelator, see for example, Cakmak et al., Plant Soil, 180:183-189, (1996)). Alternatively, the secreted compound is a root
30 exudate, such as an organic acid (e.g., lactic, acetic, formic, pyruvic, succinic, tartaric, oxalic, citric, isocitric, aconitic, etc.), carbohydrate, amino acid, or polysaccharide capable of assimilating carbon (see, for example, Paul, E.A., and Clark, F.E., in "Soil microbiology and biochemistry", Academic Press, San Diego, (1989)).

In another embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may be able to modulate plant nutrient levels by
35 modulating, either directly or indirectly, the activity, kinetics, and/or expression of phosphatase enzymes, nitrate reductase enzymes, citrate synthesis enzymes, etc.

In another embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may be able to modulate plant nutrient levels by

5 modulating the active transport and/or passive transport mechanisms of the plant. Alternatively, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may be able to modulate plant nutrient levels by modulating the inter- and intra-tissue and/or cellular transport of nutrients in the plant (e.g., transport through the phloem, xylem, desmosomes, etc.). Additional mechanisms of
10 modulating plant nutrient transport are known in the art (see, for example, Lambers, H., et al., in "Plant Physiological Ecology", Springer-Verlag, New York, (1998); which is hereby incorporated herein by reference in its entirety).

Biotic Associations

15 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention may increase the plants ability, either directly or indirectly, to initiate and/or maintain biotic associations with other organisms. Such associations may be symbiotic, nonsymbiotic, endosymbiotic, macrosymbiotic, and/or microsymbiotic in nature. In general, a polynucleotide or polypeptide and/or agonist or antagonist of the
20 present invention may increase the plants ability to form biotic associations with any member of the fungal, bacterial, lichen, mycorrhizal, cyanobacterial, dinoflagellate, and/or algal, kingdom, phylums, families, classes, genuses, and/or species.

Specific, non-limiting, examples of organisms known to form biotic associations with a plant are: ectomycorrhizas (e.g., members of the Dimerocarpaceae,
25 Pinaceae, Fagaceae, Myrtaceae, Salicaceae, Betulaceae, Fabaceae, etc.), endomycorrhizas, vesicular arbuscular mycorrhiza (e.g., members of the Glomales), nonmycorrhizal (e.g., members of the Brassicaceae, Caryophyllaceae, Chenopodiaceae, Lecythideaceae, Proceae, Restionaceae, Sapotaceae, Urticaceae, Zygophyllaceae, etc.), symbiotic N₂ fixing organisms (e.g., members of the
30 Rhizobium, Bradyrhizobium, Sinorhizobium, Mesorhizobium, Azorhizobium, etc.), nonsymbiotic N₂ fixing organisms (e.g., Azospirillum, etc.), endosymbiotic organisms (e.g., members of the Clavicipitaceae, Ascomycetes, etc.), etc. Additional organisms capable of forming biotic associations with plants are known in the art and are encompassed by the invention (see, for example, Lambers, H., et al., in "Plant
35 Physiological Ecology", Springer-Verlag, New York, (1998); Raven, P.H., et al., in "Biology of Plants", 5th Edition, Worth Publishers, New York, (1992); which are hereby incorporated herein by reference in its entirety

5 The mechanism by which a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may increase the plants ability, either directly or indirectly, to initiate and/or maintain biotic associations is variable, though may include, modulated secretions of organic acids, phenolic compounds, nutrients, or the increased expression of a protein required for host-biotic organisms interactions (e.g.,
10 a receptor, ligand, etc.). Additional mechanisms are known in the art and are encompassed by the invention.

 In an alternative embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may decrease a plants ability to form biotic associations with a plant. Such a decrease may be due to increasing the plants ability
15 to utilize, obtain, store, and/or synthesize essential nutrients that the biotic organism supplied to the plant through the association.

 Mechanisms by which a plant could inhibit colonization of biotic associating organisms are known in the art (see, for example, Smith, S.E., and Read, D.J., in “Mycorrhizal Symbiosis”, Academic Press, London, 1997, etc.). For example, in low
20 phosphorus containing soils, a plant may form beneficial symbiosis colonization with a species capable of providing phosphorus. However, in soils of high phosphorus content, such colonization could be inhibited by the plant. Thus, a polynucleotide or polypeptide of the invention could increase the plants ability to live in phosphorus depleted soils by enabling the plant to assimilate phosphorus through a mechanism
25 previously not endogenous to the plant, for example.

Pheromones

 In another embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may increase the plants ability to synthesize and/or
30 release a pheromone. Such a pheromone may attract predatory organisms to the plant that may feed on plant pests or infectious agents. For example, recent studies of the aphid *Acyrtosiphon pisum* have shown that feeding by the aphid alters the composition of volatiles released by the plant and that these compounds act as synomones for the aphids foraging parasitoid, *Aphidius ervi*. Additional studies have
35 shown that other pheromones released by the plant in response to aphid foraging, namely(E)-beta-farnesene, may also attract other aphid predators, such as the lacewing *Chrysoperla carnea* and the seven-spot ladybird, *Coccinella septempunctata*.

5 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention may modulate the biosynthesis and/or release of pheromones, volatiles having pheromone-like effects, volatiles having kairomone effects, volatiles having synomones effects, and/or may releases (E)-beta-farnesene specifically. Alternatively, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may modulate the release of pheromones, indirectly, through the induction of the following, non-limiting, compounds, compounds from the octadecanoid signalling pathway, structurally non-related amino acid conjugates such as the bacterial phytotoxin coronatine, the synthetic indanoyl-isoleucine, or amino acid conjugates of linolenic, and/or volatile terpenoids, such as members of the sesqui- and diterpenoids. Preferably, any of the pheromones, and/or volatiles released from the plant, or induced, by a polynucleotide or polypeptide and/or agonist or antagonist of the invention have behavioral effects on predators and/or pests of plants. Other examples of known pheromones or volatiles released by a plant having behavioral modulating effects on plant pest predators and/or plant pests are known in the art (see, Boland W., et al., Novartis Found Symp, 223:110-26 (1999); Wadhams L.J., Novartis Found Symp, 223:60-7 (1999); Tumlinson JH, et al., Novartis Found Symp, 223:95-105 (1999)).

 In yet another embodiment, a polynucleotide or polypeptide and/or agonists or antagonists of the present invention may modulate the biosynthesis of furanones. Furanones are naturally occurring compounds found in a variety of plants that have been shown to have pheromone-like effects, anti-bacterial effects, anti-viral effects, etc., on a variety of organisms. For example, 5-Methyl-4-hydroxy-3(2H)-furanone is a male pheromone in the cockroach *Eurycolis florionda* (Walker) and the 2,5-dimethyl derivative deters fungal growth on strawberries and is an important component of the attractive aroma of the fruit. The red seaweed *Delisea pulchra* (Greville) Montagne produces a range of brominated furanones which prevent colonisation of the plant by bacteria by interfering with the acylated homoserine lactone (AHL) signalling system used by the bacteria for quorum sensing. In addition, furanones have been shown to have mutagenic properties in bacteria and viruses, and thus could serve as anti-bacterials and anti-virals (Colin, Slaughter J, Biol Rev Camb Philos Soc, 74(3):259-76 (1999)).

5 **Chemotaxis**

 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention may have chemotaxis activity. A chemotaxic molecule attracts or mobilizes cells to a particular site in the plant or animal body, such as inflammation, infection, or site of hyperproliferation. The mobilized cells can then fight off and/or heal the particular trauma or abnormality.

 A polynucleotide or polypeptide and/or agonist or antagonist of the present invention may increase chemotaxic activity of particular cells. These chemotactic molecules can then be used to detect, prevent, and/or alleviate inflammation, infection, hyperproliferative diseases, disorders, and/or conditions, or any plant disorder by increasing the number of cells targeted to a particular location in the plants body. For example, chemotaxic molecules can be used to treat, prevent, and/or diagnose wounds and other trauma to tissues by attracting cells to the injured location.

 It is also contemplated that a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may inhibit chemotactic activity. These molecules could also be used to detect, prevent, and/or alleviate diseases, disorders, and/or conditions. Thus, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention could be used as an inhibitor of chemotaxis.

 In an alternative embodiment, a polynucleotide or polypeptide and/or agonist or antagonist of the present invention may modulate the ability of a plant to sense the presence of plant neighbors (i.e., chemoperception). Such chemoperceptive modulation may come in the form of sensing the presence of plant hormones (e.g., jasmonate, etc.), heat exchange differential, and/or allelochemical detection (see, for example, Boller, *Ann. Rev. Plant Physiol. Plant Mol. Biol.*, 46:189-214, (1995)).

30 **Binding Activity**

 A polypeptide of the present invention may be used to screen for molecules that bind to the polypeptide or for molecules to which the polypeptide binds. The binding of the polypeptide and the molecule may activate (agonist), increase, inhibit (antagonist), or decrease activity of the polypeptide or the molecule bound. Examples of such molecules include antibodies, oligonucleotides, proteins (e.g., receptors), or small molecules.

 Preferably, the molecule is closely related to the natural ligand of the polypeptide, e.g., a fragment of the ligand, or a natural substrate, a ligand, a structural

5 or functional mimetic. (See, Coligan et al., Current Protocols in Immunology 1(2):Chapter 5 (1991).) Similarly, the molecule can be closely related to the natural receptor to which the polypeptide binds, or at least, a fragment of the receptor capable of being bound by the polypeptide (e.g., active site). In either case, the molecule can be rationally designed using known techniques.

10 Preferably, the screening for these molecules involves producing appropriate cells that express the polypeptide, either as a secreted protein or on the cell membrane. Preferred cells include cells from plants, yeast, or E. coli. Cells expressing the polypeptide (or cell membrane containing the expressed polypeptide) are then preferably contacted with a test compound potentially containing the
15 molecule to observe binding, stimulation, or inhibition of activity of either the polypeptide or the molecule. Moreover, the identification of such molecules may be obtained through the application of the yeast 2 or 3 hybrid system (see for example, Wallach D, et al., Curr Opin Immunol., 10(2):131-6, (1998); Young KH., Biol Reprod., 58(2):302-11, (1998); and Fernandes, PB., Curr Opin Chem Biol., 2(5):597-
20 603 (1998); which are hereby incorporated by reference in their entirety herein, including the methods disclosed and the references cited therein). Further, the identification of such molecules may be obtained through the application of additional screening technologies which include, but are not limited to, the following issued US Patents: 5,284,746; 5,576,210; 5,691,188; 5,846,819; and International Publication
25 No. WO 95/34646; which are hereby incorporated by reference in their entirety herein, including the methods disclosed and references cited therein.

The assay may simply test binding of a candidate compound to the polypeptide, wherein binding is detected by a label, or in an assay involving competition with a labeled competitor. Further, the assay may test whether the
30 candidate compound results in a signal generated by binding to the polypeptide.

Alternatively, the assay can be carried out using cell-free preparations, polypeptide/molecule affixed to a solid support, chemical libraries, or natural product mixtures. The assay may also simply comprise the steps of mixing a candidate compound with a solution containing a polypeptide, measuring polypeptide/molecule
35 activity or binding, and comparing the polypeptide/molecule activity or binding to a standard.

Preferably, an ELISA assay can measure polypeptide level or activity in a sample (e.g., biological sample) using a monoclonal or polyclonal antibody. The

5 antibody can measure polypeptide level or activity by either binding, directly or indirectly, to the polypeptide or by competing with the polypeptide for a substrate.

As an alternative approach for receptor identification, the labeled polypeptides can be photoaffinity linked with cell membrane or extract preparations that express the receptor molecule. Cross-linked material is resolved by PAGE analysis and
10 exposed to X-ray film. The labeled complex containing the receptors of the polypeptides can be excised, resolved into peptide fragments, and subjected to protein microsequencing. The amino acid sequence obtained from microsequencing would be used to design a set of degenerate oligonucleotide probes to screen a cDNA library to identify the genes encoding the putative receptors.

15 Moreover, the techniques of gene-shuffling, motif-shuffling, exon-shuffling, and/or codon-shuffling (collectively referred to as "DNA shuffling") may be employed to modulate the activities of polypeptides of the invention thereby effectively generating agonists and antagonists of polypeptides of the invention. See generally, U.S. Patent Nos. 5,605,793, 5,811,238, 5,830,721, 5,834,252, and
20 5,837,458, and Patten, P. A., et al., *Curr. Opinion Biotechnol.* 8:724-33 (1997); Harayama, S. *Trends Biotechnol.* 16(2):76-82 (1998); Hansson, L. O., et al., *J. Mol. Biol.* 287:265-76 (1999); and Lorenzo, M. M. and Blasco, R. *Biotechniques* 24(2):308-13 (1998) (each of these patents and publications are hereby incorporated by reference). The techniques of DNA shuffling are known in the art and more
25 particularly described elsewhere herein.

All of the assays referenced above, and elsewhere herein, can be used as diagnostic or prognostic markers. The molecules discovered using these assays may be used to detect, prevent, and/or confer resistance to a disease or to bring about a particular result in an organism (e.g., vessel growth, etc.) by activating or inhibiting
30 the polypeptide/molecule. Moreover, the assays can discover agents which may inhibit or enhance the production of the polypeptides of the invention from suitably manipulated cells or tissues.

Therefore, the invention includes a method of identifying compounds which bind to the polypeptides of the invention comprising the steps of: (a) incubating a
35 candidate binding compound with the polypeptide; and (b) determining if binding has occurred. Moreover, the invention includes a method of identifying agonists/antagonists comprising the steps of: (a) incubating a candidate compound

- 5 with the polypeptide, (b) assaying a biological activity, and (b) determining if a biological activity of the polypeptide has been altered.

Antisense and Ribozyme (Antagonists)

10 In preferred embodiments, the invention encompasses antagonists which correspond to the polynucleotide sequences shown in SEQ ID NO:X, the complementary strand thereof, and/or to the polynucleotide sequences contained within the deposited clone. Antisense technology results in modulation (i.e., complete or partial inhibition), of the expression of a particular protein through direct inhibition of the proteins mRNA. Antisense nucleic acids may be in the form of DNA, RNA,
15 PNA, triple helix, quad helix, a chimeric mixture of any of these aforementioned types (e.g., DNA:RNA, PNA:RNA, PNA:DNA, etc.), and may be single or double stranded. Antisense nucleic acids modulate gene expression by binding to the RNA of the gene of interest, effectively inhibiting translation. Such interactions may rely follow typical Watson-Crick base pair recognition, or the case of a triple or quad
20 helix, may rely upon Hoogsteen basepair recognition.

The antisense nucleic acids may be transiently generated within the organism (e.g., sequence contained within an inducible or constitutively expressed vector introduced into the cells of an organism), stably generated within the organism (e.g., sequence contained within an inducible or constitutively expressed vector introduced
25 into the cells of an organism using transgenic methods, including viral integration, etc.) or may be exogenously administered. For a nucleic acid to serve an antisense role, it is only necessary that it has sequence homology to the sense RNA product of the gene of interest. A number of methods of administering antisense nucleic acids, their compositions, and designs are known in the art and encompassed by the invention (see for example, Agrawal S, et al., Mol Med Today. 2000 Feb;6(2):72-81; Yacyshyn BR, et al, Can J Gastroenterol. 1999 Nov;13(9):745-51; Mrsny RJ, J Drug Target. 1999;7(1):1-10; Toulme JJ, et al, Nucleic Acids Symp Ser. 1997;(36):39-41.), Okano, Neurochem., 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988); and Cooper SR, et al.,
35 Pharmacol Ther. 1999 May-Jun;82(2-3):427-35). Likewise, a number of methods have been developed regarding the application of triple helix antisense technology to modulating gene expression (see, for example, Gowers DM, et al, Nucleic Acids Res. 1999 Apr 1;27(7):1569-77; and Chan PP, et al., J Mol Med. 1997 Apr;75(4):267-82).

5 Antisense technology has wide-ranging applications in plants. For example, antisense RNA has been shown to effectively downregulate a variety of plant genes as described by Shimada, et al., *Theor. Appl. Genet.*, 86:665-672, (1993); Kull, et al., *J. Genet. Breed.*, 49:67-76, (1995)., Slabas and Elborough, WO 97/07222; Knutzon et al., *Proc. Natl. Acad. Sci. USA*, 89:2624-2628, (1992), and Baulcombe
10 DC., *Plant Mol Biol.* 1996 Oct;32(1-2):79-88)..

 The antisense nucleic acids of the invention comprise a sequence complementary to at least a portion of an RNA transcript of a gene of interest. However, absolute complementarity, although preferred, is not required. A sequence “complementary to at least a portion of an RNA,” referred to herein, means a
15 sequence having sufficient complementarity to be able to hybridize with the RNA, forming a stable duplex; in the case of double stranded antisense nucleic acids of the invention, a single strand of the duplex DNA may thus be tested, or triplex formation may be assayed. The ability to hybridize will depend on both the degree of complementarity and the length of the antisense nucleic acid. Generally, the larger the
20 hybridizing nucleic acid, the more base mismatches with a RNA sequence of the invention it may contain and still form a stable duplex (or triplex as the case may be). One skilled in the art can ascertain a tolerable degree of mismatch by use of standard procedures to determine the melting point of the hybridized complex.

 Antisense oligonucleotides that are complementary to the 5' end of the
25 message, e.g., the 5' untranslated sequence up to and including the AUG initiation codon, should work most efficiently at inhibiting translation. However, sequences complementary to the 3' untranslated sequences of mRNAs have been shown to be effective at inhibiting translation of mRNAs as well. See generally, Wagner, R., *Nature*, 372:333-335 (1994). Thus, oligonucleotides complementary to either the 5' -
30 or 3' - non- translated, non-coding regions of a polynucleotide sequence of the invention could be used in an antisense approach to inhibit translation of endogenous mRNA. Oligonucleotides complementary to the 5' untranslated region of the mRNA should include the complement of the AUG start codon. Antisense oligonucleotides complementary to mRNA coding regions are less efficient inhibitors of translation but
35 could be used in accordance with the invention. Whether designed to hybridize to the 5' -, 3' - or coding region of mRNA, antisense nucleic acids should be at least six nucleotides in length, and are preferably oligonucleotides ranging from 6 to about 50

5 nucleotides in length. In specific aspects the oligonucleotide is at least 10 nucleotides, at least 17 nucleotides, at least 25 nucleotides or at least 50 nucleotides.

The antisense oligonucleotide may be modified at the base moiety, sugar moiety, or phosphate backbone. The oligonucleotide may include other appending groups such as peptides, or agents facilitating transport across the cell membrane (see, 10 e.g., Letsinger et al., 1989, Proc. Natl. Acad. Sci. USA 86:6553-6556; Lemaitre et al., 1987, Proc. Natl. Acad. Sci.84:648-652; PCT Publication No. WO 88/09810, published Dec. 15, 1988) or blood-brain barrier (see, e.g., PCT Publication No. WO 89/10134, published Apr. 25, 1988), hybridization-triggered cleavage agents (see, e.g., Krol et al., 1988, BioTechniques 6:958-976) or intercalating agents (see, e.g., 15 Zon, 1988, Pharm. Res. 5:539-549).

The oligonucleotide can also be a peptido-nucleic acid ("PNA") based on a linked N-(2-aminoethyl)glycine backbone to which normal DNA bases have been attached (Egholm et al., 1993, Nature 365:566-67). This PNA obeys specific Watson-Crick base pairing, but with greater free energy of binding and correspondingly higher 20 melting temperatures. Suitable oligomers may be constructed entirely from PNAs or from mixed PNA and DNA and/or RNA oligomers. In fact, PNA:DNA chimeras have increased solubility characteristics, as compared to DNA:DNA or DNA:RNA chimeras of the same sequence. Most notably, PNAs have the unique ability to displace one strand of a DNA double-helix thus making them highly suitable in 25 antisense applications (Uhlmann E., Biol Chem. 1998 Aug-Sep;379(8-9):1045-52).

In a preferred embodiment, the oligonucleotide comprises at least one modified phosphate backbone selected from the group consisting of a phosphorothioate, a phosphorodithioate, a phosphoramidothioate, a phosphoramidate, a phosphordiamidate, a methylphosphonate, an alkyl phosphotriester, and a 30 formacetal or analog thereof.

In another embodiment, the anti-sense oligonucleotide of the invention may comprise at least one modified base moiety which is selected from the group including, but not limited to, 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) 35 uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-

5 methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, 3-(3-amino-3-N-2-
10 carboxypropyl) uracil, and
2,6-diaminopurine.

In another embodiment, the anti-sense oligonucleotide comprises at least one modified sugar moiety selected from the group including, but not limited to, arabinose, 2-fluoroarabinose, xylulose, and hexose.

15 In yet another embodiment, the anti-sense oligonucleotide may be conjugated to another molecule, e.g., a peptide, hybridization triggered cross-linking agent, transport agent, hybridization-triggered cleavage agent, etc.

The anti-sense oligonucleotides of the invention may be synthesized by standard methods known in the art, e.g., by use of an automated DNA synthesizer
20 (such as are commercially available from Biosearch, Applied Biosystems, etc.). As examples, phosphorothioate oligos may be synthesized by the method of Stein et al. (1988, Nucl. Acids Res. 16:3209), methylphosphonate oligos can be prepared by use of controlled pore glass polymer supports (Sarin et al., 1988, Proc.Natl. Acad. Sci. USA 85:7448-7451), etc.

25 In a specific embodiment, the oligonucleotide comprises catalytic RNA, or a ribozyme (see, e.g., PCT International Publication WO 90/11364, published Oct. 4, 1990; Sarver et al., 1990, Science 247:1222-1225; Hasselhoff, et al., Nature 342:76-79 (1988)). Ribozymes have been used to downregulate gene expression, and more recently in the downregulation of plant proteins (seem e.g., PCT International
30 Publication WO 97/10328). In another embodiment, the oligonucleotide is a 2'-O-methylribonucleotide (Inoue et al., 1987, Nucl. Acids Res. 15:6131-6148), or a chimeric RNA-DNA analogue (Inoue et al., 1987, FEBS Lett. 215:327-330).

Other Activities

35 In another embodiment, the polypeptides, polynucleotides encoding these polypeptides, variants, and/or fragments thereof, may be used to inhibit gene expression using co-suppression methodology. The mechanism of co-suppression is not known, though its application to inhibiting plant gene expression has been

5 documented and described (e.g., Seymour, et al., *Plant. Mol. Biol.*, 23:1-9, (1993),
Brusslan, et al., *Plant Cell*, 5:667-677, (1993); Vaucheret, et al., *Mol. Gen. Genet.*,
248:311-317, (1995); and Jorgensen, et al., *Plant Mol. Biol.*, 31:957-973, (1996)).
Co-suppression involves creating a constitutively expressed vector construct
comprising, for example, a CaMV 35S promoter, the 5' coding region of a first gene
10 (R) for which inhibited expression is desired, in-frame and upstream from, the entire
coding region of a second gene (S) for which inhibited expression is desired, and a
terminator. Upon positive transformation of plants with this vector (i.e., transgenic
plants), no detectable mRNA expression will be detected for either R nor S (see
Seymour, supra).

15 The polypeptide or polynucleotides and/or agonist or antagonists of the
present invention may also increase or decrease the differentiation or proliferation of
protoplast cells, embryoblast cells, etc.

The polypeptide or polynucleotides and/or agonist or antagonists of the
present invention may be useful in modulating programmed cell death in plants, plant
20 cells, and eukaryotic cells and organisms, in general. Such modulation would either
be through direct or indirect interaction between a polypeptide or polynucleotides
and/or agonist or antagonists of the present invention with the gene or protein critical
in modulating programmed cell death in the organism. Specific targets of interaction
for programmed cell death, particularly in a plant, are provided in International
25 Publication Number WO 00/04173 (e.g., poly-ADP-ribose polymerase (PARP) genes,
specifically PARP genes of the ZAP class, etc.).

The polypeptide or polynucleotides and/or agonist or antagonists of the
present invention may also be used to modulate plant characteristics, such as stem
height, pigmentation, stress tolerance, etc. Similarly, polypeptides or polynucleotides
30 and/or agonist or antagonists of the present invention may be used to modulate plant
metabolism affecting catabolism, anabolism, processing, utilization, and storage of
energy.

Polypeptide or polynucleotides and/or agonist or antagonists of the present
invention may also be used as a food additive or preservative, such as to increase or
35 decrease storage capabilities, fat content, lipid, protein, carbohydrate, vitamins,
minerals, cofactors or other nutritional components.

5 Other Preferred Embodiments

Other preferred embodiments of the claimed invention include an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 50 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1.

10 Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the “5’ NT of Clone” and ending with the nucleotide at about the position of the “3’ NT of Clone” as defined for SEQ ID NO:X in Table 1.

15 Also preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the “5’ NT of Start Codon of Clone ORF” and ending with the nucleotide at about the position of the “3’ NT of Clone ORF” as defined for SEQ ID NO:X in Table 1.

20 Similarly preferred is a nucleic acid molecule wherein said sequence of contiguous nucleotides is included in the nucleotide sequence of SEQ ID NO:X in the range of positions beginning with the nucleotide at about the position of the “5’ NT of the First AA of the Signal Peptide” and ending with the nucleotide at about the position of the “3’ NT of Clone ORF” as defined for SEQ ID NO:X in Table 1.

25 Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 150 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

Further preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least about 500
30 contiguous nucleotides in the nucleotide sequence of SEQ ID NO:X.

A further preferred embodiment is a nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the nucleotide sequence of SEQ ID NO:X beginning with the nucleotide at about the position of the “5’ NT of Clone ORF” and ending with the nucleotide at about the position of the “3’ NT of Clone
35 ORF” as defined for SEQ ID NO:X in Table 1.

A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence of SEQ ID NO:X.

5 Also preferred is an isolated nucleic acid molecule which hybridizes under stringent hybridization conditions to a nucleic acid molecule, wherein said nucleic acid molecule which hybridizes does not hybridize under stringent hybridization conditions to a nucleic acid molecule having a nucleotide sequence consisting of only A residues or of only T residues.

10 Also preferred is a composition of matter comprising a DNA molecule which comprises a cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the material deposited with the American Type Culture Collection and given the ATCC Deposit Number shown in Table 1 for said cDNA Clone Identifier.

15 Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a sequence of at least 50 contiguous nucleotides in the nucleotide sequence of a cDNA clone identified by a cDNA Clone Identifier in Table 1, which DNA molecule is contained in the deposit given the ATCC Deposit Number shown in Table 1.

20 Also preferred is an isolated nucleic acid molecule, wherein said sequence of at least 50 contiguous nucleotides is included in the nucleotide sequence of the complete open reading frame sequence encoded by said cDNA clone.

 Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 150 contiguous
25 nucleotides in the nucleotide sequence encoded by said cDNA clone.

 A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to sequence of at least 500 contiguous nucleotides in the nucleotide sequence encoded by said cDNA clone.

30 A further preferred embodiment is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to the complete nucleotide sequence encoded by said cDNA clone.

 A further preferred embodiment is a method for detecting in a biological sample a nucleic acid molecule comprising a nucleotide sequence which is at least
35 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the

5 deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1; which method comprises a step of comparing a nucleotide sequence of at least one nucleic acid molecule in said sample with a sequence selected from said group and determining whether the sequence of said nucleic acid molecule in said sample is at least 95% identical to said selected sequence.

10 Also preferred is the above method wherein said step of comparing sequences comprises determining the extent of nucleic acid hybridization between nucleic acid molecules in said sample and a nucleic acid molecule comprising said sequence selected from said group. Similarly, also preferred is the above method wherein said step of comparing sequences is performed by comparing the nucleotide sequence
15 determined from a nucleic acid molecule in said sample with said sequence selected from said group. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

A further preferred embodiment is a method for identifying the species, tissue or cell type of a biological sample which method comprises a step of detecting nucleic
20 acid molecules in said sample, if any, comprising a nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the
25 deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for identifying the species, tissue or cell type of a biological sample can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least 95% identical to a sequence of at least 50
30 contiguous nucleotides in a sequence selected from said group.

Also preferred is a method for diagnosing in a subject a pathological condition associated with abnormal structure or expression of a gene encoding a protein identified in Table 1, which method comprises a step of detecting in a biological sample obtained from said subject nucleic acid molecules, if any, comprising a
35 nucleotide sequence that is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a cDNA clone identified by a cDNA Clone

5 Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

The method for diagnosing a pathological condition can comprise a step of detecting nucleic acid molecules comprising a nucleotide sequence in a panel of at least two nucleotide sequences, wherein at least one sequence in said panel is at least
10 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from said group.

Also preferred is a composition of matter comprising isolated nucleic acid molecules wherein the nucleotide sequences of said nucleic acid molecules comprise a panel of at least two nucleotide sequences, wherein at least one sequence in said
15 panel is at least 95% identical to a sequence of at least 50 contiguous nucleotides in a sequence selected from the group consisting of: a nucleotide sequence of SEQ ID NO:X wherein X is any integer as defined in Table 1; and a nucleotide sequence encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone
20 in Table 1. The nucleic acid molecules can comprise DNA molecules or RNA molecules.

Also preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1.

25 Also preferred is a polypeptide, wherein said sequence of contiguous amino acids is included in the amino acid sequence of SEQ ID NO:Y in the range of positions beginning with the residue at about the position of the "Total AA of the Open Reading Frame (ORF)" as set forth for SEQ ID NO:Y in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence
30 at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of SEQ ID NO:Y.

35 Further preferred is an isolated polypeptide comprising an amino acid sequence at least 95% identical to the complete amino acid sequence of SEQ ID NO:Y.

5 Further preferred is an isolated polypeptide comprising an amino acid sequence at least 90% identical to a sequence of at least about 10 contiguous amino acids in the complete amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

10 Also preferred is a polypeptide wherein said sequence of contiguous amino acids is included in the amino acid sequence of a portion of the protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence
15 at least 95% identical to a sequence of at least about 30 contiguous amino acids in the amino acid sequence of the portion of the protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence
20 at least 95% identical to a sequence of at least about 100 contiguous amino acids in the amino acid sequence of the protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated polypeptide comprising an amino acid sequence
25 at least 95% identical to the amino acid sequence of the protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Further preferred is an isolated antibody which binds specifically to a polypeptide comprising an amino acid sequence that is at least 90% identical to a
30 sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

35 Further preferred is a method for detecting in a biological sample a polypeptide comprising an amino acid sequence which is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as

5 defined in Table 1; and a complete amino acid sequence of a protein encoded by a
cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the
deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1;
which method comprises a step of comparing an amino acid sequence of at least one
polypeptide molecule in said sample with a sequence selected from said group and
10 determining whether the sequence of said polypeptide molecule in said sample is at
least 90% identical to said sequence of at least 10 contiguous amino acids.

Also preferred is the above method wherein said step of comparing an amino
acid sequence of at least one polypeptide molecule in said sample with a sequence
selected from said group comprises determining the extent of specific binding of
15 polypeptides in said sample to an antibody which binds specifically to a polypeptide
comprising an amino acid sequence that is at least 90% identical to a sequence of at
least 10 contiguous amino acids in a sequence selected from the group consisting of:
an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in
Table 1; and a complete amino acid sequence of a protein encoded by a cDNA clone
20 identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the
ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is the above method wherein said step of comparing sequences
is performed by comparing the amino acid sequence determined from a polypeptide
molecule in said sample with said sequence selected from said group.

25 Also preferred is a method for identifying the species, tissue or cell type of a
biological sample which method comprises a step of detecting polypeptide molecules
in said sample, if any, comprising an amino acid sequence that is at least 90%
identical to a sequence of at least 10 contiguous amino acids in a sequence selected
from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is
30 any integer as defined in Table 1; and a complete amino acid sequence of a protein
encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and
contained in the deposit with the ATCC Deposit Number shown for said cDNA clone
in Table 1.

Also preferred is the above method for identifying the species, tissue or cell
35 type of a biological sample, which method comprises a step of detecting polypeptide
molecules comprising an amino acid sequence in a panel of at least two amino acid
sequences, wherein at least one sequence in said panel is at least 90% identical to a

5 sequence of at least 10 contiguous amino acids in a sequence selected from the above group.

Also preferred is a method for diagnosing a pathological condition associated with an organism with abnormal structure or expression of a gene encoding a protein identified in Table 1, which method comprises a step of detecting in a biological
10 sample obtained from said subject polypeptide molecules comprising an amino acid sequence in a panel of at least two amino acid sequences, wherein at least one sequence in said panel is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a
15 complete amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

In any of these methods, the step of detecting said polypeptide molecules includes using an antibody.

20 Also preferred is an isolated nucleic acid molecule comprising a nucleotide sequence which is at least 95% identical to a nucleotide sequence encoding a polypeptide wherein said polypeptide comprises an amino acid sequence that is at least 90% identical to a sequence of at least 10 contiguous amino acids in a sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y
25 wherein Y is any integer as defined in Table 1; and a complete amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

Also preferred is an isolated nucleic acid molecule, wherein said nucleotide
30 sequence encoding a polypeptide has been optimized for expression of said polypeptide in a prokaryotic host.

Also preferred is an isolated nucleic acid molecule, wherein said polypeptide comprises an amino acid sequence selected from the group consisting of: an amino acid sequence of SEQ ID NO:Y wherein Y is any integer as defined in Table 1; and a
35 complete amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1.

5 Further preferred is a method of making a recombinant vector comprising inserting any of the above isolated nucleic acid molecule(s) into a vector. Also preferred is the recombinant vector produced by this method. Also preferred is a method of making a recombinant host cell comprising introducing the vector into a host cell, as well as the recombinant host cell produced by this method.

10 Also preferred is a method of making an isolated polypeptide comprising culturing this recombinant host cell under conditions such that said polypeptide is expressed and recovering said polypeptide. Also preferred is this method of making an isolated polypeptide, wherein said recombinant host cell is a eukaryotic cell and said polypeptide is a protein comprising an amino acid sequence selected from the
15 group consisting of: an amino acid sequence of SEQ ID NO:Y comprising the range depicted in "Total AA of ORF" of SEQ ID NO:Y wherein Y is an integer set forth in Table 1; and an amino acid sequence of a protein encoded by a cDNA clone identified by a cDNA Clone Identifier in Table 1 and contained in the deposit with the ATCC Deposit Number shown for said cDNA clone in Table 1. The isolated polypeptide
20 produced by this method is also preferred.

Also preferred is a method of treatment of an individual in need of an increased level of protein activity, which method comprises administering to such an individual a pharmaceutical composition comprising an amount of an isolated polypeptide, polynucleotide, or antibody of the claimed invention effective to increase
25 the level of said protein activity in said individual.

The above-recited applications have uses in a wide variety of hosts. Such hosts include, but are not limited to, barley, oats, rye, sorghum, pea, sunflower, tobacco, cotton, petunia, tomato, broccoli, lettuce, apple, plum, orange, and lemon, and more preferably rice, maize, conola, wheat, sugarbeet, sugarcane, and soybean,
30 in addition to other hosts referenced elsewhere herein.

Having generally described the invention, the same will be more readily understood by reference to the following examples, which are provided by way of illustration and are not intended as limiting.

35 **Examples**

Description of the Preferred Embodiments

5 Example 1: Culturing *Physcomitrella patens*

Plants of the species *Physcomitrella patens* (Hedw.) B.S.G. from the collection of the genetic studies section of the University of Hamburg were used. They originate from the strain 16/14 collected by H.L.K. Whitehouse in Gransden Wood, Huntingdonshire (England), which was subcultured from a spore by Engel (1968, Am
10 J Bot 55, 438-446). Proliferation of the plants was carried out by means of spores and by means of regeneration of the gametophytes. The protonema developed from the haploid spore as a chloroplast-rich chloronema and chloroplast-low caulonema, on which buds formed after approximately 12 days. These grew to give gametophores bearing antheridia and archegonia. After fertilization, the diploid sporophyte with a
15 short seta and the spore capsule resulted, in which the meiospores mature.

Culturing was carried out in a climatic chamber at an air temperature of 25°C and light intensity of 55 $\mu\text{mol s}^{-1}\text{m}^{-2}$ (white light; Philips TL 65W/25 fluorescent tube) and a light/dark change of 16/8 hours. The moss was either modified in liquid culture using Knop medium according to Reski and Abel (1985, Planta 165, 354-358) or
20 cultured on Knop solid medium using 1% oxoid agar (Unipath, Basingstoke, England).

The protonemas used for RNA and DNA isolation were cultured in aerated liquid cultures. The protonemas were comminuted every 9 days and transferred to fresh culture medium.

25

Example 2: Construction of the cDNA library from *Physcomitrella patens*

For cDNA library construction first strand synthesis was achieved using Murine Leukemia Virus reverse transcriptase (Roche, Mannheim, Germany) and oligo-d(T)-primers, second strand synthesis by incubation with DNA polymerase I,
30 Klenow enzyme and RNaseH digestion at 12 degree C (2h), 16 degree C (1h) and 22 degree C (1h). The reaction was stopped by incubation at 65 degree C (10 min) and subsequently transferred to ice. Double stranded DNA molecules were blunted by T4-DNA-polymerase (Roche, Mannheim) at 37 degree C (30 min). Nucleotides were removed by phenol/chloroform extraction and Sephadex G50 spin columns. EcoRI
35 adapters (Pharmacia, Freiburg, Germany) were ligated to the cDNA ends by T4-DNA-ligase (Roche, 12 degree C, overnight) and phosphorylated by incubation with polynucleotide kinase (Roche, 37 degree C, 30 min). This mixture was subjected to separation on a low melting agarose gel. DNA molecules larger than 300 basepairs

5 were eluted from the gel, phenol extracted, concentrated on Elutip-D-columns (Schleicher and Schuell, Dassel, Germany) and were ligated to vector arms and packed into lambda ZAPII phages or lambda ZAP-Express phages using the Gigapack Gold Kit (Stratagene, Amsterdam, Netherlands) using material and following the instructions of the manufacturer.

10

Example 3 : Sequencing and function annotation of *Physcomitrella patens* ESTs

cDNA libraries as described in Example 4 were used for DNA sequencing according to standard methods, in particular by the chain termination method using the ABI PRISM Big Dye Terminator Cycle Sequencing Ready Reaction Kit (Perkin-
15 Elmer, Weiterstadt, Germany). Random Sequencing was carried out subsequent to preparative plasmid recovery from cDNA libraries via in vivo mass excision, retransformation, and subsequent plating of DH10B on agar plates (material and protocol details from Stratagene, Amsterdam, Netherlands. Plasmid DNA was prepared from overnight grown *Escherichia coli* cultures grown in Luria-Broth (LB)
20 medium containing ampicillin (see Sambrook et al. (1989) (Cold Spring Harbor Laboratory Press: ISBN 0-87969-309-6)) on a Qiagen DNA preparation robot (Qiagen, Hilden) according to the manufacturers protocols. Sequencing primers with the following nucleotide sequences were used:

Qiagen1: 5'-CAGGAAACAGCTATGACC-3' (SEQ ID NO:6)

25 Qiagen2: 5'-CTAAAGGGAACAAAAGCTG-3' (SEQ ID NO:7)

Qiagen3: 5'-TGTAACACGACGGCCAGT-3' (SEQ ID NO:8)

Sequences were processed and annotated using the software package EST-MAX commercially provided by Bio-Max (Munich, Germany). The program incorporates practically all bioinformatics methods important for functional and
30 structural characterization of protein sequences. For reference see <http://pedant.mips.biochem.mpg.de>.

The most important algorithms incorporated in EST-MAX are:

FASTA: Very sensitive protein sequence database searches with estimates of statistical significance; Pearson W.R. (1990) Rapid and sensitive sequence
35 comparison with FASTP and FASTA. *Methods Enzymol.* 183:63-98.

BLAST: Very sensitive protein sequence database searches with estimates of statistical significance. Altschul S.F., Gish W., Miller W., Myers E.W., and Lipman D.J. Basic local alignment search tool. *Journal of Molecular Biology* 215:403-10 .

- 5 PREDATOR: High-accuracy secondary structure prediction from single and multiple sequences. Frishman, D. and Argos, P. (1997) 75% accuracy in protein secondary structure prediction. *Proteins*, 27:329-335.
- CLUSTALW: Multiple sequence alignment. Thompson, J.D., Higgins, D.G. and Gibson, T.J. (1994) CLUSTAL W: improving the sensitivity of progressive multiple
10 sequence alignment through sequence weighting, positions-specific gap penalties and weight matrix choice. *Nucleic Acids Research*, 22:4673-4680.
- TMAP: Transmembrane region prediction from multiply aligned sequences. Persson, B. and Argos, P. (1994) Prediction of transmembrane segments in proteins utilising multiple sequence alignments. *J. Mol. Biol.* 237:182-192.
- 15 ALOM2: Transmembrane region prediction from single sequences. Klein, P., Kanehisa, M., and DeLisi, C. Prediction of protein function from sequence properties: A discriminant analysis of a database. *Biochim. Biophys. Acta* 787:221-226 (1984).
Version 2 by Dr. K. Nakai.
- PROSEARCH: Detection of PROSITE protein sequence patterns. Kolakowski L.F.
20 Jr., Leunissen J.A.M., Smith J.E. (1992) ProSearch: fast searching of protein sequences with regular expression patterns related to protein structure and function. *Biotechniques* 13, 919-921.
- BLIMPS: Similarity searches against a database of ungapped blocks. J.C. Wallace and Henikoff S., (1992)
- 25 PATMAT: A searching and extraction program for sequence, pattern and block queries and databases, *CABIOS* 8:249-254. Written by Bill Alford.

Example 4: Binary vector construction

The pNC53 vector was digested with HindIII (Roche) and blunt-end filled
30 with Klenow enzyme and 0.1mM dNTPs (Roche) according to manufacture's instructions. This fragment was extracted from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) according to manufacture's instructions. The purified fragment was then digested with EcoRI (Roche) according to manufacture's instructions. This fragment was extracted from agarose gel with a QIAquick Gel
35 Extraction Kit (Qiagen) according to manufacture's instructions. The resulting 1.4kilobase fragment, the gentamycin cassette, included the nos promoter, aacCI gene and the g7 terminator.

5 The vector pBlueScript was digested with EcoRI and SmaI (Roche) according to manufacture's instructions. The resulting fragment was extracted from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) according to manufacture's instructions. The digested pBlueScript vector and the gentamycin cassette fragments were ligated with T4 DNA Ligase (Roche) according to manufacture's instructions, joining the two
10 respective EcoRI sites and joining the blunt-ended HindIII site with the SmaI site. The recombinant vector (pGMBS) was transformed into Top10 cells (Invitrogen) using standard conditions. Transformed cells were selected for on LB agar containing 50 µg/ml Kanamycin, grown overnight at 37°C and used to inoculate 3ml of liquid LB containing 50 µg/ml Kanamycin and grown overnight at 37°C. Plasmid DNA was
15 extracted using the QIAprep Spin Miniprep Kit (Qiagen) following manufacture's instructions. Analyses of subsequent clones and restriction mapping was performed according to standard molecular biology techniques (Sambrook et al. 1989).

 Both the pGMBS vector and p1bxSuperGUS vector were digested with XbaI and KpnI (Roche) according to manufacture's instructions, excising the gentamycin
20 casstte from pGMBS and producing the backbone from the p1bxSuperGUS vector. The resulting fragments were extracted from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) according to manufacture's instructions. These two fragments were ligated with T4 DNA ligase (Roche) according to manufactures's instructions. The resulting recombinant vector (pGMSG) was transformed into Top10 cells
25 (Invitrogen) using standard conditions. Transformed cells were selected for on LB agar containing 50 µg/ml Kanamycin, grown overnight at 37°C and used to inoculate 3ml of liquid LB containing 50 µg/ml Kanamycin and grown overnight at 37°C. Plasmid DNA was extracted using the QIAprep Spin Miniprep Kit (Qiagen) following manufacture's instructions. Analyses of subsequent clones and restriction mapping was
30 performed according to standard molecular biology techniques (Sambrook et al. 1989).

Example 5: Total RNA isolation from *Physcomitrella patens*

 The total RNA was obtained from wild-type 14 day old *Physcomitrella patens*
35 following the Van Slogteren method (Van Slogteren, C.M.S., Hoge, J.H.C., Hooykaas, P.J.J., and Schilperoort, R.A.1983. Clonal analysis of heterogenous crown gall tumor tissues induced by wild-types and shooter mutant strains of *Agrobacterium*

5 *tumefaciens*-expression of T-DNA genes. Plant Mol. Biol. 2: 321-333.1983) with slight modifications. The plant tissue (200mg) was frozen with liquid nitrogen and ground to a fine powder with a mortar and pestle. The powder was placed in a microfuge tube and the RNA was extracted with 500ul of extraction buffer (phenyl:0.1M LiCl, 100mM Tris-HCl [pH8.0], 10mM EDTA, 1% SDS (w/v) [1:1]
10 preheated to 90oC. The mixture was heated further for 1 min at 90oC and then vortexed for 5 mins. Proteins were extracted by adding 250 ul of chloroform:isoamyl alcohol (24:1) and the mixture was vortexed for 5 mins and centrifuged for 15 mins at 13,000 rpm in an Eppendorf centrifuge 5414 at 4oC. The aqueous layer was removed and the protein extraction was repeated twice more. One vol of 4mM LiCl was added
15 and the RNA was allowed to precipitate overnight at 4oC. To collect the RNA, the mixture was centrifuged for 15 min at 4oC at 13,000 rpm in an Eppendorf centrifuge 5414. The pellet was resuspended in 250 ul sterile, deionized water. To precipitate the RNA, 0.1 vols of 3M sodium acetate (pH5.2) and 2 vols 100% ethanol were added. An aliquote was taken and centrifuged for 20 mins at 4oC at 13,000 rpm in an
20 Eppendorf centrifuge 5414. The pellet was washed with 70% ethanol to remove salts from the pellet and dried using a speed vac. The pellet was resuspended in 25 ul DEPC H2O and analyzed for integrity via electrophoresis. The RNA was stored at –70° degree C.

25 **Example 6: Construction of cDNA library of *Physcomitrella patens***

To isolate the full-length clones encoding Na⁺/H⁺ Antiporter from *Physcomitrella patens* a cDNA library was created with SMART RACE cDNA Amplification kit (Clontech Laboratories) following manufacture's instructions. Total RNA created as described in Example 5 was used as the template.

30

Example 7: Obtaining the full length clone of the vacuole-type Na⁺/H⁺ antiporter from *Physcomitrella patens*

The Physcomitrella patens full-length and partial clones (EST#156 and EST#610) for
35 PpNHX1 (SEQ ID NO: 1) and PpNHX2 (SEQ ID NO: 3) were identified in the *Physcomitrella patens* EST sequencing program using the program EST-MAX through BLAST analysis. These particular clone were chosen for further analyses since they encoded for Na⁺/H⁺ antiporter.

5

For PpNHX1A full-length sequence corresponding to EST #156 was obtained by performing polymerase chain reaction (PCR) with a gene-specific EST as the template DNA. The synthetic oligonucleotide primers (MWG-Biotech) for the reaction were: 156F: 5'-CTGCCCGGGTACCGGAAGGTTGAAGATGGCGAC-3'

10 (SEQ ID NO:9) and 156R: 5'-GACGAGCTCTAGACACTAAGTTGATTCAACTGTG-3' (SEQ ID NO:10). The primers designed contained a SmaI and a KpnI site in the 5' region and a SacI and a XbaI site in the 3' region for cloning purposes. The conditions for the reaction were standard conditions with PWO DNA polymerase (Roche). PCR was performed

15 according to standard conditions and to manufacture's protocols (Sambrook et al. 1989, Biometra T3 Thermocycler). The parameters for the reaction were: five minutes at 94°C followed by five cycles of one minute at 94°C, one minute at 50°C and 1.5 minutes at 72°C. This was followed by twentyfive cycles of one minute at 94°C, one minute at 65°C and 1.5 minutes at 72°C. These parameters generated a fragment 1.7

20 kilobases long.

The fragment was extracted from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) and ligated into the TOPO pCR 2.1 vector (Invitrogen) following manufacture's instructions. Recombinant vectors were transformed into Top10 cells (Invitrogen) using standard conditions (Sambrook et al. 1989). Transformed cells

25 were selected for on LB agar containing 100 µg/ml carbenicillin, 0.8mg X-gal (5-bromo-4-chloro-3-indolyl-β-D-galactoside) and 0.8mg IPTG (isopropylthio-β-D-galactoside) grown overnight at 37°C. White colonies were selected and used to inoculate 3ml of liquid LB containing 100 µg/ml ampicillin and grown overnight at 37°C. Plasmid DNA was extracted using the QIAprep Spin Miniprep Kit (Qiagen)

30 following manufacture's instructions. Analyses of subsequent clones and restriction mapping was performed according to standard molecular biology techniques (Sambrook et al. 1989).

For PpNHX2

35 To isolate full-length clone encoding for PpNHX2 from *Physcomitrella patens*, cDNA libraries was used as a template for RACE.

- 5' and 3' RACE Protocol

5

The EST sequence (EST#610) identified from the database search as described in Example 3 was used to design oligos (CGTTTGCGCATGCCCC TGGTCGCTT SEQ ID NO: 15; CAGGACGAAGGAAACACCGACACCT SEQ ID NO: 16) for RACE. The extended sequences for the gene were obtained by performing Rapid Amplification of cDNA Ends polymerase chain reaction (RACE PCR) using the Advantage 2 PCR kit (Clontech Laboratories) and the SMART RACE cDNA amplification kit (Clontech Laboratories) using a Biometra T3 Thermocycler following the manufacturer's instructions.

15 **The sequences obtained from the RACE reactions corresponded to full-length coding regions of PpNHX2 and was used to design oligos for full-length cloning of the respective genes (see below full-length amplification).**

- Full-length Amplification

20

Full-length clones corresponding PpNHX2 (SEQ ID NO: 4) was obtained by performing polymerase chain reaction (PCR) with gene-specific primers (610F: CTGCCCGGGTACCATTGGCACCAGCAAGATGGCGGACGCT SEQ ID NO: 17 610R: GCGTCTAGAC CCGGGTCATA TACACGCGGT GGTCAGGAG SEQ ID NO: 18).

25 The fragment of the full-length clone was isolated and sub-cloned into the TOPO pCR 2.1 vector (Invitrogen) by following the same procedure for PpNHX1 except for following several steps. For cloning purposes the primers designed contained a SmaI and a KpnI site in the 5' region and a SmaI and a XbaI site in the 3' region. The parameters for the PCR reaction were the same as that for PpNHX1 except for the extension time for 3 minute at 72 °C. This extension time allowed to amplify a fragment 2.6 kb.

35 **Example 8: Subcloning the Na⁺/H⁺ antiporter from *Physcomitrella patens* into pGMSG.**

For PpNHX1

5 The fragment containing the Na⁺/H⁺ antiporter was excised from the recombinant TOPO pCR vector by digestion with SacI and SmaI (Roche) according to manufacture's instructions. The subsequent fragment was excised from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) according to manufacture's instructions. The target binary vector was also cleaved with SacI and SmaI (Roche) and
10 dephosphorylated with shrimp alkaline phosphatase (Roche) according to manufacture's instructions. The resulting fragment was extracted from agarose gel with a QIAquick Gel Extraction Kit (Qiagen). The binary vector and Na⁺/H⁺ antiporter fragments were ligated together with Rapid DNA Ligation Kit (Roche), joining the respective SacI sites and SmaI sites.

15 Recombinant vectors were transformed into Top10 cells (Invitrogen) using standard conditions (Sambrook et al. 1989). Transformed cells were selected for on LB agar containing 50 µg/ml Kanamycin (Roche). Plasmid DNA was extracted using the QIAprep Spin Miniprep Kit (Qiagen) following manufacture's instructions. Analyses of subsequent clones and restriction mapping was performed according to standard
20 molecular biology techniques (Sambrook et al. 1989). The recombinant vector was transformed into Agrobacterium tumefaciens according to standard conditions (Hoefgen and Willmitzer, 1990 Biochemical and genetic analysis of different patatin isoforms expressed in various organs of potato (*solanum tuberosum*); Plant Science 66, 221-230). The resultant vector is named as pBPSNVT001.

25 **For PpNHX2**

The full-length fragment of the Na⁺/H⁺ antiporter was subcloned into the binary vector by following the same procedure for PpNHX1 except for the following steps. For the excision of the fragment from the TOPO pCR vector KpnI and SmaI (Roche) were used. The resultant fragment was blunt ended and ligated into the binary vector
30 that was digested by SmaI (Roche) and dephosphorylated with shrimp alkaline phosphatase (Roche) according to manufacture's instructions. The resultant vector is named as pBPSNVT003.

Example 9: Transformation of Agrobacterium tumefaciens

35 The recombinant vectors were transformed into Agrobacterium tumefaciens C58C1 and PMP90 according to standard conditions (Hoefgen and Willmitzer, 1990).

5 **Example 10: Subcloning the Na⁺/H⁺ antiporter from *Physcomitrella patens* into pYES.**

The fragments containing the Na⁺/H⁺ antiporter (PpNHX1 and PpNHX2) was excised respectively from the recombinant TOPO pCR vector by digestion with KpnI and XbaI (Roche) according to manufacture's instructions. The subsequent fragment
10 was excised from agarose gel with a QIAquick Gel Extraction Kit (Qiagen) according to manufacture's instructions. The target binary vector was also cleaved with KpnI and XbaI (Roche) and dephosphorylated with shrimp alkaline phosphatase (Roche) according to manufacture's instructions. The vector and Na⁺/H⁺ antiporter fragments were ligated together with Rapid DNA Ligation Kit (Roche) and transformed into
15 Top10 Cells (Invitrogen) according to standard protocols (Sambrook et al. 1989). Transformed cells were selected for on LB agar containing 50 µg/ml Kanamycin. Plasmid DNA was extracted using the QIAprep Spin Miniprep Kit (Qiagen) following manufacture's instructions. Analyses of subsequent clones and restriction mapping was performed according to standard molecular biology techniques (Sambrook et al.
20 1989).

Example 11: *In vivo* complementation of the *NHX1 Saccharomyces cerevisiae* mutant

The recombinant expression pYES2 vector (Invitrogen) containing the Na⁺/H⁺
25 antiporters, PpNHX1 and PpNHX2, respectively in the sense orientation under the *Saccharomyces cerevisiae* GAL1 promoter was transformed into the yeast mutant BY4741 (MATa his3::D1 leu2::D0 lys2D0 ura3::D0) following manufacture's instructions. The transformed cells were selected for on Complete Supplement Mixture (CSM) minus Uracil 0.8% agar (Bio 101, Inc.) grown at 30°C for two days.
30 The transformed colonies were selected to plate on 1) CSM plus Uracil 0.8% agar plates, 2) CSM minus Uracil 0.8% agar plates, 3) CSM minus Uracil 0.8% agar plates supplemented with 2% galactose for induction of the expression of the antiporter and 4) CSM minus Uracil 0.8% agar plates supplemented with 2% galactose for induction of the expression of the antiporter and 0.5M NaCl. The plates were incubated at 30°C
35 for two days.

5 **Example 12: Transformation of *Arabidopsis thaliana***

Arabidopsis thaliana ecotype C24 were grown and transformed according to standard conditions (Bechtold 1993 (Bechtold, N., Ellis, J., Pelletier, G. 1993. *In planta Agrobacterium* mediated gene transfer by infiltration of *Arabidopsis thaliana* plants C.R. Acad.Sci.Paris. 316:1194-1199); Bent et al. 1994 (Bent, A., Kunkel, B.N., Dahlbeck, D., Brown, K.L., Schmidt, R., Giraudat, J., Leung, J., and Staskawicz, B.J. 1994; PPCS2 of *Arabidopsis thaliana*: A leucine-rich repeat class of plant disease resistant genes; Science 265:1856-1860).

Example 13: Screening of transformed *Arabidopsis* plants

15 *Arabidopsis* T1 and T2 seeds were sterilized according to standard protocols (Xiong et al., Plant Molecular Biology 17, 150-170 (1999)). Seeds were plated on ½ MS 0.6% agar supplemented with 1 - 3% sucrose and 50 -150 µg/ml gentamycin. Seeds on plates were vernalized for two days at 4°C. The seeds were germinated in a climatic chamber at an air temperature of 22°C and light intensity of 55 µmol s⁻¹m⁻² (white light; Philips TL 65W/25 fluorescent tube) and a light/dark change of 16/8 hours. Transformed *Arabidopsis* seedlings were selected after 7 - 14 and transferred to ½ MS 0.6% agar plates supplemented with 1 % sucrose and allowed to recover for 1 - 5 days. The transgenic plants are screened for their improved salt tolerance according to the screening method described in example 17 demonstrating that transgene expression confers salt tolerance.

Example 14: Detection of the Na⁺/H⁺ antiporter Transgene in the *Arabidopsis* Transgenic Lines

30 To check for the presence of the PpNHX1 and PpNHX2 transgene in *Arabidopsis* T1 and T2 transgenic lines, PCR was performed on genomic DNA which contaminates the RNA samples taken as described in Example 5 below. 2.5 µl of RNA sample was used in a 50 µl PCR reaction using *Taq* DNA polymerase (Roche Molecular Biochemicals) according to the manufacturer's instructions. The gene specific primers (for PpNHX1:156F: 5'-CTGCCCGGGTACCGGAAGGTTGAAGATGGCGAC-30'
35 SEQ ID NO: 19 and 156R: 5'-GACGAGCTCTAGACACTAAGTTGATTCAACTGTG-3. SEQ ID NO: 20; for

5 PpNHX2: 610F: CTGCCCCGGGTACCATTGGCACCAGCAAGATGGCGGACGCT
SEQ ID NO: 21 610R: GCGTCTAGAC CCGGGTCATA TACACGCGGT
GGTCAGGAG SEQ ID NO: 22) were used for the PCR. Binary vector plasmid with
PpNHX1 and PpNHX2 gene cloned in was used as positive control, and the wild type
C24 genomic DNA was used as negative control in the PCR reactions. 10 µl PCR
10 reaction was analyzed on 0.8% agarose - ethidium bromide gel.

The transgene was successfully amplified from the T1 and T2 transgenic lines,
but not from the wild-type plants. This result indicated that the T1 and T2 transgenic
plants contain at least one copy of the PpNHX1 and PpNHX2 transgene. There was
no indication of existence of the corresponding gene or homolog of the PpNHX1
15 and PpNHX2 that could be amplified in this method in the wild-type plants.

Example 15: Detection of the Na⁺/H⁺ antiporter Transgene mRNA in Transgenic Arabidopsis Lines

Transgene expression was detected using RT-PCR. Total RNA was isolated
20 from T1 and T2 plants using a procedure adapted from (Verwoerd et al. 1989. NAR
17:2362). Leaf samples (50-100mg) were collected and ground to a fine powder in
liquid nitrogen. Ground tissue was resuspended in 500µl of a 80°C, 1:1 mixture, of
phenol to extraction buffer (100mM LiCl, 100mM Tris pH8, 10mM EDTA, 1% SDS),
followed by brief vortexing to mix. After the addition of 250 µl of chloroform each
25 sample was vortexed briefly. Samples were then centrifuged for 5 minutes at 12,000
x g. The upper aqueous phase was removed to a fresh eppendorf tube. RNA was
precipitated by adding 1/10th volume 3M sodium acetate and 2 volumes 95% ethanol.
Samples were mixed by inversion and placed on ice for 30 minutes. RNA was
pelleted by centrifugation at 12,000 x g for 10 minutes. The supernatant was removed
30 and pellets briefly air dried. RNA sample pellets were resuspended in 10 µl DEPC
treated water. To remove contaminating DNA from the samples, each was treated
with RNase-free DNase (Roche) according to the manufacturer's recommendations.
cDNA was synthesized from total RNA using the Superscript First-Strand Synthesis
System for RT-PCR (Gibco-BRL) following manufacturer's recommendations. PCR
35 amplification of a gene-specific fragment from the synthesized cDNA was performed
using *Taq* DNA polymerase (Roche) and gene-specific primers (for PpNHX1:156F:
5'-CTGCCCCGGGTACCGGAAGGTTGAAGATGGCGAC-3' SEQ ID NO: 19 and

5 156R: 5'-GACGAGCTCTAGACACTAACTTGATTCAACTGTG-3. SEQ ID NO:
20 ; for PpNHX2: 5'-CGTTTGCGCATGCCCTGGTCGCTT-3' SEQ ID NO: 21,
610R: GCGTCTAGAC CCGGGTCATA TACACGCGGT GGTCAGGAG SEQ ID
NO. 22) in the following reaction: 1X PCR buffer, 1.5mM MgCl₂, 0.2 μM each
primer, 0.2μM dNTPs, 1 unit polymerase, 5μl cDNA from synthesis reaction.
10 Amplification was performed under the following conditions: Denaturation, 95°C, 1
minute; annealing, 62°C, 30 seconds; extension, 72°C, 1 minute, 35 cycles; extension,
72°C, 5 minutes; hold, 4°C, forever. PCR products were run on a 1% agarose gel,
stained with ethidium bromide, and visualized under UV light using the Quantity-One
gel documentation system (Bio-Rad).

15

Expression of the transgenes was detected in the T1 and T2 transgenic lines. These
results indicated that the transgenes are expressed in the transgenic lines and strongly
suggested that their gene product improved plant stress tolerance in the transgenic
lines. In agreement with the previous statement, no expression of identical or very
20 similar endogenous genes could be detected by this method. These results are in
agreement with the data from Example 14.

Example 16: Transformation of soybean

Seeds of soybean are surface sterilized with 70 % ethanol for 4 minutes at
25 room temperature with continuous shaking, followed by 20% (v/v) Clorox
supplemented with 0.05 % (v/v) Tween for 20 minutes with continuous shaking.
Then, the seeds are rinsed 4 times with distilled water and placed on moistened sterile
filter paper in a Petri dish at room temperature for 6 to 39 hours. The seed coats are
peeled off, and cotyledons are detached from the embryo axis. The embryo axis is
30 examined to make sure that the meristematic region is not damaged. The excised
embryo axes are collected in a half-open sterile Petri dish and air dried to a moisture
content less than 20% (fresh weight) in a sealed Petri dish until further use.

Agrobacterium tumefaciens culture is prepared from a single colony in LB solid
medium plus appropriate antibiotics (e.g. 100 mg/l streptomycin, 50 mg/l kanamycin)
35 followed by growth of the single colony in liquid LB medium to an optical density at
600 nm of 0.8. Then, the bacteria culture is pelleted at 7000 rpm for 7 minutes at RT,
and resuspended in MS (Murashige and Skoog, 1962) medium supplemented with

5 100 μ M acetosyringone. Bacteria cultures are incubated in this pre-induction medium for 2 hours at RT before use. The axis of soybean zygotic seed embryos at approximately 15% moisture content are imbibed for 2 hours at RT with the pre-induced *Agrobacterium* suspension culture. The embryos are removed from the imbibition culture and are transferred to Petri dishes containing solid MS medium
10 supplemented with 2% sucrose and incubated for 2 days, in the dark at RT. Alternatively, the embryos are placed on top of moistened (liquid MS medium) sterile filter paper in a Petri dish and incubated under the same conditions described above. After this period, the embryos are transferred to either solid or liquid MS medium supplemented with 500 mg/L carbenicillin or 300mg/L cefotaxime to kill the
15 agrobacteria. The liquid medium is used to moisten the sterile filter paper. The embryos are incubated during 4 weeks at 25 °C, under 150 $\mu\text{mol m}^{-2}\text{sec}^{-1}$ and 12 hours photoperiod. Once the seedlings have produced roots, they are transferred to sterile metromix soil. The medium of the *in vitro* plants is washed off before transferring the plants to soil. The plants are kept under a plastic cover for 1 week to favor the
20 acclimatization process. Then the plants are transferred to a growth room where they are incubated at 25 °C, under 150 $\mu\text{mol m}^{-2}\text{sec}^{-1}$ light intensity and 12 hours photoperiod for about 80 days.

Example 17: Transformation of Canola

25 The method described in example 14 of plant transformation is also applicable to Brassica and other crops. Seeds of canola are surface sterilized with 70% ethanol for 4 minutes at room temperature with continuous shaking, followed by 20% (v/v) Clorox supplemented with 0.05 % (v/v) Tween for 20 minutes, at RT with continuous shaking. Then, the seeds are rinsed 4 times with distilled water and placed on
30 moistened sterile filter paper in a Petri dish at room temperature for 18 hours. Then the seed coats are removed and the seeds are air dried overnight in a half-open sterile Petri dish. During this period the seeds loses approx. 85% of its water content. The seeds are then stored at room temperature in a sealed Petri dish until further use. DNA constructs and embryo imbibition are as described in example 14. Samples of the
35 primary transgenic plants (T0) are analyzed by PCR to confirm the presence of T-DNA. These results are confirmed by Southern hybridization in which DNA is electrophoresed on a 1% agarose gel and transferred to a positively charged nylon

5 membrane (Roche Diagnostics). The PCR DIG Probe Synthesis Kit (Roche Diagnostics) is used to prepare a digoxigenin-labelled probe by PCR, and used as recommended by the manufacturer.

Example 18: Transformation of Corn

10 Transformation of maize (*Zea Mays L.*) is performed with the method described by Ishida et al. 1996. Nature Biotch 14745-50), Immature embryos are cocultivated with *Agrobacterium tumefaciens* that carry “super binary” vectors, and transgenic plants are recovered through organogenesis. This procedure provides a transformation efficiency between 2.5% and 20%.

15 The imbibition of dry embryos with a culture of *Agrobacterium* is also applicable to maize embryo axes. The experimental protocol is the same as described in example 16 but using maize seeds as the source of embryos.

Example 19: Salt Tolerance Screening

Salt test on MS plate

20 Seedlings were transferred to filter paper soaked in ½ MS and placed on ½ MS 0.6% agar supplemented with 2ug/ml benomyl the night before the stress screening. For the stress screening, the filter paper with the seedlings was moved to stacks of sterile filter paper, soaked in 50mM NaCl, in a petri dish. After two hours, the filter paper
25 with the seedlings was moved to stacks of sterile filter paper, soaked with 200mM NaCl, in a petri dish. After two hours, the filter paper with the seedlings was moved to stacks of sterile filter paper, soaked in 600mM NaCl, in a petri dish. After 10 hours, the seedlings were moved to petri dishes containing ½ MS 0.6% agar supplemented with 2ug/ml benomyl. The seedlings were scored after 5 days.

30

Soil test for salt tolerance

Seeds of plants to be tested are sterilized (100% bleach, 0.1% TritonX for five minutes two times and rinsed five times with ddH₂O). Seeds are plated on non-selection media (1/2 MS, 0.6% phytagar, 0.5g/L MES, 1% sucrose, 2ug/ml benamyl).
35 Seeds are allowed to germinate for approximately ten days. At the 4-5 leaf stage, transgenic plants were potted into 5.5cm diameter pots filled with loosely packed soil (Metromix 360, Scotts) wetted with 1g/L 20-20-20 fertilizer (Peters Professional, Scotts).

5 The plants are allowed to grow (22°C, continuous light) for approx. seven days, watering as needed. When the plants are just about to bolt, the water is removed from the tray and the assay is started. To begin the assay, three liters of 100mM NaCl and 1/8 MS was added to the tray under the pots. To the tray containing the control plants, three liters of 1/8 MS was added. After 10 days, the NaCl treated and the control
10 plants were given water. Ten days later, the plants were photographed.

Figures 6 and 7 show pictures of an example of the T1 PpNHX1 and PpNHX2 transgenic lines and wild-type plants after the salt stress. As can be seen from the
15 Figures the transgenic lines are more tolerant than the wildtype after salt stress treatment. On the other hand, their growth of wild-type plants were severely inhibited under the same condition. This result clearly indicates that the transgenic lines over-expressing the PpNHX1 and PpNHX2 gene acquired salt stress tolerance. The results should be better when a homozygous strong expresser will be found.

20 **Example 20: Drought Tolerance Screening**

T1 and T2 seedlings were transferred to dry, sterile filter paper in a petri dish and allowed to desiccate for two hours at 80% RH (relative humidity) in a Sanyo Growth Cabinet MLR-350H, micromols⁻¹m² (white light; Philips TL 65W/25 fluorescent tube). The RH was then decreased to 60% and the seedlings were desiccated further
25 for eight hours. Seedlings were then removed and placed on ½ MS 0.6% agar plates supplemented with 2ug/ml benomyl and scored after five days.

The transgenic plants are screened for their improved drought tolerance demonstrating that transgene expression confers drought tolerance.

30

Example 21: Freezing Tolerance Screening

Seedlings were moved to petri dishes containing ½ MS 0.6% agar supplemented with 2% sucrose and 2ug/ml benomyl. After four days, the seedlings were incubated at +4°C for 1 hour and then covered with shaved ice. The seedlings
35 were then placed in an Environmental Specialist ES2000 Environmental Chamber and incubated for 3.5 hours beginning at -1.0°C decreasing -1°C hour. The seedlings were

5 then incubated at -5.0°C for 24 hours and then allowed to thaw at $+5^{\circ}\text{C}$ for 12 hours.
The water was poured off and the seedlings were scored after 5 days.

The transgenic plants are screened for their improved cold tolerance demonstrating that transgene expression confers cold tolerance.

10

Example 22: Engineering salt-tolerant Arabidopsis plants by overexpression of Na^+/H^+ antiporters

The construct pBPSNVT001 and pBPSNVT003 were used to transform Arabidopsis as described below.

15 Transgenic plants overexpressing the Na^+/H^+ antiporters were obtained and screened as described in the examples 12 and 13, respectively. The transgenic plants are screened for their improved salt tolerance according to the screening method described in example 19 demonstrating that transgene expression confers salt tolerance.

20

Example 23: Engineering salt-tolerant rapeseed plants by overexpression of Na^+/H^+ antiporters

The construct pBPSNVT001 and pBPSNVT003 were used to transform rapeseed as described below.

25 Transgenic plants overexpressing the Na^+/H^+ antiporters were obtained and screened as described in the example 17. The transgenic plants are screened for their improved salt tolerance according to the screening method described in example 19 demonstrating that transgene expression confers salt tolerance.

30 **Example 24: Engineering salt-tolerant soybean plants by overexpression of Na^+/H^+ antiporters**

The construct pBPSNVT001 and pBPSNVT003 were used to transform soybean as described below.

35 Transgenic plants overexpressing the Na^+/H^+ antiporters were obtained and screened as described in the example 16. The transgenic plants are screened for their improved salt tolerance according to the screening method described in example 19 demonstrating that transgene expression confers salt tolerance.

5 **Example 25: Engineering salt-tolerant corn plants by overexpression of Na⁺/H⁺ antiporters**

Transgenic plants overexpressing the Na⁺/H⁺ antiporters were obtained and screened as described in the example 18. The transgenic plants are screened for their improved salt tolerance according to the screening method described in example 19
10 demonstrating that transgene expression confers salt tolerance.

Example 26: Engineering stress-tolerant wheat plants plants by over-expressing the Na⁺/H⁺ antiporters

15 The construct pBPSNVT001 and pBPSNVT003 were used to transform wheat as described below.

Transformation of wheat is performed with the method described by Ishida et al. 1996. Nature Biotech 14745-50), Immature embryos are cocultivated with
20 *Agrobacterium tumefaciens* that carry “super binary” vectors, and transgenic plants are recovered through organogenesis. This procedure provides a transformation efficiency between 2.5% and 20%.

The transgenic plants are screened for their improved salt tolerance according to the
25 screening method described in Example 19 demonstrating that transgene expression confers stress tolerance.

Example 27 - Isolation of a Specific Clone from the Deposited Sample.

The deposited material in the sample assigned the ATCC Deposit Number
30 cited in Table 1 for any given cDNA clone also may contain one or more additional plasmids, each comprising a cDNA clone different from that given clone. Thus, deposits sharing the same ATCC Deposit Number contain at least a plasmid for each cDNA clone identified in Table 1. Typically, each ATCC deposit sample cited in Table 1 comprises a mixture of approximately equal amounts (by weight) of about 1-
35 10 plasmid DNAs, each containing a different cDNA clone and/or partial cDNA clone; but such a deposit sample may include plasmids for more or less than 2 cDNA clones.

5 Two approaches can be used to isolate a particular clone from the deposited sample of plasmid DNAs cited for that clone in Table 1. First, a plasmid is directly isolated by screening the clones using a polynucleotide probe corresponding to SEQ ID NO:X.

10 Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The oligonucleotide is labeled, for instance, with ^{32}P -(-ATP using T4 polynucleotide kinase and purified according to routine methods. (E.g., Maniatis et al., *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Press, Cold Spring, NY (1982).) The plasmid mixture is transformed into a suitable host, as indicated above (such as
15 XL-1 Blue (Stratagene)) using techniques known to those of skill in the art, such as those provided by the vector supplier or in related publications or patents cited above. The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for
20 bacterial colony screening (e.g., Sambrook et al., *Molecular Cloning: A Laboratory Manual*, 2nd Edit., (1989), Cold Spring Harbor Laboratory Press, pages 1.93 to 1.104), or other techniques known to those of skill in the art.

Alternatively, two primers of 17-20 nucleotides derived from both ends of the SEQ ID NO:X (i.e., within the region of SEQ ID NO:X bounded by the 5' NT and
25 the 3' NT of the clone defined in Table 1) are synthesized and used to amplify the desired cDNA using the deposited cDNA plasmid as a template. The polymerase chain reaction is carried out under routine conditions, for instance, in 25 ul of reaction mixture with 0.5 ug of the above cDNA template. A convenient reaction mixture is 1.5-5 mM MgCl_2 , 0.01% (w/v) gelatin, 20 uM each of dATP, dCTP, dGTP, dTTP, 25
30 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94 degree C for 1 min; annealing at 55 degree C for 1 min; elongation at 72 degree C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cyclor. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product
35 is verified to be the selected sequence by subcloning and sequencing the DNA product.

Several methods are available for the identification of the 5' or 3' non-coding and/or coding portions of a gene which may not be present in the deposited clone.

5 These methods include but are not limited to, filter probing, clone enrichment using specific probes, and protocols similar or identical to 5' and 3' "RACE" protocols that are well known in the art. For instance, a method similar to 5' RACE is available for generating the missing 5' end of a desired full-length transcript. (Fromont-Racine et al., *Nucleic Acids Res.* 21(7):1683-1684 (1993)).

10 Briefly, a specific RNA oligonucleotide is ligated to the 5' ends of a population of RNA presumably containing full-length gene RNA transcripts. A primer set containing a primer specific to the ligated RNA oligonucleotide and a primer specific to a known sequence of the gene of interest is used to PCR amplify the 5' portion of the desired full-length gene. This amplified product may then be
15 sequenced and used to generate the full-length gene.

This above method starts with total RNA isolated from the desired source, although poly-A+ RNA can be used. The RNA preparation can then be treated with phosphatase if necessary to eliminate 5' phosphate groups on degraded or damaged RNA that may interfere with the later RNA ligase step. The phosphatase should then
20 be inactivated and the RNA treated with tobacco acid pyrophosphatase in order to remove the cap structure present at the 5' ends of messenger RNAs. This reaction leaves a 5' phosphate group at the 5' end of the cap cleaved RNA which can then be ligated to an RNA oligonucleotide using T4 RNA ligase.

This modified RNA preparation is used as a template for first strand cDNA
25 synthesis using a gene specific oligonucleotide. The first strand synthesis reaction is used as a template for PCR amplification of the desired 5' end using a primer specific to the ligated RNA oligonucleotide and a primer specific to the known sequence of the gene of interest. The resultant product is then sequenced and analyzed to confirm that the 5' end sequence belongs to the desired gene. Moreover, it may be
30 advantageous to optimize the RACE protocol to increase the probability of isolating additional 5' or 3' coding or non-coding sequences. Various methods of optimizing a RACE protocol are known in the art, though a detailed description summarizing these methods can be found in B.C. Schaefer, *Anal. Biochem.*, 227:255-273, (1995).

35 **Example 28 - Tissue Distribution of Polypeptide.**

Tissue distribution of mRNA expression of polynucleotides of the present invention is determined using protocols for Northern blot analysis, described by, among others, Sambrook et al. For example, a cDNA probe produced by the method

5 described in Example 1 is labeled with p32 using the rediprimetm DNA labeling system (Amersham Life Scinece), according to manufacturer's instructions. After labeling, the probe is purified using CHROMA SPIN0-100 column (Clontech Laboratories, Inc.) according to manufacturer's protocol number PT1200-1. The purified labeled probe is then used to examine various tissues for mRNA expression.

10 Tissue Northern blots containing the bound mRNA of various tissues are examined with the labeled probe using ExpressHybtm hybridization solution (Clonetech according to manufacturers protocol number PT1190-1. Northern blots can be produced using various protocols well known in the art (e.g., Sambrook et al). Following hybridization and washing, the blots are mounted and exposed to film at –
15 70C overnight, and the films developed according to standard procedures.

Example 29 Chromosomal Mapping of the Polynucleotides.

An oligonucleotide primer set is designed according to the sequence at the 5' end of SEQ ID NO:X. This primer preferably spans about 100 nucleotides. This
20 primer set is then used in a polymerase chain reaction under the following set of conditions: 30 seconds,95 degree C; 1 minute, 56 degree C; 1 minute, 70 degree C. This cycle is repeated 32 times followed by one 5 minute cycle at 70 degree C. Plant DNA is used as template in addition to a somatic cell hybrid panel containing individual chromosomes or chromosome fragments (Bios, Inc). The reactions are
25 analyzed on either 8% polyacrylamide gels or 3.5 % agarose gels. Chromosome mapping is determined by the presence of an approximately 100 bp PCR fragment in the particular somatic cell hybrid.

Example 30 - Bacterial Expression of a Polypeptide.

30 A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers used to amplify the cDNA insert should preferably contain restriction sites, such as BamHI and XbaI, at the 5' end of the primers in order to clone the amplified product
35 into the expression vector. For example, BamHI and XbaI correspond to the restriction enzyme sites on the bacterial expression vector pQE-9. (Qiagen, Inc., Chatsworth, CA). This plasmid vector encodes antibiotic resistance (Ampr), a bacterial origin of replication (ori), an IPTG-regulatable promoter/operator (P/O), a

5 ribosome binding site (RBS), a 6-histidine tag (6-His), and restriction enzyme cloning sites.

The pQE-9 vector is digested with BamHI and XbaI and the amplified fragment is ligated into the pQE-9 vector maintaining the reading frame initiated at the bacterial RBS. The ligation mixture is then used to transform the E. coli strain
10 M15/rep4 (Qiagen, Inc.) which contains multiple copies of the plasmid pREP4, that expresses the lacI repressor and also confers kanamycin resistance (Kanr). Transformants are identified by their ability to grow on LB plates and ampicillin/kanamycin resistant colonies are selected. Plasmid DNA is isolated and confirmed by restriction analysis.

15 Clones containing the desired constructs are grown overnight (O/N) in liquid culture in LB media supplemented with both Amp (100 ug/ml) and Kan (25 ug/ml). The O/N culture is used to inoculate a large culture at a ratio of 1:100 to 1:250. The cells are grown to an optical density 600 (O.D.600) of between 0.4 and 0.6. IPTG (Isopropyl-B-D-thiogalacto pyranoside) is then added to a final concentration of 1
20 mM. IPTG induces by inactivating the lacI repressor, clearing the P/O leading to increased gene expression.

Cells are grown for an extra 3 to 4 hours. Cells are then harvested by centrifugation (20 mins at 6000Xg). The cell pellet is solubilized in the chaotropic agent 6 Molar Guanidine HCl by stirring for 3-4 hours at 4 degree C. The cell debris
25 is removed by centrifugation, and the supernatant containing the polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (available from QIAGEN, Inc., supra). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure (for details see: The QIAexpressionist (1995) QIAGEN, Inc., supra).

30 Briefly, the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl.
35 Alternatively, the protein can be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4,

5 containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins are eluted by the addition of 250 mM imidazole. Imidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4 degree C or frozen at -80 degree C.

10

Example 31 - Purification of a Polypeptide from an Inclusion Body.

The following alternative method can be used to purify a polypeptide expressed in E coli when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10 degree C.

15 Upon completion of the production phase of the E. coli fermentation, the cell culture is cooled to 4-10 degree C and the cells harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution
20 containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells are then lysed by passing the solution through a microfluidizer (Microfluidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by
25 centrifugation at 7000 xg for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 xg centrifugation for 15 min., the pellet is discarded and the polypeptide containing supernatant is incubated at 4 degree
30 C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 xg) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4 degree C
35 without mixing for 12 hours prior to further purification steps.

To clarify the refolded polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 um membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The

5 filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 nm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

10 Fractions containing the polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40
15 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A280 monitoring of the effluent. Fractions containing the polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

20 The resultant polypeptide should exhibit greater than 95% purity after the above refolding and purification steps. No major contaminant bands should be observed from Commassie blue stained 16% SDS-PAGE gel when 5 ug of purified protein is loaded. The purified protein can also be tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL
25 assays.

Example 32 - Cloning and Expression of a Polypeptide in a Baculovirus Expression System.

In this example, the plasmid shuttle vector pAc373 is used to insert a
30 polynucleotide into a baculovirus to express a polypeptide. A typical baculovirus expression vector contains the strong polyhedrin promoter of the *Autographa californica* nuclear polyhedrosis virus (AcMNPV) followed by convenient restriction sites, which may include, for example BamHI, Xba I and Asp718. The polyadenylation site of the simian virus 40 ("SV40") is often used for efficient
35 polyadenylation. For easy selection of recombinant virus, the plasmid contains the beta-galactosidase gene from *E. coli* under control of a weak *Drosophila* promoter in the same orientation, followed by the polyadenylation signal of the polyhedrin gene. The inserted genes are flanked on both sides by viral sequences for cell-mediated

5 homologous recombination with wild-type viral DNA to generate a viable virus that express the cloned polynucleotide.

Many other baculovirus vectors can be used in place of the vector above, such as pVL941 and pAcIM1, as one skilled in the art would readily appreciate, as long as the construct provides appropriately located signals for transcription, translation,
10 secretion and the like, including a signal peptide and an in-frame AUG as required. Such vectors are described, for instance, in Luckow et al., *Virology* 170:31-39 (1989).

A polynucleotide encoding a polypeptide of the present invention is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' ends of the DNA sequence, as outlined in Example 1, to synthesize insertion fragments. The primers
15 used to amplify the cDNA insert should preferably contain restriction sites at the 5' end of the primers in order to clone the amplified product into the expression vector. Specifically, the cDNA sequence contained in the deposited clone, including the AUG initiation codon and the naturally associated leader sequence identified elsewhere herein (if applicable), is amplified using the PCR protocol described in Example 1. If
20 the naturally occurring signal sequence is used to produce the protein, the vector used does not need a second signal peptide. Alternatively, the vector can be modified to include a baculovirus leader sequence, using the standard methods described in Summers et al., "A Manual of Methods for Baculovirus Vectors and Insect Cell Culture Procedures," Texas Agricultural Experimental Station Bulletin No. 1555
25 (1987).

The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

30 The plasmid is digested with the corresponding restriction enzymes and optionally, can be dephosphorylated using calf intestinal phosphatase, using routine procedures known in the art. The DNA is then isolated from a 1% agarose gel using a commercially available kit ("GeneClean" BIO 101 Inc., La Jolla, Ca.).

The fragment and the dephosphorylated plasmid are ligated together with T4
35 DNA ligase. *E. coli* HB101 or other suitable *E. coli* hosts such as XL-1 Blue (Stratagene Cloning Systems, La Jolla, CA) cells are transformed with the ligation mixture and spread on culture plates. Bacteria containing the plasmid are identified by digesting DNA from individual colonies and analyzing the digestion product by

5 gel electrophoresis. The sequence of the cloned fragment is confirmed by DNA sequencing.

Five ug of a plasmid containing the polynucleotide is co-transformed with 1.0 ug of a commercially available linearized baculovirus DNA ("BaculoGoldtm baculovirus DNA", Pharmingen, San Diego, CA), using the lipofection method
10 described by Felgner et al., Proc. Natl. Acad. Sci. USA 84:7413-7417 (1987). One ug of BaculoGoldtm virus DNA and 5ug of the plasmid are mixed in a sterile well of a microtiter plate containing 50ul of serum-free Grace's medium (Life Technologoes Inc., Gaithersburg, MD). Afterwards, 10 ul Lipofectin plus 90 ul Grace's medium are added, mixed and incubated for 15 minutes at room temperature. Then the
15 transfection mixture is added drop-wise to Sf9 insect cells (ATCC CRL 1711) seeded in a 35 mm tissue culture plate with 1 ml Grace's medium without serum. The plate is then incubated for 5 hours at 27 degrees C. The transfection solution is then removed from the plate and 1 ml of Grace's insect medium supplemented with 10% fetal calf serum is added. Cultivation is then continued at 27 degrees C for four days.

20 After four days the supernatant is collected and a plaque assay is performed, as described by Summers and Smith, supra. An agarose gel with "Blue Gal" (Life Technologies Inc., Gaithersburg) is used to allow easy identification and isolation of gal-expressing clones, which produce blue-stained plaques. (A detailed description of a "plaque assay" of this type can also be found in the user's guide for insect cell
25 culture and baculovirology distributed by Life Technologies Inc., Gaithersburg, page 9-10.) After appropriate incubation, blue stained plaques are picked with the tip of a micropipettor (e.g., Eppendorf). The agar containing the recombinant viruses is then resuspended in a microcentrifuge tube containing 200 ul of Grace's medium and the suspension containing the recombinant baculovirus is used to infect Sf9 cells seeded
30 in 35 mm dishes. Four days later the supernatants of these culture dishes are harvested and then they are stored at 4 degree C.

To verify the expression of the polypeptide, Sf9 cells are grown in Grace's medium supplemented with 10% heat-inactivated FBS. The cells are infected with the recombinant baculovirus containing the polynucleotide at a multiplicity of
35 infection ("MOI") of about 2. If radiolabeled proteins are desired, 6 hours later the medium is removed and is replaced with SF900 II medium minus methionine and cysteine (available from Life Technologies Inc., Rockville, MD). After 42 hours, 5 uCi of 35S-methionine and 5 uCi 35S-cysteine (available from Amersham) are added.

5 The cells are further incubated for 16 hours and then are harvested by centrifugation. The proteins in the supernatant as well as the intracellular proteins are analyzed by SDS-PAGE followed by autoradiography (if radiolabeled).

Microsequencing of the amino acid sequence of the amino terminus of purified protein may be used to determine the amino terminal sequence of the
10 produced protein.

Example 33 - Expression of a Polypeptide in Mammalian Cells.

The polypeptide of the present invention can be expressed in a mammalian cell. A typical mammalian expression vector contains a promoter element, which
15 mediates the initiation of transcription of mRNA, a protein coding sequence, and signals required for the termination of transcription and polyadenylation of the transcript. Additional elements include enhancers, Kozak sequences and intervening sequences flanked by donor and acceptor sites for RNA splicing. Highly efficient transcription is achieved with the early and late promoters from SV40, the long
20 terminal repeats (LTRs) from Retroviruses, e.g., RSV, HTLV1, HIV1 and the early promoter of the cytomegalovirus (CMV). However, cellular elements can also be used (e.g., the human actin promoter).

Suitable expression vectors for use in practicing the present invention include, for example, vectors such as pSVL and pMSG (Pharmacia, Uppsala, Sweden),
25 pRSVcat (ATCC 37152), pSV2dhfr (ATCC 37146), pBC12MI (ATCC 67109), pCMVSPORT 2.0, and pCMVSPORT 3.0. Mammalian host cells that could be used include, human HeLa, 293, H9 and Jurkat cells, mouse NIH3T3 and C127 cells, Cos 1, Cos 7 and CV1, quail QC1-3 cells, mouse L cells and Chinese hamster ovary (CHO) cells.

30 Alternatively, the polypeptide can be expressed in stable cell lines containing the polynucleotide integrated into a chromosome. The co-transformation with a selectable marker such as dhfr, gpt, neomycin, hygromycin allows the identification and isolation of the transformed cells.

The transformed gene can also be amplified to express large amounts of the
35 encoded protein. The DHFR (dihydrofolate reductase) marker is useful in developing cell lines that carry several hundred or even several thousand copies of the gene of interest. (See, e.g., Alt, F. W., et al., J. Biol. Chem. 253:1357-1370 (1978); Hamlin, J. L. and Ma, C., Biochem. et Biophys. Acta, 1097:107-143 (1990); Page, M. J. and

5 Sydenham, M. A., *Biotechnology* 9:64-68 (1991.) Another useful selection marker is the enzyme glutamine synthase (GS) (Murphy et al., *Biochem J.* 227:277-279 (1991); Bebbington et al., *Bio/Technology* 10:169-175 (1992). Using these markers, the mammalian cells are grown in selective medium and the cells with the highest resistance are selected. These cell lines contain the amplified gene(s) integrated into a
10 chromosome. Chinese hamster ovary (CHO) and NSO cells are often used for the production of proteins.

A polynucleotide of the present invention is amplified according to the protocol outlined in Example 1. If the naturally occurring signal sequence is used to produce the protein, the vector does not need a second signal peptide. Alternatively,
15 if the naturally occurring signal sequence is not used, the vector can be modified to include a heterologous signal sequence. (See, e.g., WO 96/34891.) The amplified fragment is isolated from a 1% agarose gel using a commercially available kit ("GeneClean," BIO 101 Inc., La Jolla, Ca.). The fragment then is digested with appropriate restriction enzymes and again purified on a 1% agarose gel.

20 The amplified fragment is then digested with the same restriction enzyme and purified on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC6 using, for instance, restriction enzyme analysis.

25 Chinese hamster ovary cells lacking an active DHFR gene is used for transformation. Five μ g of an expression plasmid is cotransformed with 0.5 μ g of the plasmid pSVneo using lipofectin (Felgner et al., supra). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded
30 in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM,
35 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100 -

5 200 uM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

In addition to the method provided above, other methods of expressing polypeptides, preferably polypeptides of the present invention are known in the art. For example, US Patent No. 6,066,781 (which is hereby incorporated by reference
10 herein in its entirety) describes a chimeric gene consisting of the encoding polynucleotides for the N-terminal moiety corresponding to a portion of the rice alpha-amylase signal sequence peptide (MKNTSSLCLLLLVLCSLTCNSGQA SEQ ID NO:33 operatively linked, to the encoding polynucleotide sequence of the protein of interest – in this case, a polypeptide of the present invention. This signal
15 sequence peptide may be operatively substituted for the native signal sequence of a polypeptide of the present invention as a heterologous signal sequence. Such a method of producing mature forms of the inventive polypeptides is encompassed by the present invention and may be used either alone or in conjunction with other methods known in the art and/or disclosed herein.

20 Moreover, enhanced protein production of recombinant proteins in higher plants has recently been obtained by N-terminal fusion of a ubiquitin or a cucumber mosaic virus coat protein peptide (see International Publication No. WO 00/36129). Such methods may be applied to increase the expression of a polypeptide of the present invention in a suitable plant host.

25

Example 34 - Protein Fusions.

The polypeptides of the present invention are preferably fused to other proteins. These fusion proteins can be used for a variety of applications. For example, fusion of the present polypeptides to His-tag, HA-tag, protein A, IgG
30 domains, and maltose binding protein facilitates purification. (See Example described herein; see also EP A 394,827; Traunecker, et al., Nature 331:84-86 (1988).) Similarly, fusion to IgG-1, IgG-3, and albumin increases the halflife time in vivo. Nuclear localization signals fused to the polypeptides of the present invention can target the protein to a specific subcellular localization, while covalent heterodimer or
35 homodimers can increase or decrease the activity of a fusion protein. Fusion proteins can also create chimeric molecules having more than one function. Finally, fusion proteins can increase solubility and/or stability of the fused protein compared to the non-fused protein. All of the types of fusion proteins described above can be made by

5 modifying the following protocol, which outlines the fusion of a polypeptide to an IgG molecule.

Briefly, the human Fc portion of the IgG molecule can be PCR amplified, using primers that span the 5' and 3' ends of the sequence described below. These primers also should have convenient restriction enzyme sites that will facilitate
10 cloning into an expression vector, preferably a mammalian expression vector. Note that the polynucleotide is cloned without a stop codon, otherwise a fusion protein will not be produced.

The naturally occurring signal sequence may be used to produce the protein. Alternatively, if the naturally occurring signal sequence is not used, the vector can be
15 modified to include a heterologous signal sequence. (See, e.g., WO 96/34891 and/or US Patent No. 6,066,781, supra.)

Human IgG Fc region:

GGGATCCGGAGCCCAAATCTTCTGACAAAACACACATGCCACCGTGC
20 CCAGCACCTGAATTCGAGGGTGCACCGTCAGTCTTCTTCCCCCAAAA
CCCAAGGACACCCTCATGATCTCCCGGACTCCTGAGGTCACATGCGTGGT
GGTGGACGTAAGCCACGAAGACCCTGAGGTCAAGTTCAACTGGTACGTGG
ACGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGCGGGAGGAGCAGTA
CAACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGACT
25 GGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGCCCTCCCA
ACCCCATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAAC
CACAGGTGTACACCCTGCCCCATCCCGGGATGAGCTGACCAAGAACCAG
GTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCAAGCGACATCGCCGT
GGAGTGGGAGAGCAATGGGCAGCCGAGAAACAACACTACAAGACCACGCCT
30 CCCGTGCTGGACTCCGACGGTCTCTTCTTCTTCTACAGCAAGCTCACCGTG
GACAAGAGCAGGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCA
TGAGGCTCTGCACAACCACTACACGCAGAAGAGCCTCTCCCTGTCTCCGG
GTAAATGAGTGCGACGGCCGCGACTCTAGAGGAT (SEQ ID NO:11)

35 **Example 35 - Production of an Antibody from a Polypeptide.**

The antibodies of the present invention can be prepared by a variety of methods. (See, Current Protocols, Chapter 2.) As one example of such methods, cells expressing a polypeptide of the present invention are administered to an animal to

5 induce the production of sera containing polyclonal antibodies. In a preferred method, a preparation of the protein is prepared and purified to render it substantially free of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In the most preferred method, the antibodies of the present invention are
10 monoclonal antibodies (or protein binding fragments thereof). Such monoclonal antibodies can be prepared using hybridoma technology. (Köhler et al., *Nature* 256:495 (1975); Köhler et al., *Eur. J. Immunol.* 6:511 (1976); Köhler et al., *Eur. J. Immunol.* 6:292 (1976); Hammerling et al., in: *Monoclonal Antibodies and T-Cell Hybridomas*, Elsevier, N.Y., pp. 563-681 (1981).) In general, such procedures
15 involve immunizing an animal (preferably a mouse) with polypeptide or, more preferably, with a polypeptide-expressing cell. Such cells may be cultured in any suitable tissue culture medium; however, it is preferable to culture cells in Earle's modified Eagle's medium supplemented with 10% fetal bovine serum (inactivated at about 56 degrees C), and supplemented with about 10 g/l of nonessential amino acids,
20 about 1,000 U/ml of penicillin, and about 100 ug/ml of streptomycin.

The splenocytes of such mice are extracted and fused with a suitable myeloma cell line. Any suitable myeloma cell line may be employed in accordance with the present invention; however, it is preferable to employ the parent myeloma cell line (SP2O), available from the ATCC. After fusion, the resulting hybridoma cells are
25 selectively maintained in HAT medium, and then cloned by limiting dilution as described by Wands et al. (*Gastroenterology* 80:225-232 (1981).) The hybridoma cells obtained through such a selection are then assayed to identify clones which secrete antibodies capable of binding the polypeptide.

Alternatively, additional antibodies capable of binding to the polypeptide can
30 be produced in a two-step procedure using anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and therefore, it is possible to obtain an antibody that binds to a second antibody. In accordance with this method, protein specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells,
35 and the hybridoma cells are screened to identify clones that produce an antibody whose ability to bind to the protein-specific antibody can be blocked by the polypeptide. Such antibodies comprise anti-idiotypic antibodies to the protein-

5 specific antibody and can be used to immunize an animal to induce formation of further protein-specific antibodies.

It will be appreciated that Fab and F(ab')₂ and other fragments of the antibodies of the present invention may be used according to the methods disclosed herein. Such fragments are typically produced by proteolytic cleavage, using
10 enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')₂ fragments). Alternatively, protein-binding fragments can be produced through the application of recombinant DNA technology or through synthetic chemistry.

For in vivo use of antibodies in humans, it may be preferable to use "humanized" chimeric monoclonal antibodies. Such antibodies can be produced using
15 genetic constructs derived from hybridoma cells producing the monoclonal antibodies described above. Methods for producing chimeric antibodies are known in the art. (See, for review, Morrison, Science 229:1202 (1985); Oi et al., BioTechniques 4:214 (1986); Cabilly et al., U.S. Patent No. 4,816,567; Taniguchi et al., EP 171496; Morrison et al., EP 173494; Neuberger et al., WO 8601533; Robinson et al., WO
20 8702671; Boulianne et al., Nature 312:643 (1984); Neuberger et al., Nature 314:268 (1985).)

Moreover, in a more preferred method, the antibodies directed against the polypeptides of the present invention may be produced in plants. Specific methods are disclosed in US Patent Nos. 5,959,177, and 6,080,560, which are hereby
25 incorporated in their entirety herein. The methods not only describe methods of expressing antibodies, but also the means of assembling foreign multimeric proteins in plants (i.e., antibodies, etc.), and the subsequent secretion of such antibodies from the plant.

30 **Example 36 – Antibody Mediated Down-Regulation of Plant Proteins.**

The process of genetically modifying a plant to modulate specific characteristics, to introduce novel traits, or to inhibit endogenous traits represents a significant area of research in the agricultural field. Recently, a new method of modulating endogenous gene expression using antibodies has been elucidated (see,
35 International Publication Number WO 00/05391, which is hereby incorporated in its entirety herein). In this example, the researchers were able to achieve 40-70% inhibition of an endogenous plant protein through the use of a single-chain antibody construct directed towards the plant protein.

5 The method is directed towards the production of monoclonal antibodies, specifically, single chain antibodies, specific to endogenous transit peptides in a plant in an effort to decrease steady state levels of such transit peptides within the plant. The method is comprised of the following steps: I) generating monoclonal antibodies to a specific plant, II) cloning the gene for said monoclonal antibody, III) creating an
10 expression vector comprising a fusion of the heavy-chain and light chain gene sequences of said monoclonal antibody gene downstream of the p67 leader peptide, and under the control of a constitutive plant promoter, IV) optimizing the codons of said heavy-chain and light chain fusion vector for efficient expression of the gene encoded thereof in a plant, and V), transforming a plant with said heavy-chain and
15 light chain fusion expression vector.

 The skilled artisan would appreciate the methods described therein (WO 00/05391), and would have the ability to apply such methods to inhibiting the steady-state expression levels of any of the polypeptides of the present invention, including variants, and fragments, thereof. The skilled artisan would appreciate that any leader
20 peptide (i.e., signal sequence) from a plant protein could be used in creating the heavy-chain and light chain fusion vector. The skilled artisan would also appreciate that different plant species may have different codon usage requirements, and thus, the decision to optimize the codons of the heavy-chain and light chain fusion vector would be affected according to the codons required for that particular plant species.

25 The method could not only be applied to transit peptides, but also to secreted proteins, membrane proteins, receptors, and ligands. The method could also be applied in combination with other antibody production methods in plants. For example, antibodies directed towards polypeptides of the present invention may inhibit specific traits in a plant which could increase the plants defense mechanisms to
30 pathogens. Thus, where such an antibody was expressed, another antibody could be expressed in combination with the first, to inhibiting the pathogenicity of a plant pathogen by directing the expression of antibodies directed towards pathogenic proteins (e.g., those proteins critical to the initiating events of infection, such as the BUF1 gene from *M.grisea*, stage two juvenile salivary gland proteins which include, svp30, scp31a, scp31b, scp32, scp32, scp39, and scp49 from *G.rostochiensis* (WO
35 96/22372), etc.). Such a combination would also be of value where the second “anti-pathogenic” antibody is an antibody directed towards a pathogen and fused to a toxic

5 protein wherein such a toxin could be chitinase, glucanase, lysozyme, BT, or colicin F, for example (see WO 96/09398), etc.).

As described elsewhere herein, the method could also be used as a means of inhibiting allergic reactions to plant antigens in humans, mammals, animals, etc., by directing the production of a single chain antibody construct specific towards said
10 plant antigen in the plant (via transgenic methodology). In the latter example, the plant would not be limited to edible plants, as inhibiting the production of such a plant antigen would provide benefit to a human by removing the antigen from the humans environment, for example, irrespective of whether the plant was ingested.

Of particular interest to this example, is the fact that secretion of functional
15 antibody through the plasma membrane of plant cells has been reported for protoplasts isolated from transgenic plants and for callus cells adapted to suspension culture (Hein et al., *Biotechnol. Prog.* 7:455-561, 1991). However, the levels of secreted antibody detected in both culture systems were extremely low. In other studies, cultured tobacco cells were transformed with a gene encoding a synthetic
20 antibody derivative expressed as a single chain consisting of both the heavy- and light-chain variable domains of the intact immunoglobulin joined together by a flexible peptide linker (Pluckthun, *Immunol. Rev.* 130:151-188, 1991; and Bird et al., *Science* 242:423-426, 1988). This synthetic single-chain antibody retained the full antigen-binding potential of the intact immunoglobulin but accumulated in the
25 extracellular apoplastic space of the transformed cells (Firek et al., *Plant Molecular Biology* 23:861-870, 1993), indicating that the antibody was being transported through the plasma membrane but not through the cell wall to the external environment. Moreover, recent studies have shown that increased antibody production in a plant, and heterologous protein expression, in general, could be
30 increased by including in the plant culture medium a protein stabilizing agent (e.g., polyvinylpyrrolidone), see US Patent No. 6,020,169, which is hereby incorporated by reference in its entirety herein.

Example 37 - Transformation of Dicotyledons.

35 The polynucleotides of the present invention, including the polynucleotides encoding the polypeptides of the present invention, in addition to the polynucleotides encoding antibodies directed against the polypeptides of the present invention may be used to transform monocotyledons in an effort to confer specific traits into the plant.

5 Such polynucleotides may be either the full-length polynucleotide, fragments, the complementary strand, or variants thereof, and may be either by themselves or operably fused to heterologous polynucleotides as described in more detail elsewhere herein.

Transformation techniques for dicotyledons are well known in the art and
10 include *Agrobacterium*-based techniques and techniques which do not require *Agrobacterium*. Non-*Agrobacterium* techniques involve the uptake of exogenous genetic material directly by protoplasts or cells. This can be accomplished by PEG or electroporation mediated uptake, particle bombardment-mediated delivery, or microinjection. Examples of these techniques are described by Paszkowski et al.,
15 *EMBO J* 3: 2717-2722 (1984), Potrykus et al., *Mol. Gen. Genet.* 199: 169-177 (1985), Reich et al., *Biotechnology* 4: 1001-1004 (1986), and Klein et al., *Nature* 327: 70-73 (1987). In each case the transformed cells are regenerated to whole plants using standard techniques known in the art.

Agrobacterium-mediated transformation is a preferred technique for
20 transformation of dicotyledons because of its high efficiency of transformation and its broad utility with many different species. The many crop species which are routinely transformable by *Agrobacterium* include tobacco, tomato, sunflower, cotton, oilseed rape, potato, soybean, alfalfa and poplar (EP 0 317 511 (cotton), EP 0 249 432 (tomato, to Calgene), WO 87/07299 (*Brassica*, to Calgene), U.S. Pat. No. 4,795,855 (poplar)). *Agrobacterium* transformation typically involves the transfer of the binary
25 vector carrying the foreign DNA of interest (e.g. pCIB200 or pCIB2001) to an appropriate *Agrobacterium* strain which may depend of the complement of vir genes carried by the host *Agrobacterium* strain either on a co-resident Ti plasmid or chromosomally (e.g. strain CIB542 for pCIB200 and pCIB2001 (Uknes et al. *Plant*
30 *Cell* 5: 159-169 (1993)). The transfer of the recombinant binary vector to *Agrobacterium* is accomplished by a triparental mating procedure using *E. coli* carrying the recombinant binary vector, a helper *E. coli* strain which carries a plasmid such as pRK2013 and which is able to mobilize the recombinant binary vector to the target *Agrobacterium* strain. Alternatively, the recombinant binary vector can be
35 transferred to *Agrobacterium* by DNA transformation (Hofgen & Willmitzer, *Nucl. Acids Res.* 16: 9877(1988)).

Transformation of the target plant species by recombinant *Agrobacterium* usually involves co-cultivation of the *Agrobacterium* with explants from the plant and

5 follows protocols well known in the art. Transformed tissue is regenerated on selectable medium carrying the antibiotic or herbicide resistance marker present between the binary plasmid.T-DNA borders.

Other methods for the transformation of dicotyledons are known in the art. Thus, this example should be not construed as limiting the scope of the invention to
10 only those examples illustrated above or elsewhere herein.

Example 38- Transformation of Monocotyledons.

The polynucleotides of the present invention, including the polynucleotides encoding the polypeptides of the present invention, in addition to the polynucleotides
15 encoding antibodies directed against the polypeptides of the present invention may be used to transform monocotyledons in an effort to confer specific traits into the plant. Such polynucleotides may be either the full-length polynucleotide, fragments, the complementary strand, or variants thereof, and may be either by themselves or operably fused to heterologous polynucleotides as described in more detail elsewhere
20 herein.

Transformation of most monocotyledon species has now also become routine. Preferred techniques include direct gene transfer into protoplasts using PEG or electroporation techniques, and particle bombardment into callus tissue. Transformations can be undertaken with a single DNA species or multiple DNA
25 species (i.e. co-transformation) and both these techniques are suitable for use with this invention. Co-transformation may have the advantage of avoiding complex vector construction and of generating transgenic plants with unlinked loci for the gene of interest and the selectable marker, enabling the removal of the selectable marker in subsequent generations, should this be regarded desirable. However, a disadvantage
30 of the use of co-transformation is the less than 100% frequency with which separate DNA species are integrated into the genome (Schocher et al. *Biotechnology* 4: 1093-1096 (1986)).

Patent Applications EP 0 292 435 (to Ciba-Geigy), EP 0 392 225 (to Ciba-Geigy) and WO 93/07278 (to Ciba-Geigy) describe techniques for the preparation of
35 callus and protoplasts from an elite inbred line of maize, transformation of protoplasts using PEG or electroporation, and the regeneration of maize plants from transformed protoplasts. Gordon-Kamm et al., *Plant Cell* 2: 603-618 (1990)) and Fromm et al., *Biotechnology* 8: 833-839 (1990)) have published techniques for transformation of

5 A188-derived maize line using particle bombardment. Furthermore, application WO
93/07278 (to Ciba-Geigy) and Koziel et al., *Biotechnology* 11: 194-200 (1993))
describe techniques for the transformation of elite inbred lines of maize by particle
bombardment. This technique utilizes immature maize embryos of 1.5-2.5 mm length
excised from a maize ear 14-15 days after pollination and a PDS-1000He Biolistics
10 device for bombardment.

Transformation of rice can also be undertaken by direct gene transfer
techniques utilizing protoplasts or particle bombardment. Protoplast-mediated
transformation has been described for Japonica-types and Indica-types (Zhang et al.,
Plant Cell Rep 7: 379-384 (1988); Shimamoto et al. *Nature* 338: 274-277 (1989);
15 Datta et al. *Biotechnology* 8: 736-740 (1990)). Both types are also routinely
transformable using particle bombardment (Christou et al. *Biotechnology* 9: 957-962
(1991)).

Patent Application EP 0 332 581 (to Ciba-Geigy) describes techniques for the
generation, transformation and regeneration of Pooideae protoplasts. These techniques
20 allow the transformation of *Dactylis* and wheat. Furthermore, wheat transformation
was been described by Vasil et al., *Biotechnology* 10: 667-674 (1992)) using particle
bombardment into cells of type C long-term regenerable callus, and also by Vasil et
al., *Biotechnology* 11: 1553-1558 (1993)) and Weeks et al., *Plant Physiol.* 102: 1077-
1084 (1993) using particle bombardment of immature embryos and immature
25 embryo-derived callus. A preferred technique for wheat transformation, however,
involves the transformation of wheat by particle bombardment of immature embryos
and includes either a high sucrose or a high maltose step prior to gene delivery. Prior
to bombardment, any number of embryos (0.75-1 mm in length) are plated onto MS
medium with 3% sucrose (Murashige & Skoog, *Physiologia Plantarum* 15: 473-497
30 (1962)) and 3 mg/l 2,4-D for induction of somatic embryos which is allowed to
proceed in the dark. On the chosen day of bombardment, embryos are removed from
the induction medium and placed onto the osmoticum (i.e. induction medium with
sucrose or maltose added at the desired concentration, typically 15%). The embryos
are allowed to plasmolyze for 2-3 h and are then bombarded. Twenty embryos per
35 target plate is typical, although not critical. An appropriate gene-carrying plasmid
(such as pCIB3064 or pSG35) is precipitated onto micrometer size gold particles
using standard procedures. Each plate of embryos is shot with the DuPont Biolistics,
helium device using a burst pressure of .about.1000 psi using a standard 80 mesh

5 screen. After bombardment, the embryos are placed back into the dark to recover for
about 24 h (still on osmoticum). After 24 hrs, the embryos are removed from the
osmoticum and placed back onto induction medium where they stay for about a
month before regeneration. Approximately one month later the embryo explants with
developing embryogenic callus are transferred to regeneration medium (MS+1
10 mg/liter NAA, 5 mg/liter GA), further containing the appropriate selection agent (10
mg/l basta in the case of pCIB3064 and 2 mg/l methotrexate in the case of pSOG35).
After approximately one month, developed shoots are transferred to larger sterile
containers known as "GA7s" which contained half-strength MS, 2% sucrose, and the
same concentration of selection agent.

15 An additional method of transforming monocots is found in International
Publication No. WO 00/12734, and describes the application of the Ac-Ds transposon
system to the insertion of transgenes into plants.

Other methods for the transformation of monocotyledons are known in the art.
Thus, this example should be not construed as limiting the scope of the invention to
20 only those examples illustrated above or elsewhere herein.

Example 39 - Method of Enhancing the Biological Activity/Functional Characteristics of Invention through Molecular Evolution.

Although many of the most biologically active proteins known are highly
25 effective for their specified function in an organism, they often possess characteristics
that make them undesirable for transgenic, therapeutic, agricultural, and/or industrial
applications. Among these traits, a short physiological half-life is the most prominent
problem, and is present either at the level of the protein, or the level of the proteins
mRNA. The ability to extend the half-life, for example, would be particularly
30 important for a proteins use in gene therapy, transgenic plant or animal production,
the bioprocess production and purification of the protein, and use of the protein as a
chemical modulator (e.g., herbicide, insectide, etc.), among others. Therefore, there is
a need to identify novel variants of isolated proteins possessing characteristics which
enhance their application as a therapeutic for treating diseases of both plant and
35 animal origin, in addition to the proteins applicability to common industrial and
agricultural applications.

Thus, one aspect of the present invention relates to the ability to enhance
specific characteristics of invention through directed molecular evolution. Such an

5 enhancement may, in a non-limiting example, benefit the inventions utility as an essential component in a kit, the inventions physical attributes such as its solubility, structure, or codon optimization, the inventions specific biological activity, including any associated enzymatic activity, the proteins enzyme kinetics, the proteins K_i , K_{cat} , K_m , V_{max} , K_d , protein-protein activity, protein-DNA binding activity, 10 antagonist/inhibitory activity (including direct or indirect interaction), agonist activity (including direct or indirect interaction), the proteins antigenicity (e.g., where it would be desirable to either increase or decrease the antigenic potential of the protein), the immunogenicity of the protein, the ability of the protein to form dimers, trimers, or multimers with either itself or other proteins, the antigenic efficacy of the invention, 15 including its subsequent use a preventative treatment for disease or disease states, or as an effector for targeting diseased genes. Moreover, the ability to enhance specific characteristics of a protein may also be applicable to changing the characterized activity of an enzyme to an activity completely unrelated to its initially characterized activity. Other desirable enhancements of the invention would be specific to each 20 individual protein, and would thus be well known in the art and contemplated by the present invention.

Directed evolution is comprised of several steps. The first step is to establish a library of variants for the gene or protein of interest. The most important step is to then select for those variants that entail the activity you wish to identify. The design 25 of the screen is essential since your screen should be selective enough to eliminate non-useful variants, but not so stringent as to eliminate all variants. The last step is then to repeat the above steps using the best variant from the previous screen. Each successive cycle, can then be tailored as necessary, such as increasing the stringency of the screen, for example.

30 Over the years, there have been a number of methods developed to introduce mutations into macromolecules. Some of these methods include, random mutagenesis, "error-prone" PCR, chemical mutagenesis, site-directed mutagenesis, and other methods well known in the art (for a comprehensive listing of current mutagenesis methods, see Maniatis, Molecular Cloning: A Laboratory Manual, Cold 35 Spring Harbor Press, Cold Spring, NY (1982)). Typically, such methods have been used, for example, as tools for identifying the core functional region(s) of a protein or the function of specific domains of a protein (if a multi-domain protein). However,

5 such methods have more recently been applied to the identification of macromolecule variants with specific or enhanced characteristics.

Random mutagenesis has been the most widely recognized method to date. Typically, this has been carried out either through the use of “error-prone” PCR (as described in Moore, J., et al, Nature Biotechnology 14:458, (1996), or through the application of randomized synthetic oligonucleotides corresponding to specific regions of interest (as described by Derbyshire, K.M. et al, Gene, 46:145-152, (1986), and Hill, DE, et al, Methods Enzymol., 55:559-568, (1987). Both approaches have limits to the level of mutagenesis that can be obtained. However, either approach enables the investigator to effectively control the rate of mutagenesis. This is particularly important considering the fact that mutations beneficial to the activity of the enzyme are fairly rare. In fact, using too high a level of mutagenesis may counter or inhibit the desired benefit of a useful mutation.

While both of the aforementioned methods are effective for creating randomized pools of macromolecule variants, a third method, termed “DNA Shuffling”, or “sexual PCR” (WPC, Stemmer, PNAS, 91:10747, (1994)) has recently been elucidated. DNA shuffling has also been referred to as “directed molecular evolution”, “exon-shuffling”, “directed enzyme evolution”, “in vitro evolution”, and “artificial evolution”. Such reference terms are known in the art and are encompassed by the invention. This new, preferred, method apparently overcomes the limitations of the previous methods in that it not only propagates positive traits, but simultaneously eliminates negative traits in the resulting progeny.

DNA shuffling accomplishes this task by combining the principal of in vitro recombination, along with the method of “error-prone” PCR. In effect, you begin with a randomly digested pool of small fragments of your gene, created by Dnase I digestion, and then introduce said random fragments into an “error-prone” PCR assembly reaction. During the PCR reaction, the randomly sized DNA fragments not only hybridize to their cognate strand, but also may hybridize to other DNA fragments corresponding to different regions of the polynucleotide of interest – regions not typically accessible via hybridization of the entire polynucleotide. Moreover, since the PCR assembly reaction utilizes “error-prone” PCR reaction conditions, random mutations are introduced during the DNA synthesis step of the PCR reaction for all of the fragments -further diversifying the potential hybridation sites during the annealing step of the reaction.

5 A variety of reaction conditions could be utilized to carry-out the DNA shuffling reaction. However, specific reaction conditions for DNA shuffling are provided, for example, in PNAS, 91:10747, (1994). Briefly:

Prepare the DNA substrate to be subjected to the DNA shuffling reaction. Preparation may be in the form of simply purifying the DNA from contaminating
10 cellular material, chemicals, buffers, oligonucleotide primers, deoxynucleotides, RNAs, etc., and may entail the use of DNA purification kits as those provided by Qiagen, Inc., or by the Promega, Corp., for example.

Once the DNA substrate has been purified, it would be subjected to Dnase I digestion. About 2-4ug of the DNA substrate(s) would be digested with .0015 units
15 of Dnase I (Sigma) per ul in 100ul of 50mM Tris-HCL, pH 7.4/1mM MgCl₂ for 10-20 min. at room temperature. The resulting fragments of 10-50bp could then be purified by running them through a 2% low-melting point agarose gel by electrophoresis onto DE81 ion-exchange paper (Whatman) or could be purified using Microcon concentrators (Amicon) of the appropriate molecular weight cutoff, or
20 could use oligonucleotide purification columns (Qiagen), in addition to other methods known in the art. If using DE81 ion-exchange paper, the 10-50bp fragments could be eluted from said paper using 1M NaCl, followed by ethanol precipitation.

The resulting purified fragments would then be subjected to a PCR assembly reaction by re-suspension in a PCR mixture containing: 2mM of each dNTP, 2.2mM
25 MgCl₂, 50 mM KCl, 10mM Tris•HCL, pH 9.0, and 0.1% Triton X-100, at a final fragment concentration of 10-30ng/ul. No primers are added at this point. *Taq* DNA polymerase (Promega) would be used at 2.5 units per 100ul of reaction mixture. A PCR program of 94 C for 60s; 94 C for 30s, 50-55 C for 30s, and 72 C for 30s using 30-45 cycles, followed by 72 C for 5min using an MJ Research (Cambridge, MA)
30 PTC-150 thermocycler. After the assembly reaction is completed, a 1:40 dilution of the resulting primerless product would then be introduced into a PCR mixture (using the same buffer mixture used for the assembly reaction) containing 0.8um of each primer and subjecting this mixture to 15 cycles of PCR (using 94 C for 30s, 50 C for 30s, and 72 C for 30s). The referred primers would be primers corresponding to the
35 nucleic acid sequences of the polynucleotide(s) utilized in the shuffling reaction. Said primers could consist of modified nucleic acid base pairs using methods known in the

5 art and referred to else where herein, or could contain additional sequences (i.e., for adding restriction sites, mutating specific base-pairs, etc.).

The resulting shuffled, assembled, and amplified product can be purified using methods well known in the art (e.g., Qiagen PCR purification kits) and then subsequently cloned using appropriate restriction enzymes.

10 Although a number of variations of DNA shuffling have been published to date, such variations would be obvious to the skilled artisan and are encompassed by the invention. The DNA shuffling method can also be tailored to the desired level of mutagenesis using the methods described by Zhao, et al. (*Nucl Acid Res.*, 25(6):1307-1308, (1997).

15 As described above, once the randomized pool has been created, it can then be subjected to a specific screen to identify the variant possessing the desired characteristic(s). Once the variant has been identified, DNA corresponding to the variant could then be used as the DNA substrate for initiating another round of DNA shuffling. This cycle of shuffling, selecting the optimized variant of interest, and then
20 re-shuffling, can be repeated until the ultimate variant is obtained. Examples of model screens applied to identify variants created using DNA shuffling technology may be found in the following publications: J. C., Moore, et al., *J. Mol. Biol.*, 272:336-347, (1997), F.R., Cross, et al., *Mol. Cell. Biol.*, 18:2923-2931, (1998), and A. Cramer, et al., *Nat. Biotech.*, 15:436-438, (1997).

25 DNA shuffling has several advantages. First, it makes use of beneficial mutations. When combined with screening, DNA shuffling allows the discovery of the best mutational combinations and does not assume that the best combination contains all the mutations in a population. Secondly, recombination occurs simultaneously with point mutagenesis. An effect of forcing DNA polymerase to
30 synthesize full-length genes from the small fragment DNA pool is a background mutagenesis rate. In combination with a stringent selection method, enzymatic activity has been evolved up to 16000 fold increase over the wild-type form of the enzyme. In essence, the background mutagenesis yielded the genetic variability on which recombination acted to enhance the activity.

35 A third feature of recombination is that it can be used to remove deleterious mutations. As discussed above, during the process of the randomization, for every one beneficial mutation, there may be at least one or more neutral or inhibitory mutations. Such mutations can be removed by including in the assembly reaction an

5 excess of the wild-type random-size fragments, in addition to the random-size fragments of the selected mutant from the previous selection. During the next selection, some of the most active variants of the polynucleotide/polypeptide/enzyme, should have lost the inhibitory mutations.

Finally, recombination enables parallel processing. This represents a
10 significant advantage since there are likely multiple characteristics that would make a protein more desirable (e.g. solubility, activity, etc.). Since it is increasingly difficult to screen for more than one desirable trait at a time, other methods of molecular evolution tend to be inhibitory. However, using recombination, it would be possible to combine the randomized fragments of the best representative variants for the
15 various traits, and then select for multiple properties at once.

DNA shuffling can also be applied to the polynucleotides and polypeptides of the present invention to decrease their immunogenicity in a specified host. For example, a particular variant of the present invention may be created and isolated using DNA shuffling technology. Such a variant may have all of the desired
20 characteristics, though may be highly immunogenic in a host due to its novel intrinsic structure. Specifically, the desired characteristic may cause the polypeptide to have a non-native structure which could no longer be recognized as a “self” molecule, but rather as a “foreign”, and thus activate a host immune response directed against the novel variant. Such a limitation can be overcome, for example, by including a copy
25 of the gene sequence for a xenobiotic ortholog of the native protein in with the gene sequence of the novel variant gene in one or more cycles of DNA shuffling. The molar ratio of the ortholog and novel variant DNAs could be varied accordingly. Ideally, the resulting hybrid variant identified would contain at least some of the coding sequence which enabled the xenobiotic protein to evade the host immune
30 system, and additionally, the coding sequence of the original novel variant that provided the desired characteristics.

Likewise, the invention encompasses the application of DNA shuffling technology to the evolution of polynucleotides and polypeptides of the invention, wherein one or more cycles of DNA shuffling include, in addition to the gene
35 template DNA, oligonucleotides coding for known allelic sequences, optimized codon sequences, known variant sequences, known polynucleotide polymorphism sequences, known ortholog sequences, known homolog sequences, additional

5 homologous sequences, additional non-homologous sequences, sequences from another species, and any number and combination of the above.

In addition to the described methods above, there are a number of related methods that may also be applicable, or desirable in certain cases. Representative among these are the methods discussed in PCT applications WO 98/31700, and WO
10 98/32845, which are hereby incorporated by reference. Furthermore, related methods can also be applied to the polynucleotide sequences of the present invention in order to evolve invention for creating ideal variants for use in gene therapy, protein engineering, evolution of whole cells containing the variant, or in the evolution of entire enzyme pathways containing polynucleotides of the invention as described in
15 PCT applications WO 98/13485, WO 98/13487, WO 98/27230, WO 98/31837, and Cramer, A., et al., Nat. Biotech., 15:436-438, (1997), respectively.

Additional methods of applying "DNA Shuffling" technology to the polynucleotides and polypeptides of the present invention, including their proposed applications, may be found in US Patent No. 5,605,793; PCT Application No. WO
20 95/22625; PCT Application No. WO 97/20078; PCT Application No. WO 97/35966; and PCT Application No. WO 98/42832; PCT Application No. WO 00/09727 specifically provides methods for applying DNA shuffling to the identification of herbicide selective crops which could be applied to the polynucleotides and polypeptides of the present invention; additionally, PCT Application No. WO
25 00/12680 provides methods and compositions for generating, modifying, adapting, and optimizing polynucleotide sequences that confer detectable phenotypic properties on plant species; each of the above are hereby incorporated in their entirety herein for all purposes.

30 **Example 40 – Functional Determination of Proteins Using Microarray.**

Preparation of DNA Microarrays.

Clones in YTK12 yeast cells are grown on CMD-W plates at 30°C. PCR in 96 wells plates is used to amplify the inserts as follow: per clone, a 100 ul reaction mix consisting of 80ul sterile water, 10ul 10X RedTaq Buffer (Sigma), 2ul 10mM dNTP
35 Mix (Boehringer Mannheim), 2ul of a forward primer corresponding to a polynucleotide of the present invention, 100ng/ul, 2ul T7 Primer 100ng/ul, 4ul RedTaq DNA Polymerase (Sigma) and an aliquot of the yeast colony are mixed and subjected to 95°C for 2 minutes (once) then 94°C for 30 seconds, followed by 50°C

5 for 30 seconds, and 72°C for 30 seconds (35 to 40 cycles) and finally 72°C step for 7 minutes. The PCR products are purified using gel filtration cartridges (Edge BioSystems) twice to eliminate completely excess primers. All PCR products are evaluated by agarose gel electrophoresis. Using a robot (TECAN), the PCR products are transferred to 384 well plates and samples are then dried under vacuum and
10 resuspended in 20ul of 50% DMSO.

The DNA samples are arrayed onto amino silinated coated glass microscopic slides using a microarray GEN III spotter from Molecular Dynamics/Amersham Pharmacia Biotech. After the samples are air dried, UV crosslinking of the samples to the slide can be done by exposing the slides to 50 mJ pulse of light of wavelength 254nm.

15

RNA isolation and preparation of labeled mRNA by in vitro transcription of cDNA.

Total RNA is isolated from various tissue (leaf, root, flower, stem, meristems, cell cultures, treated or untreated with chemicals or submitted or not to biotic or abiotic stresses) using a RNeasy kit from Qiagen. Total RNA is converted to cDNA as described below. First
20 strand cDNA synthesis can be performed as follows:

3-5 ug of total RNA is mixed with 1 ul of T7-oligo(dT)₂₄ primer 5'-GGCCAGTGAATTGTAATACGCTCACTATAGGGAGGCGGTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT-3' (SEQ ID NO:12) and the final volume is brought to 12 ul with DEPC-H₂O. After heating the tube at 70°C for 5 minutes, 4ul of 5 X first Strand Buffer (Life technology), 1 ul
25 of 0.1 M DTT and 1 ul of 10 mM dNTP are added. After 2 minutes incubation at 37°C, 3 ul of reverse transcriptase (SSII RT, 200-unit/ul, Life Technology) is added and incubation at 37°C for 1 hour followed.

The second strand synthesis is performed as follows: to the first strand reaction mixture 89 ul of DEPC-H₂O, 3 ul of 10mM dNTP, 1 ul of 10 U/ul *E.coli* DNA ligase, 4 ul of
30 10 U/ul *E.coli* DNA polymerase, and 2 ul of 2 U/ul *E.coli* RNase H are added and incubated at 16 °C for 2 hour. Then 2 ul of 5 U/ul T4 DNA polymerase are added and incubated at 16 °C for 5 min. The reaction is stopped by one phenol/chloroform extraction and the supernatant precipitated in the presence of 2.5 volume of 100% ethanol and 1/10 volume of 5M NH₄Ac. After mixing the sample, it is centrifuged immediately at 12 000g for 5 minutes
35 at room temperature. The pellet is washed twice with 80% ethanol and air dried before being dissolved in 2 ul of RNase free water. In vitro transcription with Cy dye labeling is performed as follow:

5 per reaction, 1 ul T7 10 X reaction buffer (Ambion), 3 ul of 25 mM ATP, CTP, GTP mix, 1 ul of 20 mM UTP (Ambion), 1 ul of cDNA, 3 ul of 5 mM of Cy3 or Cy5 (Amersham), 0.5 ul of the internal control PCR product (i.e. human CCR5), 1 ul of enzyme mix (Ambion, MEGAscript™) and 1 ul of T7 RNA polymerase (USB, 100 units/ul) are combined and incubated 37 °C for 4 – 6 hours in a water bath in the dark. The template cDNA is then
10 removed by incubating at 37 °C for 15 minutes with 1 ul of RNase-free DNaseI. The RNA probe is then purified using the RNeasy mini protocol for RNA probe purification (Qiagen). Each probe is store protected from light at – 70 °C.

Microarray hybridization and data analysis.

15 The two mRNA pools to be compared are mixed and applied to a microarray slide in 36 ul of a hybridization mixture containing 50% formamide, 5X SSC, 0.1% SDS. Hybridization can take place under a glass coverslip in a humidified plastic tube with a lid, in a 42°C oven for 14-18 hours. The slides are then washed twice for 5 minutes at 42°C in 0.5X SSC, 0.1% SDS, once for 10 minutes at 42°C in 0.25X SSC, 0.1% SDS, once in 0.25X SSC,
20 0.1% SDS at 55°C for 30 minutes, before being dried under a gentle stream of N₂.

Scanning of the slides is performed with a Generation III Array Scanner (Molecular Dynamic) at a PMT of 665V for Cy3 labeled probes and a PTM of 685V for Cy5 labeled probes. Capture of the data and critical analysis (background removal, normalization, and mean values...) is done using the software ArrayVision™ (Imaging Research, Inc.). The
25 Cy5/Cy3 fluorescence ratio and the log₁₀-transformed ratio are calculated from the normalized values. The software SpotFire™ (Spotfire, Inc.) and GeneSpring™ (Silicon Genetics, Inc.) are used for data visualization and clustering analysis.

Example 41 – Method of Transforming Plants Using Vacuum Infiltration

30 Transformation of plants may serve as a vital tool in assessing the biological function of a particular polynucleotide or polypeptide. For example, a plant may be transformed with a vector capable of downregulating a particular gene via anti-sense regulation (i.e., the vector may express a transcript of the gene of interest in the anti-sense direction), or the vector may simply be capable of overexpressing a particular
35 polypeptide, for example. By observing the resulting phenotypes of the transformant, one may derive protein function using techniques known in the art and described elsewhere herein. Other uses for transformation methods are described elsewhere herein.

5 The following method of transforming plant material may be applicable to any plant species, though is particularly suited for use in Arabidopsis.

Grow Arabidopsis at 20°C, 8 hr light, 18°C 16 dark until needed for transformation. Fertilize once a week from below. Thin plants to ~1 per square inch. Use immediately upon bolting. Short days allow stronger vegetative plant growth and
10 increase seed yield.

Transfer plants to 20°C, 16 hr light, 18 °C 8 hr dark. The plants should bolt quickly, and they are ready to infiltrate when the primary inflorescences are 10-15 cm tall and the secondary inflorescences are appearing at the rosette.

In the meantime, transform your constructs into *Agrobacterium tumefaciens* strain
15 EHA105 (Koncz and Schell, 1986) (see Direct *Agrobacterium* Transformation: Freeze-Thaw Method below). When the plants are ready to transform use 1 ml of an overnight culture to inoculate a 500 ml culture of YEB medium (2 L flask) containing the appropriate antibiotic for your construct and 50 ug/mL rifampicin (C58 Agro_ and or 25 mg/mL gentamycin (pMP90). Grow cultures 2 days
20 at 28 °C, ~275 rpm. YEB media is described below.

When OD₆₀₀ is greater than 2.0, centrifuge the culture 30 min, 3500 rpm and resuspend it in 0.5-1.0 ml of infiltration medium described below.

Place resuspended culture in a container with a large bell jar, and invert pots containing plants to be infiltrated into the infiltration medium so that the entire plant
25 is covered (including the rosette, but not too much soil). Remove any large air bubbles under the plants. Draw a vacuum (~700 Hg). Close the suction, and allow the plants to sit under vacuum for 5 min. Quickly release the vacuum pressure. Briefly drain the pots.

Grow infiltrated plants as before in 20°C, 16 hr light, 18 °C 8 hr dark. Stake plants
30 as the bolts grow. When the plants are finished flowering, harvest the T₀ seeds.

Sterilize seeds and screen for transformants on the selective medium described below. Transfer dark green (resistant) plants to secondary selection plate a week after germination, then to soil after 6-10 days. Keep new transplants covered for several days.

35

Plant media:

	Vacuum Infiltration Medium	Selection Medium
For 1 L		
MS Salts	2.2 g	4.3 g
B5Vitamins, 1000X	1.0 mL	1.0 mL
40 Sucrose	50 g	10g

5	MES, 200mg/mL pH 5.7 with KOH	2.5 mL	2.5 mL
	Benzylammonipurine (BAP, 1mg/mL)	44 μ L	-
	Silweet L-77	200 μ L	-
	Phytagar	-	8 g
10	<u>Bacterial Media</u>	<u>YEP</u>	
	Yeast extract	1.0 g	
	Beef Extract	5.0 g	
	Peptone	5.0 g	
	Sucrose	5.0 g	
15	MgSO ₄	0.5 g	

The skilled artisan would appreciate that the above transformation method could be modified to apply to other species of plants. Such modification may include the addition of new steps, the deletion of any of the steps described, and/or substitution of reagents.

Direct *Agrobacterium* Transformation: Freeze-Thaw Method.

Grow an *Agrobacterium* strain containing the appropriate helper Ti plasmid in 5 mL of YEP medium (YEP medium is described elsewhere herein) overnight at 28 °C. Add 2 ml of the overnight culture to 50 ml YEP medium in a 259 mL flask and shake vigorously (250 rpm) at 28 °C until the culture grows to an OD₆₀₀ of 0.5 to 1.0. Chill the culture on ice. Centrifuge the cell suspension at 3000 g for 5 min at 4 °C.

Discard the supernatant solution. Resuspend the cells in 1 mL of 20 mM CaCl₂ solution (ice-cold). Dispense 0.1 mL aliquots into prechilled Eppendorf tubes. Add about 1 ug of plasmid DNA to the cells.

Freeze the cells in liquid nitrogen Thaw the cells by incubating the test tube in a 37 °C water bath for 5 min. Add 1 mL of YEP medium to the tube and incubate at 28 °C for 2-4 hr with gentle rocking. This period allows the bacteria to express the antibiotic resistance genes. Centrifuge the tubes for 30 s in an Eppendorf centrifuge. Discard the supernatant solution. Resuspend the cells in 0.1 mL YEP medium

Spread the cells onto a YEP agar plate containing 3-5 ug/mL tetracycline and 10-25 ug/mL kanamycin. Incubate the plate at 28 °C. Transformed colonies should appear in 2-3 days.

The skilled artisan would appreciate that the above transformation method could be modified to apply to other species of plants. Such modification may include the addition of new steps, the deletion of any of the steps described, and/or substitution of reagents.

5 **Example 42– Functional Determination of Proteins Using Metabolite Profiling.**

The present invention encompasses the application of metabolite profiling to the identification of gene function for the polypeptides of the present invention. In one example, transgenic plants could be produced which are either incapable of expressing a protein of the present invention, or that have decreased expression levels
10 of a protein of the present invention. Such transgenic plants could be produced by creating knockout constructs to inactivate or delete the endogenous gene, for example, using methods known in the art and described elsewhere herein. Alternatively, the transgenic plants could be produced by inserting into the plant a construct that expresses antagonists of a protein of the present invention (e.g., antisense
15 oligonucleotides, antisense genes, antibodies, etc.). Other examples of methods of producing transgenic plants, including specific strategies, are known in the art, some of which are described elsewhere herein.

Once a protein of the present invention is inactivated, or its expression inhibited, the resulting metabolite profile of the plant can be ascertained, and the
20 function of the protein assigned. Some of the anticipated metabolic profiles of inhibiting or inactivating the expression of a protein of the present invention in a plant may resemble known nutritional deficiencies, pathogenic diseases, biotic stresses, or abiotic stresses, for example, many of which are disclosed elsewhere herein. In addition, the metabolic profile of a transgenic plant of the present invention may be
25 useful in identifying the specific pathways the polypeptide of the present invention is a member of, in addition, to identifying the potential downstream and/or upstream effectors or effectors, respectively. In addition, it may be possible to identify the mode of action of a polypeptide of the present invention.

A number of methods are known in the art for identifying the metabolic
30 profile of a plant. A non-limiting example is provided by Sauter, H., et al., in “Metabolic Profiling of Plants: A New Diagnostic Technique”, Synthesis and Chemistry of Agrochemicals II, Baker, D.R., Fenyves, J.G., and Moberg, W.K., eds, ACS Symposium Series, 433, Chapter 24, pp. 288-299, (1991). Briefly, transgenic plants of the invention, or plants in which the expression of a polypeptide of the
35 present invention is inhibited or inactivated, are grown in growth chambers. The shoots are harvested and immediately deep frozen until further treatment. The frozen plant samples are weighed and a threefold amount (W:W) of ethanol is added. The mixture is then macerated in a mixer and the resulting suspension is left for 2 hours

5 for extraction. The next steps are filtration, evaporation, and silylation with N-Methyl-N-(trimethylsilyl)trifluoroacetamide (MSTFA). Internal standard alkanes are also added, thus allowing for the calculation of retention coefficients, as well as, quantification. The crude mixture is then subjected to gas chromatography on a methyl silicon gum fused silica capillary column (30 m DB-1. Injection temperature
10 230°C. Oven temperature 100°-320°C, 4°C/min; 15 min 320°C). Retention coefficients are then calculated relative to internal standards (n-C₁₀H₂₂=1000, n-C₂₈H₅₈=2800).

The above protocol can be applied to numerous test plants, in addition, to controls. The data from the resulting profiles are then grouped together (i.e., one
15 group for the test plants, another group for the controls) to arrive at an average profile for each group. In the latter step, the corresponding peaks (i.e., those peaks with equal retention coefficients) are grouped together and the peak heights are subjected to statistical analysis.

The differences in metabolic profiles between the test and control plants are
20 determined by calculating the “difference profile” between the two groups. The difference profile is calculated by dividing the peak heights. This difference profile provides a semiquantitative estimate of the change in magnitude of one metabolic with respect to the other.

Once the above is completed, the peaks are then associated with particular
25 metabolites (i.e., the metabolite identify of each peak is determined). By comparing the metabolite profile of proteins known to modulate specific pathways in a plant, to those of the present invention, clues for and/or identification of the function of a polypeptide of the present invention may be determined. Other methods are known in the art, and any one or more steps, may be equally substituted with such methods.

30

Example 43 – Functional Determination of Proteins Using Morphological Phenotyping.

The present invention encompasses the application of morphological
phenotyping to the identification of gene function for the polypeptides of the present
35 invention. In one example, transgenic plants could be produced which are either incapable of expressing a protein of the present invention, or that have decreased expression levels of a protein of the present invention. Such transgenic plants could

5 be produced by creating knockout constructs to inactivate or delete the endogenous gene, for example, using methods known in the art and described elsewhere herein. Alternatively, the transgenic plants could be produced by inserting into the plant a construct that expresses antagonists of a protein of the present invention (e.g., antisense oligonucleotides, antisense genes, antibodies, etc.). Other examples of
10 methods of producing transgenic plants, including specific strategies, are known in the art, some of which are described elsewhere herein.

Once a protein of the present invention is inactivated, or its expression inhibited, the resulting morphological phenotype of the plant can be ascertained, and the function of the protein assigned. Some of the anticipated phenotypes of inhibiting
15 or inactivating the expression of a protein of the present invention in a plant may resemble known nutritional deficiencies, pathogenic diseases, biotic stresses, or abiotic stresses, for example, many of which are disclosed elsewhere herein.

**Example 44 – Functional Determination of Proteins Using TransGenomic
20 Technology (TGT).**

The present invention encompasses the application of TransGenomic Technology (TGT) to the identification of gene function for the polypeptides of the present invention (Cambia Biosystems, Bellevue, WA). TGT combines both genomics and functional genomics, enabling the association of gene sequences (e.g.,
25 the polynucleotide sequences of the present invention) with specific phenotypes or traits. The technology is based upon the random integration of reporter gene [e.g. beta-Glucuronidase (GUS), BoGUS, GFP] transposons throughout a plant's genome (specifically rice and Arabidopsis).

The transposon may comprise an enhancer trap vector containing a
30 transcriptional activator and a reporter gene capable of responding to the transactivator. Since the integration is random, it is expected that the transposon may integrate near a gene regulatory element - effectively turning on the transactivator gene, the activity of said gene promoting activation of the reporter gene. Thus, by repeating the process above, a large population of cell lines with unique expression
35 patterns can be generated. These "pattern lines" are genetically characterized in an effort to select lines with desired expression patterns. The pattern lines can be used for reverse genetics studies to identify lines with mutations in genes of interest and may also be used as a genetic background to obtain gain-of-function (GOF) and loss-of-

5 function (LOF) mutations. Moreover, novel expression (or lack of expression) patterns may uncover novel phenotypes and traits. In addition, the pattern lines showing desirable expression patterns could serve as a genetic foundation to direct the expression of genes in target tissues or at desired developmental stages.

The TGT technology described above can be used to identify the function of
10 the polynucleotides and/or polypeptides of the present invention. For example, the polynucleotide sequences of the present invention can be used to generate PCR primers suitable for the screening and identification of cell lines positively tagged with the TGT technology upstream of a polynucleotide of the present invention. Once identified, the function of the polynucleotide sequence may be ascertained by linking
15 the genes of the present invention tagged with the TGT technology with specific phenotypes or traits observed in a particular cell line, or pattern line. Another possible application of TGT to the polynucleotides of the present invention would be in targeting gene expression of selected polynucleotides of the present invention. For example, polynucleotide sequences of the present invention flanked by an upstream
20 regulatory sequence may be introduced into pattern lines showing desired expression patterns using techniques known in the art of molecular biology and described elsewhere herein.

A number of variations of the above technology are known in the art and would be appreciated by the skilled artisan. For example, the skilled artisan would
25 appreciate that other transposons, reporter genes, and activating sequences, etc., could be substituted or used in conjunction with the TGT technology. Specific methods of applying TGT technology can be found, for example, in Wilson KJ, et al., *Microbiol*, 141:1691-1705, (1995), and in US Patent Nos. 5,879,906, 5,599,670, and 5,432,081; which are hereby incorporated by reference in their entirety herein.

30

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CLAIMS

WHAT IS CLAIMED IS:

1. An isolated Na^+/H^+ antiporter coding nucleic acid, wherein the nucleic acid
10 comprises a polynucleotide selected from the group consisting of:
 - a) a polynucleotide of SEQ ID NO:1;
 - b) a polynucleotide of SEQ ID NO:4;
 - c) a polynucleotide encoding a polypeptide of SEQ ID NO:2;
 - d) a polynucleotide encoding a polypeptide of SEQ ID NO:5;
 - 15 e) a polynucleotide comprising at least 60 consecutive nucleotides of any of a) through d) above;
 - f) a polynucleotide having at least 70% sequence identity with the nucleotide sequence of SEQ ID NO:1 or SEQ ID NO:4;
 - g) a polynucleotide encoding a polypeptide having at least 70% sequence identity
20 with the polypeptide of SEQ ID NO:2 or SEQ ID NO:5; and
 - h) a polynucleotide complementary to a polynucleotide of any of a) through g).
2. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid comprises a polynucleotide encoding the polypeptide of SEQ ID NO:2.
25
3. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid comprises a polynucleotide encoding the polypeptide of SEQ ID NO:5.
4. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid comprises a polynucleotide encoding a polypeptide having at least 90% sequence
30 identity to the polypeptide of SEQ ID NO:2 or SEQ ID NO:5.
5. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid comprises at least 60 consecutive nucleotides of any of the polynucleotides of a) through d) of Claim 1.
35
6. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid comprises a polynucleotide having at least 90% sequence identity with the nucleotide sequence of SEQ ID NO:1 or SEQ ID NO:4.
40
7. A vector comprising a nucleic acid of Claim 1.

- 5 8. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid encodes a polypeptide that functions as a modulator of an environmental stress response in a plant.
9. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein overexpression
10 of the nucleic acid in a plant increases the plant's tolerance to an environmental stress.
10. The Na^+/H^+ antiporter coding nucleic acid of Claim 9, wherein the environmental stress is salt.
- 15 11. The Na^+/H^+ antiporter coding nucleic acid of Claim 1, wherein the nucleic acid encodes a polypeptide that functions in a sodium transport system.
12. A first isolated nucleic acid that hybridizes under stringent conditions to a second nucleic acid selected from the group consisting of:
- 20 a) a second nucleic acid comprising a polynucleotide of SEQ ID NO:1 or SEQ ID NO:4; and
- b) a second nucleic acid encoding a polypeptide of SEQ ID NO:2 or SEQ ID NO:5, wherein the first nucleic acid encodes a polypeptide that functions as a modulator of an environmental stress response in a plant.
- 25 13. A transgenic plant cell comprising a Na^+/H^+ antiporter coding nucleic acid of Claim 1.
14. A transgenic plant comprising a plant cell of Claim 13.
- 30 15. The plant of Claim 14, wherein the plant is a monocot.
16. The plant of Claim 14, wherein the plant is a dicot.
- 35 17. The plant of Claim 14, wherein the plant is selected from the group consisting of maize, wheat, rye, oat, triticale, rice, barley, soybean, peanut, cotton, rapeseed, canola, manihot, pepper, sunflower, tagetes, solanaceous plants, potato, tobacco, eggplant, tomato, Vicia species, pea, alfalfa, coffee, cacao, tea, Salix species, oil palm, coconut, perennial grass and a forage crop.
- 40 18. A plant seed produced by the plant of Claim 14.

5 19. The seed of Claim 18, wherein the seed is true breeding for an increased tolerance to environmental stress as compared to a wild type variety of the seed.

20. A method of modulating a plant's tolerance to an environmental stress comprising, modifying the expression of a Na^+/H^+ antiporter coding nucleic acid in
10 the plant, wherein the nucleic acid is selected from the group consisting of:

- a) a polynucleotide of SEQ ID NO:1;
- b) a polynucleotide of SEQ ID NO:4;
- c) a polynucleotide encoding a polypeptide of SEQ ID NO:2;
- d) a polynucleotide encoding a polypeptide of SEQ ID NO:5;
- 15 e) a polynucleotide comprising at least 60 consecutive nucleotides of any of a) through d) above;
- f) a polynucleotide having at least 70% sequence identity with the nucleotide sequence of SEQ ID NO:1 or SEQ ID NO:4;
- g) a polynucleotide encoding a polypeptide having at least 70% sequence identity
20 with the polypeptide of SEQ ID NO:2 or SEQ ID NO:5; and
- h) a polynucleotide complementary to a polynucleotide of any of a) through g).

21. The method of Claim 20, wherein the environmental stress is selected from the group consisting of high salinity, drought and low temperature.

25

22. The method of Claim 20, wherein the nucleic acid comprises a polynucleotide encoding the polypeptide of SEQ ID NO:2.

23. The method of Claim 20, wherein the nucleic acid comprises a polynucleotide
30 encoding the polypeptide of SEQ ID NO:5.

24. The method of Claim 20, wherein the nucleic acid comprises a polynucleotide encoding a polypeptide having at least 90% sequence identity to the polypeptide of SEQ ID NO:2 or SEQ ID NO:5.

35

25. The method of Claim 20, wherein the nucleic acid comprises at least 60 consecutive nucleotides of any of the polynucleotides of a) through d).

26. The method of Claim 20, wherein the nucleic acid comprises a polynucleotide
40 having at least 90% sequence identity with the nucleotide sequence of SEQ ID NO:1 or SEQ ID NO:4.

- 5 27. The method of Claim 20, wherein the plant's stress tolerance is increased by increasing expression of the nucleic acid in the plant.
28. The method of Claim 20, wherein the nucleic acid encodes a polypeptide that functions in sodium transport system.
- 10 29. The method of Claim 20, wherein the plant is not transgenic.
30. The method of Claim 20, wherein the plant is transgenic.
- 15 31. The method of Claim 30, wherein the plant is transformed with a promoter that directs expression of the nucleic acid.
32. The method of Claim 31, wherein the promoter is tissue specific.
- 20 33. The method of Claim 31, wherein the promoter is developmentally regulated.
34. A method of producing a transgenic plant containing a Na^+/H^+ antiporter coding nucleic acid wherein the plant has an increased tolerance to an environmental stress as compared to a wild type variety of the plant comprising, transforming a plant
- 25 cell with an expression vector comprising the nucleic acid and generating from the plant cell the transgenic plant, wherein the nucleic acid is selected from the group consisting of:
- a) a polynucleotide of SEQ ID NO:1;
- b) a polynucleotide of SEQ ID NO:4;
- 30 c) a polynucleotide encoding a polypeptide of SEQ ID NO:2;
- d) a polynucleotide encoding a polypeptide of SEQ ID NO:5;
- e) a polynucleotide comprising at least 60 consecutive nucleotides of any of a) through d) above;
- f) a polynucleotide having at least 70% sequence identity with the nucleotide
- 35 sequence of SEQ ID NO:1 or SEQ ID NO:4;
- g) a polynucleotide encoding a polypeptide having at least 70% sequence identity with the polypeptide of SEQ ID NO:2 or SEQ ID NO:5; and
- h) a polynucleotide complementary to a polynucleotide of any of a) through g).
- 40 35. The method of Claim 34, wherein the environmental stress is selected from the group consisting of high salinity, drought and low temperature.

- 5 36. The method of Claim 34, wherein the plant is a monocot.
37. The method of Claim 34, wherein the plant is a dicot.
38. The method of Claim 34, wherein the plant is selected from the group
10 consisting of maize, wheat, rye, oat, triticale, rice, barley, soybean, peanut, cotton, rapeseed, canola, manihot, pepper, sunflower, tagetes, solanaceous plants, potato, tobacco, eggplant, tomato, Vicia species, pea, alfalfa, coffee, cacao, tea, Salix species, oil palm, coconut, perennial grass and a forage crop.
- 15 39. The method of Claim 38, wherein the nucleic acid comprises a polynucleotide encoding the polypeptide of SEQ ID NO:2.
40. The method of Claim 38, wherein the nucleic acid comprises a polynucleotide encoding the polypeptide of SEQ ID NO:5.
- 20 41. The method of Claim 38, wherein the nucleic acid comprises a polynucleotide encoding a polypeptide having at least 90% sequence identity to the polypeptide of SEQ ID NO:2 or SEQ ID NO:5.
- 25 42. The method of Claim 38, wherein the nucleic acid comprises at least 60 consecutive nucleotides of any of the polynucleotides of a) through d).
43. The method of Claim 38, wherein the nucleic acid comprises a polynucleotide having at least 90% sequence identity with the nucleotide sequence of SEQ ID NO:1
30 or SEQ ID NO:4.
44. The method of Claim 38, wherein the plant's stress tolerance is increased by increasing expression of the nucleic acid in the plant.
- 35 45. The method of Claim 38, wherein the nucleic acid encodes a polypeptide that functions in a sodium transport system.
46. The method of Claim 38, wherein the plant is transformed with a promoter that directs expression of the nucleic acid.
- 40 47. The method of Claim 46, wherein the promoter is tissue specific.
48. The method of Claim 46, wherein the promoter is developmentally regulated.

Figure 1

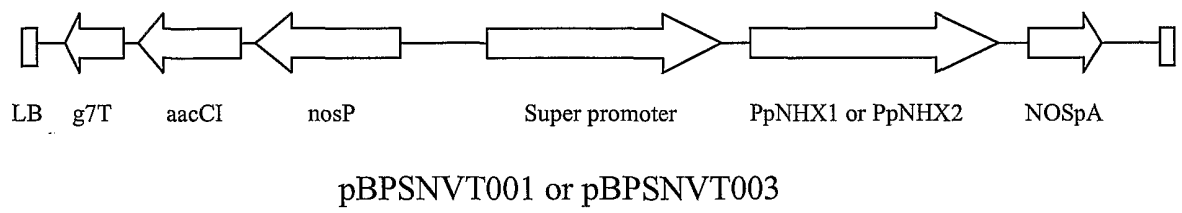


Figure 2A

Nucleotide sequence of full length PpNHX1 SEQ ID NO: 1

1 ATGGCGACCA ATGATGTCGT GAGCGTTTCG CATTCTATGC TACTAAAAGC
51 TACAGATCTC AAAGATGACC GAATCGATGT AATTTCAATCTGTCTCTTTG
101 TATTTTTACT CTGTGCGTGC ATTGTGCTGG GGCACCTTCTGGAGGAAAAT
151 CGGTGGATGA ATGAGTCTAT TACTGCTCTT CTTCTGGGAC TCTTTACTGG
201 ATCTATAGTG TTGATTTCAA GCAAAGGTCA AGGTTCTCAT ATTCTGGAGT
251 TTGATGAAGA GCTTTTCTTC ATATACCTTC TTCCACCTAT AATCTTCAAT
301 GCTGGGTTCC AGGTTAAGAA GAAGGAATTC TTTCGGAATT TCATAACAAT
351 CATGTTTTTT GGAGTTATAG GAGTCTTTAT TTCTTTCGGA ATTATCTCAA
401 CAGGAAGTTG GTATTTCTTC TCCAAGTTCG GACTTAAGAA CCTGCCTATT
451 CGAGATATCC TAGCTATTGG AGTCATCTTT TCTGCTACCG ATTCCGTCTG
501 CACGTTGCAG GTGCTGAACC AAGATGAAAC CCCTCTACTT TACAGTTTGG
551 TCTTTGGGGA AGGAGTCGTA AATGATGCTA CTTCTGTGGT TCTGTCTCGA
601 GCTGTTCAAA CATACAACTT TGACAATTTT ACATCCTTAG AAGGCTTACA
651 AATTGGAGGC AGTTTCTTGT ACTTATTCTT CTCGAGTTGC ATCCTGGGAA
701 TCGCGTCGGG CTTAATAAGC GCATATATCA TCAAGACAAT GTACTTTGGC
751 AGGCATTCCA CGGATCGTGA AATAGCAATC ATGACATTGA TGGCGTATTT
801 ATCTTACGTC TTTGCAGAGC TTTTCTACTT GAGTGGAATT CTCTCAGTGT
851 TCTTTTGCGG CATTGTAATG TCTCATTACA CTTGGCATAA CGTCACGGAG
901 AATTCTCGAA TCACAAGCAA GCATTCCTTT GCAACGATGT CATTCAATTGC
951 AGAGACGTTT ATATTTCTAT ATGTTGGAAT GGATGCTCTG GATTTCGAAA
1001 AATGGAAGAT GATGCAATCC AGTTTCACGG AATCTGCGGG CTATTTGGT
1051 AGCTTGTTGT TTCTGGTCCT GTTAGGGAGG GCCGCATTTG TGTTCCCACT
1101 CTCTGCTTTG TCCA ACTACA GCACAAAGTC TCCAGACGCG AAGATTAATT
1151 TACGCCAAAT GGTTATTATC TGGTGGGCTG GACTAATGCG AGGTGCTGTC

Figure 2B

1201 TCAATAGCAC TGGCGTTCAA CCAGGGTGGT GATGCAAAGG CTCAAACCA
 1251 AGCCACGCTA ATGGTCATTA CTATCATCAT TGTCTCTTC AGCACTATTG
 1301 TGTTTCGGCAC TGCAACCAAG CCTCTTATTA GCTGGCTACT TCCACCTCAT
 1351 TTCAGATCAA ATTACAGTGA TTCAGCCAGT CTCTCCCCAA AAGCGTCTCT
 1401 TGATGCTGAC TTTCATATAC CACTCCTTAT GGATACAGAG CGTGAAGAAT
 1451 TAGAAGCAAA TGATCGATCT ACGATAAATC AAATCCTAAA TGGTCTTCTT
 1501 TGTCTCAGT CAATAGGCAT GCTGCTGACT GCACCAAGAT CAACCATCCA
 1551 CCATGTATGG AGAAAATTTG ATGATTCTTA CATGCGGCC ACCTTTGGTG
 1601 GGAGAGGATA TGTTAGGTTG GTGTCACGGC GTGATATGGA ATACAAGAA
 1651 GATGAAATCC TTGAAGATCA CAGTTGA

Nucleotid sequence of partial PpNHX2 SEQ ID NO: 3

CGGCACGAGAACCTTTGCAGCTGCTAAGGCATGCAGGGTGGATGCCCGCA
 AAGCTTTAACTCTCGGCATCCTGATGAATACCAAAGGATTGGTGGAGCTT
 ATTGTTCTGAACATCGGTTTAGATCGTGGAGTTCTGAATTCGGAGACTTT
 TGCAATCATGGTGTGATGGCTCTCTCACAACGTTTCATGACAAACACCTC
 TGTAATGGCTATATATAAACCAGCCACGAATCCCACTCCTTACACTCGT
 AGGACTTTGGAAATGGAGGACTCGAAGGATGACTTGCGAATATTGTCATG
 CGTGCACGGAATGAAGAACGTGGCTGCCATGATCAATCTTACAGAAGCGA
 CCAGGGGCATGCGCAAACGTA CTCTGCGCCTGTATATTTGCATTTGATG
 GAACTATCCGAACGTA CTCTGCCATTATGATTGTCCAGCGGGCACGTCG
 GAATGGGCGCCCTTTTTTCAATCAGAGCAAACATTTCGGACAACAAAGATC
 AAATTGTTGCGGCCCTTCGAGACATATGAACA ACTAAGCAAGGTGACTGTG
 AGGCCTATGACTGCAATTTCCGGGTTTCGACGACATGCACGAAGACATATG
 TCGACTGCTGCTGACAAGCGGACTGCCTTGATCATGCTTCTTTCCACA
 AATCACCCAAACTGGACGGGCACTTCGATTCTACTCCAGGTTTNCGAACA
 GTTAATCAAAAGTCTCAAGCATGCACCGTCTGTTGCTATTCTAAT
 CGATCGTGGAGTCGGTGGATCAACCCAAGTGNCTT

Nucleotide sequence of full length PpNHX2 SEQ ID NO: 4

TGGCACCAGCAAGATGGCGGACGCTGTGGCGTGCAAACTATGTCCGGCTACATCCAATGG
 AGTGTGGCAGGAGATGTGCCCGTTCAATTTGCTCTTCTCTGCTCATCGTTCAAATGTCC
 TCGTTTTGGCAATCACTCGGGCGTTAGCTTTTGTCTGAAGCCTTTGAAACAGCCCCGCGT

Figure 2C

CGTCGCCGAGATTATAGGCCGAATATTGCTTGGTCCATCTGCTTTTGGACGCAATAAGGAC
TACCTGCATACGATTTTCCACATGAAAGTGTTATCATTCTGGAGGTCTTTGCAGACATGG
GACTTTTATTCTTTTTGTTTCATGGTGGGGTTAGAGCTCGATATGACCCAGATTCGGAAAAC
CGGAAAGCAAGCTATGTCCATTGCTGCAGCTGGAATCACTCTGCCTTTCGTTGCAGGTGTC
GGTGTTCCTTCGTCTGCATCTTACAATTGCACCAGAGGGAGCTTTTGGTCCGTTTCTCGT
GTTTCATGGGAGTTGCTATGTCCATCACTGCTTTCCTGTTCTGGCACGTATTTTGGCGGAG
AGGAAGCTTTTACTACCGAAGTAGGGCAATTGGCGATGTCAGCAGCTGCAGTTAATGAC
GTGGTTGCTTGGGTTCTTTTACGFTGGCGGTGCGCTTGTGGGGCTCCGGAAGGAGCCAG
CAATTGTTGCATGGGTTCTGTTGTGTGGAATCGCATTTTGTCTGGCCATCTTCTTGTGGTT
CAACCATGCATGCAATGGGTTGCTCATCGATCGCCGACAATGAGCCTGTCAAAGAATAC
ATTGTAGCATTGACTTTACTTTGTGTTCTCGTTGCTGGATTCTGTACTGATGCGATAGGAGT
TCATTCCATTTTTGGCGCGTTTCTGTTTGGACTTGTATACCTAAAGAGGGTCTTTTCGCAG
CGGCTTTGGTTGAGAAATTAGAAGATTTTGTATCTATCCTCTTGTGCCTCTCTACTTTGCA
TCGAGTGGACTGAAGACCAACATTGGAGCTATTCACAGCGCGCAATCTTTTGGCCTTTTGG
TCTTGGTTATCAGCGTTGCTTGTCTGGGTAATAATTCTCGAACCTTTGCAGCTGCTAAGGC
ATGCAGGGTGGATGCCCGCAAAGCTTTAACTCTCGGCATCCTGATGAATACCAAAGGATT
GGTGGAGCTTATTGTTCTGAACATCGGTTTAGATCGTGGAGTTCTGAATTCGGAGACTTTT
GCAATCATGGTCTGATGGCTCTCTCACAACGTTTCATGACAACACCTCTGGTAATGGCTA
TATATAAACCAGCCAGGAATCCCACTCCTTACACTCGTAGGACTTTGGAAATGGAGGACT
CGAAGGATGACTTGCGAATATTGTCATGCGTGCACGGAATGAAGAACGTGGCTGCCATGA
TCAATCTTACAGAAGCGACCAGGGGCATGCGCAAACGTACTCTGCGCCTGTATATTTTGGCA
TTTGATGGAACCTATCCGAACGTACTTCTGCCATTATGATTGTCCAGCGGGCACGTCGGAAT
GGGCGCCCTTTTTTCAATCAGAGCAAACATTCGGACAACAAAGATCAAATTGTTGCGGCC
TTCGAGACATATGAACAATAAGCAAGGTGACTGTGAGGCCTATGACTGCAATTTCCGGG
TTCGACGACATGCACGAAGACATATGTGCGACTGCTGCTGACAAGCGGACTGCCTTGATC
ATGCTTCCTTTCCACAAATCACCCAGACTGGACGGGCACTTCGATTCTACTCCAGGTTTCC
GAACAGTTAATCAGAAGGTCCTCAAGCATGCACCGTCTGCTGCTATTCTAATCGATCG
TGGAGTCGGTGGATCAGCCCAAGTGCTTCCAGCAACGTTGATCACAATGTTGTCGTGTAC
TTCTTTGGTGGTCTGACGACAGGGAAGCTCTGGCATATGGTTTCCGTATGGCTGAGCATC
CGGAGTTAAGCTTCATGTTATCCGTTTCTTCTCACAGCGTCGTCATGGACGACGGCCA
TGGAGGATTAGCTTCCGTCGGATCAGAGGTATCTGAGATTGGCAAGACGGAGGTGAGCGA
TACTCGTTTCCAGTTCGCGATGCATGGTCTGGACCAAAAACAGGCAAAGGGAGTTGGATGA
AGAAGCCTTGGCCATGTGCGTAGGAGGCAAGCTTCTGAAGATGGAAGAGTCAACATACGT
AGAAATGCAGGTATCTGAGCCTTGAAGAGGTGGTGAAGATTGAGTAGCTCTCGTGAACA
CGATATTATTTGGTTGGCCGATCGAGAAGGCCAACGCCATTTTGTAGAGCGATTCCGTCGT
AAGCACGCAGAATATGCAGAGCTTGGCCCTATTGGAGATGCTCTGATGGCCCCACAGGTA
CGAGCATCTGTCTTAGTATTTAGCAGCAGCATCATGTGCTTGGCGATCCACTTCTAATA
CCTCTGAAACGGAGGCCGTCAAAGAGTTGCAGACCTTCCCATCATCCAAGGAATTGGTGG

Figure 2D

ATCGTAAAGGTGATGTACAGAAGATCGACTTGTCTTCTCCTGACCACCGCGTGTATATGAC
CCGGGTCTAGAGCGGCCGCCACCGCGGTGGAGCTCCAGCTTTTGTCCCTTTAGTGAGGGT
TAATTGCGCG

Figure 3

Amino acid sequence PpNHX1 SEQ ID NO: 2

1 MATNDVVSVS HSMLLKATDL KDDRIDVISI CLFVFLLCAC IVLGHLLLEN
 51 RWMNESITAL LLGLFTGSIV LISSKGQGS ILEFDEELFF IYLLPPIIFN
 101 AGFQVKKKEF FRNFITIMFF GVIGVFISFG IISTGSWYFF SKFGLKNLPI
 151 RDILAIGVIF SATDSVCTLQ VLNQDETPLL YSLVFGEGVVNDATSVVLSR
 201 AVQTYNFDNF TSLEGLQIGG SFLYLFFSSC ILGIASGLIS AYIIKTMYFG
 251 RHSTDREIAI MTLMAYLSYV FABLEFYLSGI LSVFFCGIVMSHYTWHNVTE
 301 NSRITSKHSF ATMSFIAETF IFLYVGMDAL DFEKWKMMQSFTESAGLFG
 351 SLLFLVLLGR AAFVFPLSAL SNYSTKSPDA KINLRQMVIWVWAGLMRGAV
 401 SIALAFNQQG DAKDSNQATL MVITIII VLF STIVFGTATK PLISWLLPPH
 451 FRSNYSDSAS LSPKASLDAD FHIPLMDTE REELEANDRS TINQILNGLP
 501 CPQSIGMLLT APRSTIHHVWKFDDSYMRPTFGGRGYVRLVSRRDMDIQE
 551 DEILEDHS*

Amino acid sequence PpNHX2 SEQ ID NO: 5

MADAVACKTMSATSNGVWQGDVPVHFALPLLIVQIVLVLAITRALAFVLKPLKQPRVVAEIIIG
 GILLGPSAFGRNKDYLHTIPPHESVILEVFADMGLLFFLFMVGLELDMTQIRKTGKQAMSIAA
 AGITLPFVAGVGVSVFLHLTIAPEGAFGPFLVFMGVAMSITAFPVLARILAEKLLTTEVGQLA
 MSAAAVNDVVAWVLLALAVALS GSGRSPAIVA WVLLCGIAFCLAIFLVVQPCMQWVAHRSP
 DNEPVKEYIVALTLLCVLVAGFCTDAIGVHSIFGAFLFGLVIPKEGPFAAALVEKLEDFVSILL
 PLYFASSGLKTNIGAIHSAQSFGLLVLVISVACL GKILGTFAAAKACRVDARKALTLGILMNTK
 GLVELIVLNIGLDRGVLNSETFAIMVLMALFTTFMTTPLVMAIYKPARNPTPYTRRTLEMEDSK
 DDLRILSCVHGMKNVAAMINL TEATRGMRKRTLRLYLHLMELSER TSAIMIVQRARRNGRPF
 FNQSKHSDNKDQIVAAFETYEQLSKVTVRPMTAISGFDDMHEDICATAADKRTALIMLPFHKS
 PRLDGHFDSTPGFRTVNQVKVLKHAPCSVAILDRGVGGS AQVPSSNVVDHNVVVYFFGGPDDRE
 ALAYGFRMAEHPGVKLVIRFLSHSVVMDDGHGGLASVGSEVSEIGKTEVSDTRFQFAMHGL
 DQNRQRELDEEALGHVRRRQASEDGRVTVYVEMQVSEPLEEVRLSSREHDILVGRSRRPTP
 FLERFRRKHAEY AELGPIGDALMAPQVRASVLV FQQHDHVLADPLNTSETEAVKELQTFPSS
 KELVDRKGDVQKIDLSSPDHRVYMTRV

Figure 4A

	1	40
O. sativa @9SXJB	(1)	-----
Arabidopsis AtNhx1	(1)	-----
SEQ ID #2	(1)	-----
D.melanogaster @9VPL	(1)	-----MRVWVAYS
C. elegans @9X14	(1)	MMLSVEEQVLRNNIELMFTFNSLKILANGVTWQLWESTLN
	41	80
O. sativa @9SXJB	(1)	-----
Arabidopsis AtNhx1	(1)	-----
SEQ ID #2	(1)	-----
D.melanogaster @9VPL	(10)	AALLLVHAGPESISGQEVPEQSKTSSNTTTTDSSSSIHTV
C. elegans @9X14	(41)	QGTATSGIMRFALKTALSICIFLLIFQTVDSQSSSSASA
	81	120
O. sativa @9SXJB	(1)	-----MGMEVAAARL
Arabidopsis AtNhx1	(1)	-----MLDSLVS
SEQ ID #2	(1)	-----MATNDVVSVSHSMLL
D.melanogaster @9VPL	(50)	SDVFNVSPLGNVTPSISASGNASTTKRGNASTLVTDPPLI
C. elegans @9X14	(81)	SVVSGAVKSEDTVVAVNKTDVLEGEAIDANATSLEQHGAAI
	121	160
O. sativa @9SXJB	(11)	GALYTTSDYASVVSINLFFVALLCACIVLGHLLLEN--RWV
Arabidopsis AtNhx1	(9)	LPSLSTSDHASVVALNLFVALLCACIVLGHLLLEN--RWM
SEQ ID #2	(16)	KATDLKDDRIDVISICLFFVLLCACIVLGHLLLEN--RWM
D.melanogaster @9VPL	(90)	DSHAVEQEHNSLSLFFVICVIMLGILLIHSMLQTFQYL
C. elegans @9X14	(121)	VGNVSE-EKKRSLAIFFFILFVIMLATLVVHMLIVSKIHW
	161	200
O. sativa @9SXJB	(49)	NESITALIIGLCTGVVILLMT--KGKSSHLLVFSEDLFFI
Arabidopsis AtNhx1	(47)	NESITALLIGLGTGVTILLIS--KGKSHLLVFSEDLFFI
SEQ ID #2	(54)	NESITALLGLFTGSIVLISS--KGQGSIHLEFDEELFFI
D.melanogaster @9VPL	(130)	PESIVVVFLGAFIGLSLNVMSGQNGSWKREEVFSMPGFFL
C. elegans @9X14	(160)	PESLAIVALGALIGSILSYSR--R-DWSEIEALSPDVFFL
	201	240
O. sativa @9SXJB	(87)	YLLPPIIFNAGFQVKKKQFFRNFMITLFGAVGTMISFFT
Arabidopsis AtNhx1	(85)	YLLPPIIFNAGFQVKKKQFFRNFMITLFGAVGTIISCTI
SEQ ID #2	(92)	YLLPPIIFNAGFQVKKKQFFRNFMITMFFGVIGVFISFGI
D.melanogaster @9VPL	(170)	VLLPPIIFESGYNLHKGFFQNIQSILVFAIFGTTISALV
C. elegans @9X14	(197)	VLLPPIIFENAYNLNKGFFSNFVPILTFAIFGTTISAMV
	241	280
O. sativa @9SXJB	(127)	ISIAAIAIFSRMNIPTLDVGDFLAIGAIIFSATDSVCTLQV
Arabidopsis AtNhx1	(125)	ISLGVTQFFKLDIGTFDLGDYLAIGAIFAATDSVCTLQV
SEQ ID #2	(132)	ISTGSWYFFSKFGLKNLPIRDLAIGVIFSATDSVCTLQV
D.melanogaster @9VPL	(210)	IGAGIYLLGLGEVAFRLSFSSEFAFGSLISAVDPVATVAI
C. elegans @9X14	(237)	IGAGLYILGAIGLIFEFTFFECFAFAAMISAVDPVGTLAI
	281	320
O. sativa @9SXJB	(167)	LN-QDETPLYSLVFGGCVVNDATSIVLFNALQN----FD
Arabidopsis AtNhx1	(165)	LN-QDETPLLYSLVFGGCVVNDATSIVVFNAIQS----FD
SEQ ID #2	(172)	LN-QDETPLLYSLVFGGCVVNDATSIVLSRAVQT----YN
D.melanogaster @9VPL	(250)	FHALDVPILNMLVFGESILNDAISIVLTASITQSAN-VN
C. elegans @9X14	(277)	FQAVKVESLLYMLVFGESMLNDAVSIVLAATALRHAKPSF

Figure 4B

		321	360
<i>O. sativa</i> Q9SXJ8	(202)	LVHIDAAVVLKFLGNFFYLFLSSTFLGVFAGLLSAYIICK	
<i>Arabidopsis AtNhx1</i>	(200)	LTHLNHEAAFHLLGNFLYLFLLLSTLLGAATGLISAYVIKK	
SEQ ID #2	(207)	FDNFTSLEGLQIGGSFLYLFFSSCILGIASGLISAYIIKT	
<i>D.melanogaster</i> Q9VPL	(289)	AEASTGEAMFSALKTFAMFFASAGIGVIFALISALLLKH	
<i>C. elegans</i> Q9X14	(317)	NSLPASEIITSFAVTFTEMFFFSACLGVGIGLLSALLFKH	
		361	400
<i>O. sativa</i> Q9SXJ8	(242)	LYIGRHSTDREVALMMLMAYLSYMLAELLDLSGILTVFFC	
<i>Arabidopsis AtNhx1</i>	(240)	LYFGRHSTDREVALMMLMAYLSYMLAELFDLSGILTVFFC	
SEQ ID #2	(247)	MYFGRHSTDREIAIMTLMAYLSYVFAELFYLSGILSVFFC	
<i>D.melanogaster</i> Q9VPL	(329)	IDLRKHP-SLEFAMMLMFTYAPYVLAEGIHLSGIMAILFC	
<i>C. elegans</i> Q9X14	(357)	VDLRKTP-SLEFALLLIFSYPYGF AEALDLSGIMAILFC	
		401	440
<i>O. sativa</i> Q9SXJ8	(282)	GIVMSHYTWHNVTESSRVTTKHAFATLSFIAETFLFLYVG	
<i>Arabidopsis AtNhx1</i>	(280)	GIVMSHYTWHNVTESSRITTKHTFATLSFLAETFFIFLYVG	
SEQ ID #2	(287)	GIVMSHYTWHNVTESSRITSKHSFATMSFIAETFFIFLYVG	
<i>D.melanogaster</i> Q9VPL	(368)	GIVMSHYTHFNLSVTQTITMQTMRTLAFIAETCVFAYLG	
<i>C. elegans</i> Q9X14	(396)	GISMSQFTRHNVSPIAQITFRHTFRTISFVAETSTFAYIG	
		441	480
<i>O. sativa</i> Q9SXJ8	(322)	MDALDIEKWEFASDRPGKSIGISSILLGLVLI GRAAFVFP	
<i>Arabidopsis AtNhx1</i>	(320)	MDALDIDKWRVSDTPGTSIAVSSILMGLVMVGRAAFVFP	
SEQ ID #2	(327)	MDALDFEKWKMMQSSFTESAGLFGSLLFLVLLGRAAFVFP	
<i>D.melanogaster</i> Q9VPL	(408)	LAIFSFKHQVELSF-----VIWA--IVLCLIGRACNIFP	
<i>C. elegans</i> Q9X14	(436)	MAFFTIKLNFAPWLI FWS-----VVLCLLGRACNIFP	
		481	520
<i>O. sativa</i> Q9SXJ8	(362)	LSFLSNLTKKAPNEKITWRQVVIW WAGLMRGAVSIALAY	
<i>Arabidopsis AtNhx1</i>	(360)	LSFLSNLAKKNQSEKINFNMQVVIWWSGLMRGAVSMALAY	
SEQ ID #2	(367)	LSALSNYSTKSPDAKINLRQMVIIW WAGLMRGAVSIALAF	
<i>D.melanogaster</i> Q9VPL	(440)	LAFLVNKFR---EHKINNMQFIMWFSGLR-GAISYALS	
<i>C. elegans</i> Q9X14	(468)	LAYLVNQCR--KDVQISMKNQIIMWFSGMR-GAVCFALVL	
		521	560
<i>O. sativa</i> Q9SXJ8	(402)	NKFTRSGHTQLHGNAIMITSTITVVL FSTMVFGMMTKPLI	
<i>Arabidopsis AtNhx1</i>	(400)	NKFTRAGHTDVRGNAIMITSTITVCL FSTVVFGLTKPLI	
SEQ ID #2	(407)	N---QGGDAKDSNQATLMVITIIIVL FSTIVFGTATKPLI	
<i>D.melanogaster</i> Q9VPL	(476)	HLN----LDSQEKRHVIITTTLLIIVL FTTLVLGGSTMPLL	
<i>C. elegans</i> Q9X14	(505)	YMDLDK-----EKKSILLTTVLFLLI LFTTIFLGGSALPFI	
		561	600
<i>O. sativa</i> Q9SXJ8	(442)	RLLLPAS--GHPVTSEPSSPKS-----LHSPLLTSMQGS	
<i>Arabidopsis AtNhx1</i>	(440)	SYLLPHQ NATTSM L SDDNTPKS-----IHIPLLDQDSFIE	
SEQ ID #2	(444)	SULLPPHFRSNYSASLSPKASLDAD FHIPLLM DTEREE	
<i>D.melanogaster</i> Q9VPL	(512)	KYLKPGKRRRARGSGRNAAEEGG--RRNGSGRKRKSKSISL	
<i>C. elegans</i> Q9X14	(540)	SFINRCYP-NERQRKRRRTPRNKESTGNSSALMMSKTQEM	
		601	640
<i>O. sativa</i> Q9SXJ8	(475)	LEST-----TNIVRPSSLRMLLTKPTHTVHYYW	
<i>Arabidopsis AtNhx1</i>	(475)	PSGN-----HNVPRPDSIRGFLTRP TRTVHYYW	
SEQ ID #2	(484)	LEANDRSTI---NQILNGLPCPQSIGMLL TAPRSTIHHVW	
<i>D.melanogaster</i> Q9VPL	(550)	SKTREWQQA---IDSEHLSSELTEEDVTFTQARDRFRGM	
<i>C. elegans</i> Q9X14	(579)	SFFGSDDWGPKKSALDATSSAGRIMRQLFVRKFTAIERLE	

Figure 4C

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        641                                     680
O. sativa Q9SXJ8 (503) RKFD DALMRPMFGGR-----G
Arabidopsis AtNhx1 (503) RQFDDSFMRPVFGGR-----G
      SEQ ID #2 (521) RKFD DSYMRPTFGGR-----G
D.melanogaster Q9VP1 (587) RKYFIPFFTRRFNSQELHECKSQMAD-----LTNK
C. elegans Q9X14 (619) NRDKLAALTKRALASDQMTDSDDVEFGGGGGVGGGGRMKD

        681                                     720
O. sativa Q9SXJ8 (519) FVPFSPGSPTEQSHGGR-----
Arabidopsis AtNhx1 (519) FVPFVPGSPTERNPPDLKA-----
      SEQ ID #2 (537) YVRLVSRDMDIQEDELLEDHS-----
D.melanogaster Q9VP1 (617) WYQAIRVSPLDSDSDDEEIGLAAS--TSQIHLTRS----
C. elegans Q9X14 (659) DVPTRGRSGSRNSSDVIISAGGGGVSGEHHLLISSGSDS

        721
O. sativa Q9SXJ8 (536) -----SEQ ID NO: 23
Arabidopsis AtNhx1 (539) -----SEQ ID NO: 24
      SEQ ID #2 (559) -----SEQ ID NO: 2
D.melanogaster Q9VP1 (650) -----SEQ ID NO: 25
C. elegans Q9X14 (699) STNEF SEQ ID NO: 26
    
```

Figure 5

	(112)	112	120	130	140	150	160	170	189																							
ppNHx2(108)	LELD	MTQ	TKTKGKQ	AMSI	AGI	LEPF	VAG	GVSV	FWL	HL	II	AP	EG	AF	GP	ET	VF	MG	VAM	S	IT	AF	FP	VL	AR	LI	AE	RK	LI	TT	SIN:	27
A. thaliana C9SUQ7(110)	LELD	PKS	TKTKGKQ	AMSI	AGI	LEPF	VAG	GVSV	FWL	HL	II	AP	EG	AF	GP	ET	VF	MG	VAM	S	IT	AF	FP	VL	AR	LI	AE	RK	LI	TT	SIN:	28
O. sativa C9SX18 (71)	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	SIN:	29
A. thaliana C9SS27 (71)	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	---	SIN:	30
A. thaliana C9LUN4(112)	LELD	FAA	TKTKGKQ	AMSI	AGI	LEPF	VAG	GVSV	FWL	HL	II	AP	EG	AF	GP	ET	VF	MG	VAM	S	IT	AF	FP	VL	AR	LI	AE	RK	LI	TT	SIN:	31
Consensus(112)	LELD		IKTKGK AL	IASAGIT	LEPFV	GLG	SFLL	TIA	AG	QV	PF	LV	MG	VAM	S	IT	AF	FP	VL	AR	LI	AE	RK	LI	TT	SIN:	32					

Figure 6

Identity (similarity)	O.Sativa Q9SXJ8	SEQ ID #2	SEQ ID #5	D.Melanogaster Q9VP1	C.elegans Q9X14	AtNHX1 AAD 16946	Arabidopsis BAB02053
O.Sativa Q9SXJ8	-	-	-	-	-	-	-
SEQ ID #2	56 (67)	-	-	-	-	-	-
SEQ ID #5	10 (20)	27 (32)	-	-	-	-	-
D.Melanogaster Q9VP1	25 (40)	24 (39)	11 (22)	-	-	-	-
C.elegans Q9X14	22 (34)	24 (37)	10 (23)	36 (52)	-	-	-
AtNHX1 AAD 16946	72 (81)	56 (68)	10 (20)	24 (38)	22 (36)	-	-
Arabidopsis BAB02053	10 (22)	11 (22)	50 (63)	11 (22)	12 (26)	10 (21)	-