(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau

(43) International Publication Date
11 December 2003 (11.12.2003)

PCT

(10) International Publication Number
WO 03/101400 A2

(51) International Patent Classification[7]: A61K

(21) International Application Number: PCT/US03/17559

(22) International Filing Date: 4 June 2003 (04.06.2003)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/385,505 4 June 2002 (04.06.2002) US

(71) Applicant *(for all designated States except US)*: **AVALON PHARMACEUTICALS, INC.** [US/US]; 20358 Seneca Meadows Parkway, Germantown, MD 20876 (US).

(72) Inventors; and

(75) Inventors/Applicants *(for US only)*: **SHEA, Martin** [US/US]; 345 Market Street West #316, Gaithersburg, MD 20878 (US). **EBNER, Reinhard** [US/US]; 9906 Shelburn Terrace #316, Gaithersburg, MD 20878 (US).

(74) Agents: **GRANT, Alan, J.** et al.; Carella, Byrne, Bain, Gilfillan, Cecchi, Stewart &, Olstein, 6 Becker Farm Road, Roseland, NJ 07068 (US).

(81) Designated States *(national)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States *(regional)*: ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

**Published:**
— *without international search report and to be republished upon receipt of that report*

*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

WO 03/101400 A2

(54) Title: CANCER-LINKED GENE AS TARGET FOR CHEMOTHERAPY

(57) Abstract: Cancer-linked gene sequences, and derived amino acid sequences, are disclosed along with processes for assaying potential antitumor agents based on their modulation of the expression of these cancer-linked genes. Also disclosed are antibodies that react with the disclosed polypeptides and methods of using the antibodies to treat cancerous conditions, such as by using the antibody to target cancerous cells *in vivo* for purposes of delivering therapeutic agents thereto. Also described are methods of diagnosing using the gene sequences.

# CANCER-LINKED GENE AS TARGET FOR CHEMOTHERAPY

This application claims priority of U.S. Provisional Patent Application 60/385,505, filed 4 June 2002, the disclosure of which is hereby incorporated by reference in its entirety.

## FIELD OF THE INVENTION

The present invention relates to methods of screening cancer-linked genes and expression products for involvement in the cancer initiation and facilitation process as a means of cancer diagnosis as well as the use of such genes for screening potential anti-cancer agents, including small organic compounds and other molecules, and development of therapeutic agents.

## BACKGROUND OF THE INVENTION

Cancer-linked genes are valuable in that they indicate genetic differences between cancer cells and normal cells, such as where a gene is expressed in a cancer cell but not in a non-cancer cell, or where said gene is over-expressed or expressed at a higher level in a cancer as opposed to normal or non-cancer cell. In addition, the expression of such a gene in a normal cell but not in a cancer cell, especially of the same type of tissue, can indicate important functions in the cancerous process. For example, screening assays for novel drugs are based on the response of model cell based

systems *in vitro* to treatment with specific compounds. Such genes are also useful in the diagnosis of cancer and the identification of a cell as cancerous. Gene activity is readily measured by measuring the rate of production of gene products, such as RNAs and polypeptides encoded by such genes. Where genes encode cell surface proteins, appearance of, or alterations in, such proteins, as cell surface markers, are an indication of neoplastic activity. Some such screens rely on specific genes, such as oncogenes (or gene mutations). In accordance with the present invention, a cancer-linked gene has been identified and its putative amino acid sequence worked out. Such gene is useful in the diagnosing of cancer, the screening of anticancer agents and the treatment of cancer using such agents, especially in that these genes encode polypeptides that can act as markers, such as cell surface markers, thereby providing ready targets for anti-tumor agents such as antibodies, preferably antibodies complexed to cytotoxic agents, including apoptotic agents.

## BRIEF SUMMARY OF THE INVENTION

In accordance with the present invention, there is provided herein a cancer specific gene, linked especially to kidney cancer, or otherwise involved in the cancer initiating and facilitating process and the derived amino acid sequence thereof, including a number of different transcripts derived from said gene.

In one aspect, the present invention relates to a process for identifying an agent that modulates the activity of a cancer-related gene comprising:

(a) contacting a compound with a cell containing a gene that corresponds to a polynucleotide having a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33, and under conditions promoting the expression of said gene; and

(b) detecting a difference in expression of said gene relative to when said compound is not present

thereby identifying an agent that modulates the activity of a cancer-related gene.

In various embodiments of such a process, the cell is a cancer cell and the difference in expression is a decrease in expression. Such polynucleotides may also include those that have sequences identical to SEQ ID NO: 1-7, 14-20 and 27-33.

In another aspect, the present invention relates to a process for identifying an anti-neoplastic agent comprising contacting a cell exhibiting neoplastic activity with a compound first identified as a cancer related gene modulator using an assay process disclosed herein and detecting a decrease in said neoplastic activity after said contacting compared to when said contacting does not occur. Such neoplastic activity may include accelerated cellular replication and/or metastasis, and the decrease in neoplastic activity preferably results from the death of the cell, or senescence, terminal differentiation or growth inhibition.

The present invention also relates to a process for identifying an anti-neoplastic agent comprising administering to an animal exhibiting a cancer condition an effective amount of an agent first identified according to a process of one of one of the assays disclosed according to the invention and detecting a decrease in said cancerous condition.

The present invention further relates to a process for determining the cancerous status of a cell, comprising determining an increase in the level of expression in said cell of at least one gene that corresponds to a polynucleotide having a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33 wherein an elevated expression relative to a known non-cancerous cell indicates a cancerous state or potentially

cancerous state. Such elevated expression may be due to an increased copy number.

The present invention additionally relates to an isolated polypeptide, encoded by one of the polynucleotide transcripts disclosed herein, comprising an amino acid sequence homologous to an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39, wherein any difference between said amino acid sequence and the sequence of SEQ ID NO: 8-13, 21-26 and 34-39 is due solely to conservative amino acid substitutions and wherein said isolated polypeptide comprises at least one immunogenic fragment. In a preferred embodiment, the present invention encompasses an isolated polypeptide comprising an amino acid sequence homologous to an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39. These represent kidney cell surface antigens.

The present invention also relates to an antibody that reacts with a polypeptide as disclosed herein, preferably a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39. Such an antibody may be polyclonal, monoclonal, recombinant or synthetic in origin.

In one such embodiment, said antibody is associated, either covalently or non-covalently, with a cytotoxic agent, for example, an apoptotic agent. Thus, the present invention relates to an immunoconjugate comprising an antibody of the invention and a cytotoxic agent.

The present invention also relates to a process for treating cancer comprising contacting a cancerous cell with an agent having activity against an expression product encoded by a gene sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33. In one such embodiment, the

cancerous cell is contacted *in vivo*. In another such embodiment, said agent has affinity for said expression product. In a preferred embodiment, such agent is an antibody disclosed herein, such as an antibody that is specific or selective for, or otherwise reacts with, a polypeptide of the invention. In a preferred embodiment, the expression product is a polypeptide incorporating an amino acid sequence selected from SEQ ID NO: 8-13, 21-26 and 34-39.

The present invention further encompasses an immunogenic composition comprising a polypeptide disclosed herein, as well as compositions formed using antibodies specific for these polypeptides.

The present invention is also directed to uses of such compositions. Such uses include a method for treating cancer in an animal afflicted therewith comprising administering to said animal an amount of an immunogenic composition of one or more of the polypeptides disclosed herein where such amount is an amount sufficient to elicit the production of cytotoxic T lymphocytes specific for a polypeptide of the invention, preferably a polypeptide incorporating a sequence of SEQ ID NO: 8-13, 21-26 and 34-39. In a preferred embodiment, the animal to be so treated is a human patient.

## DEFINITIONS

As used herein, the terms "portion," "segment," and "fragment," when used in relation to polypeptides, refer to a continuous sequence of residues, such as amino acid residues, which sequence forms a subset of a larger sequence. For example, if a polypeptide were subjected to treatment with any of the common endopeptidases, such as trypsin or chymotrypsin, the oligopeptides resulting from such treatment would represent portions, segments or fragments of the starting polypeptide. When used in relation to a polynucleotides, such

terms refer to the products produced by treatment of said polynucleotides with any of the common endonucleases.

As used herein, the term "isolated" means that the material is removed from its original environment (e.g., the natural environment if it is naturally occurring). It could also be produced recombinantly and subsequently purified. For example, a naturally-occurring polynucleotide or polypeptide present in a living animal is not isolated, but the same polynucleotide or polypeptide, separated from some or all of the coexisting materials in the natural system, is isolated. Such polynucleotides, for example, those prepared recombinantly, could be part of a vector and/or such polynucleotides or polypeptides could be part of a composition, and still be isolated in that such vector or composition is not part of its natural environment. In one embodiment of the present invention, such isolated, or purified, polypeptide is useful in generating antibodies for practicing the invention, or where said antibody is attached to a cytotoxic or cytolytic agent, such as an apoptotic agent.

The term "percent identity" or "percent identical," when referring to a sequence, means that a sequence is compared to a claimed or described sequence after alignment of the sequence to be compared (the "Compared Sequence") with the described or claimed sequence (the "Reference Sequence"). The Percent Identity is then determined according to the following formula:

$$\text{Percent Identity} = 100\,[1-(C/R)]$$

wherein C is the number of differences between the Reference Sequence and the Compared Sequence over the length of alignment between the Reference Sequence and the Compared Sequence wherein (i) each base or amino acid in the Reference Sequence that does not have a corresponding aligned base or amino acid in the Compared Sequence and (ii) each gap in the Reference Sequence and (iii) each aligned base or amino acid in the Reference Sequence

that is different from an aligned base or amino acid in the Compared Sequence, constitutes a difference; and R is the number of bases or amino acids in the Reference Sequence over the length of the alignment with the Compared Sequence with any gap created in the Reference Sequence also being counted as a base or amino acid.

If an alignment exists between the Compared Sequence and the Reference Sequence for which the percent identity as calculated above is about equal to or greater than a specified minimum Percent Identity then the Compared Sequence has the specified minimum percent identity to the Reference Sequence even though alignments may exist in which the hereinabove calculated Percent Identity is less than the specified Percent Identity.

As known in the art "similarity" between two polypeptides is determined by comparing the amino acid sequence and its conserved amino acid substitutes of one polypeptide to the sequence of a second polypeptide.

In accordance with the present invention, the term "DNA segment" or "DNA sequence" refers to a DNA polymer, in the form of a separate fragment or as a component of a larger DNA construct, which has been derived from DNA isolated at least once in substantially pure form, i.e., free of contaminating endogenous materials and in a quantity or concentration enabling identification, manipulation, and recovery of the segment and its component nucleotide sequences by standard biochemical methods, for example, using a cloning vector. Such segments are provided in the form of an open reading frame uninterrupted by internal nontranslated sequences, or introns, which are typically present in eukaryotic genes. Sequences of non-translated DNA may be present downstream from the open reading frame, where the same do not interfere with manipulation or expression of the coding regions.

The term "coding region" refers to that portion of a gene which either naturally or normally codes for the expression product of that gene in its natural genomic environment, i.e., the region coding *in vivo* for the native expression product of the gene. The coding region can be from a normal, mutated or altered gene, or can even be from a DNA sequence, or gene, wholly synthesized in the laboratory using methods well known to those of skill in the art of DNA synthesis.

In accordance with the present invention, the term "nucleotide sequence" refers to a heteropolymer of deoxyribonucleotides. Generally, DNA segments encoding the proteins provided by this invention are assembled from cDNA fragments and short oligonucleotide linkers, or from a series of oligonucleotides, to provide a synthetic gene which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial, eukaryotic or viral operon.

The term "expression product" means that polypeptide or protein that is the natural translation product of the gene and any nucleic acid sequence coding equivalents resulting from genetic code degeneracy and thus coding for the same amino acid(s).

The term "active fragment," when referring to a coding sequence, means a portion comprising less than the complete coding region whose expression product retains essentially the same biological function or activity as the expression product of the complete coding region.

The term "primer" means a short nucleic acid sequence that is paired with one strand of DNA and provides a free 3'-OH end at which a DNA polymerase starts synthesis of a deoxyribonucleotide chain.

The term "promoter" means a region of DNA involved in binding of RNA polymerase to initiate transcription. The term "enhancer" refers to a region of DNA that, when present and active, has the effect of increasing expression of

a different DNA sequence that is being expressed, thereby increasing the amount of expression product formed from said different DNA sequence.

The term "open reading frame (ORF)" means a series of triplets coding for amino acids without any termination codons and is a sequence (potentially) translatable into protein.

As used herein, reference to a "DNA sequence" includes both single stranded and double stranded DNA. Thus, the specific sequence, unless the context indicates otherwise, refers to the single strand DNA of such sequence, the duplex of such sequence with its complement (double stranded DNA) and the complement of such sequence.

As used herein, "corresponding genes" refers to genes that encode an RNA that is at least 90% identical, preferably at least 95% identical, most preferably at least 98% identical, and especially identical, to an RNA encoded by one of the nucleotide sequences disclosed herein (i.e., SEQ ID NO: 1-7, 14-20 and 27-33). Such genes will also encode the same polypeptide sequence as any of the sequences disclosed herein, preferably SEQ ID NO: 1-7, 14-20 and 27-33, but may include differences in such amino acid sequences where such differences are limited to conservative amino acid substitutions, such as where the same overall three dimensional structure, and thus the same antigenic character, is maintained. Thus, amino acid sequences may be within the scope of the present invention where they react with the same antibodies that react with polypeptides comprising the sequences of SEQ ID NO: 8-13, 21-26 and 34-39. A "corresponding gene" includes splice variants thereof.

The genes identified by the present disclosure are considered "cancer-related" genes, as this term is used herein, and include genes expressed at higher levels (due, for example, to elevated rates of expression, elevated extent of expression or increased copy number) in cancer cells relative to

expression of these genes in normal (i.e., non-cancerous) cells where said cancerous state or status of test cells or tissues has been determined by methods known in the art, such as by reverse transcriptase polymerase chain reaction (RT-PCR) as described in the Examples herein. In specific embodiments, this relates to the genes whose sequences correspond to the sequences of SEQ ID NO: 1-7, 14-20 and 27-33.

As used herein, the term "conservative amino acid substitutions" are defined herein as exchanges within one of the following five groups:

I. Small aliphatic, nonpolar or slightly polar residues:
Ala, Ser, Thr, Pro, Gly;

II. Polar, negatively charged residues and their amides:
Asp, Asn, Glu, Gln;

III. Polar, positively charged residues:
His, Arg, Lys;

IV. Large, aliphatic, nonpolar residues:
Met Leu, Ile, Val, Cys

V. Large, aromatic residues:
Phe, Tyr, Trp

## DETAILED SUMMARY OF THE INVENTION

The present invention relates to processes for utilizing a nucleotide sequence for a cancer-linked gene, polypeptides encoded by such sequences and antibodies reactive with such polypeptides in methods of treating and diagnosing cancer, preferably kidney cancer, and in carrying out screening assays for agents effective in reducing the activity of cancer-linked genes and thereby treating a cancerous condition.

The polypeptides disclosed herein incorporate various polynucleotide transcripts (SEQ ID NO: 1-7, 14-20 and 27-33) and the derived amino acid sequence (SEQ ID NO: 8-13, 21-26 and 34-39) from said transcripts are available as targets for chemotherapeutic agents, especially anti-cancer agents, including antibodies specific for said polypeptides.

The cancer-related polynucleotide sequences disclosed herein correspond to gene sequences whose expression is indicative of the cancerous status of a given cell. Such sequences are substantially identical to SEQ ID NO: 1-7, 14-20 and 27-33, which represent different transcripts identified from the GenBank EST database and which exhibit cancer-specific expression. The polynucleotides of the invention are those that correspond to a sequence of SEQ ID NO: 1-7, 14-20 and 27-33. Such sequences have been searched within the GenBank database, especially the EST database, with results as follows:

| | |
|---|---|
| **Type:** | cell-surface tumor antigen<br>therapeutic antibody target |
| **Tissue:** | kidney |
| **Accession(s):** | AI479935, AI479935, AI186520 |
| **Unigene cluster-ID(s):** | Hs.61384 |
| **Chromosomal location:** | **3** |

The nucleotides and polypeptides, as gene products, used in the processes of the present invention may comprise a recombinant polynucleotide or polypeptide, a natural polynucleotide or polypeptide, or a synthetic polynucleotide or polypeptide, or a recombinant polynucleotide or polypeptide.

Fragments of such polynucleotides and polypeptides as are disclosed herein may also be useful in practicing the processes of the present invention.

For example, a fragment, derivative or analog of the polypeptide (SEQ ID NO: 8-13, 21-26 and 34-39) may be (i) one in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code, or (ii) one in which one or more of the amino acid residues includes a substituent group, or (iii) one in which the mature polypeptide is fused with another compound, such as a compound to increase the half-life of the polypeptide (for example, polyethylene glycol), or (iv) one in which the additional amino acids are fused to the mature polypeptide, such as a leader or secretory sequence or a sequence which is employed for purification of the mature polypeptide (such as a histidine hexapeptide) or a proprotein sequence. Such fragments, derivatives and analogs are deemed to be within the scope of those skilled in the art from the teachings herein.

In another aspect, the present invention relates to an isolated polypeptide, including a purified polypeptide, comprising an amino acid sequence at least 90% identical to the amino acid sequence of SEQ ID NO: 8-13 and/or 21-26 and/or 34-39. In preferred embodiments, said isolated polypeptide comprises an amino acid sequence having sequence identity of at least 95%, preferably at least about 98%, and especially is identical to, the sequence of SEQ ID NO: 8-13 and/or 21-26 and/or 34-39. The present invention also includes isolated active fragments of such polypeptides where said fragments retain the biological activity of the polypeptide or where such active fragments are useful as specific targets for cancer treatment, prevention or diagnosis. Thus, the present invention relates to any polypeptides, or fragments thereof, with sufficient sequence homology to the sequences disclosed herein as to be useful in the production of antibodies that react with (i.e., are selective or specific for) the polypeptides of SEQ ID NO: 8-13, 21-26 and 34-39 so as to be useful in targeting cells that exhibit such polypeptides, or fragments, on their surfaces, thereby providing targets for such antibodies and therapeutic agents associated with such antibodies.

The polynucleotides and polypeptides useful in practicing the processes of the present invention may likewise be obtained in an isolated or purified form. In addition, the polypeptide disclosed herein as being useful in practicing the processes of the invention are believed to be surface proteins present on cells, such as cancerous cells. Precisely how such cancer-linked proteins are used in the processes of the invention may thus differ depending on the therapeutic approach used. For example, cell-surface proteins, such as receptors, are desirable targets for cytotoxic antibodies that can be generated against the polypeptides disclosed herein.

The sequence information disclosed herein, as derived from the GenBank submissions, can readily be utilized by those skilled in the art to prepare the corresponding full-length polypeptide by peptide synthesis. The same is true for either the polynucleotides or polypeptides disclosed herein for use in the methods of the invention.

The present invention relates to an isolated polypeptide, encoded by one of the polynucleotide transcripts disclosed herein, comprising an amino acid sequence homologous to an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39, wherein any difference between amino acid sequence in the isolated polypeptide and the sequence of SEQ ID NO: 8-13, 21-26 and 34-39 is due solely to conservative amino acid substitutions and wherein said isolated polypeptide comprises at least one immunogenic fragment. In a preferred embodiment, the present invention encompasses an isolated polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39.

Methods of producing recombinant cells and vectors useful in preparing the polynucleotides and polypeptides disclosed herein are well known to those skilled in the molecular biology art. See, for example,

Sambrook, et al., Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, N.Y., (1989), Wu et al., *Methods in Gene Biotechnology* (CRC Press, New York, NY, 1997), and *Recombinant Gene Expression Protocols*, in *Methods in Molecular Biology*, Vol. 62, (Tuan, ed., Humana Press, Totowa, NJ, 1997), the disclosures of which are hereby incorporated by reference.

In one aspect, the present invention relates to a process for identifying an agent that modulates the activity of a cancer-related gene comprising:

(a) contacting a compound with a cell containing a gene that corresponds to a polynucleotide having a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33 and under conditions promoting the expression of said gene; and

(b) detecting a difference in expression of said gene relative to when said compound is not present

thereby identifying an agent that modulates the activity of a cancer-related gene.

In specific embodiments of such process the cell is a cancer cell and the difference in expression is a decrease in expression. Such polynucleotides may also include those that have sequences identical to SEQ ID NO: 1-7, 14-20 and 27-33.

In another aspect, the present invention relates to a process for identifying an anti-neoplastic agent comprising contacting a cell exhibiting neoplastic activity with a compound first identified as a cancer related gene modulator using an assay process disclosed herein and detecting a decrease in said neoplastic activity after said contacting compared to when said contacting does not occur. Such neoplastic activity may include accelerated cellular replication and/or metastasis, and the decrease in neoplastic activity preferably results from the death of the cell.

The present invention also relates to a process for identifying an anti-neoplastic agent comprising administering to an animal exhibiting a cancer condition an effective amount of an agent first identified according to a process of one of one of the assays disclosed according to the invention and detecting a decrease in said cancerous condition.

In specific embodiments of the present invention, the genes useful for the invention comprise genes that correspond to polynucleotides having a sequence selected from SEQ ID NO: 1-7, 14-20 and 27-33, or may comprise the sequence of any of the polynucleotides disclosed herein (where the latter are cDNA sequences).

In accordance with the present invention, such assays rely on methods of determining the activity of the gene in question. Such assays are advantageously based on model cellular systems using cancer cell lines, primary cancer cells, or cancerous tissue samples that are maintained in growth medium and treated with compounds at a single concentration or at a range of concentrations. At specific times after treatment, cellular RNAs are conveniently isolated from the treated cells or tissues, which RNAs are indicative of expression of selected genes. The cellular RNA is then divided and subjected to differential analysis that detects the presence and/or quantity of specific RNA transcripts, which transcripts may then be amplified for detection purposes using standard methodologies, such as, for example, reverse transcriptase polymerase chain reaction (RT-PCR), etc. The presence or absence, or concentration levels, of specific RNA transcripts are determined from these measurements. The polynucleotide sequences disclosed herein are readily used as probes for the detection of such RNA transcripts and thus the measurement of gene activity and expression.

The polynucleotides of the invention can include fully operational genes with attendant control or regulatory sequences or merely a polynucleotide

sequence encoding the corresponding polypeptide or an active fragment or analog thereof.

Because expression of the polynucleotide sequences disclosed herein are specific to the cancerous state, useful gene modulation is downward modulation, so that, as a result of exposure to an antineoplastic agent identified by the screening assays herein, the corresponding gene of the cancerous cell is expressed at a lower level (or not expressed at all) when exposed to the agent as compared to the expression when not exposed to the agent. For example, the gene sequences disclosed herein (SEQ ID NO: 1-7, 14-20 and 27-33) correspond to a gene expressed at a higher level in cells of kidney cancer than in normal kidney cells. Thus, where said chemical agent causes this gene of the tested cell to be expressed at a lower level than the same genes of the reference, this is indicative of downward modulation and indicates that the chemical agent to be tested has anti-neoplastic activity.

In carrying out the assays disclosed herein, relative antineoplastic activity may be ascertained by the extent to which a given chemical agent modulates the expression of genes present in a cancerous cell. Thus, a first chemical agent that modulates the expression of a gene associated with the cancerous state (i.e., a gene corresponding to one or more of the polynucleotide transcripts disclosed herein) to a larger degree than a second chemical agent tested by the assays of the invention is thereby deemed to have higher, or more desirable, or more advantageous, anti-neoplastic activity than said second chemical agent.

The gene expression to be measured is commonly assayed using RNA expression as an indicator. Thus, the greater the level of RNA (for example, messenger RNA or mRNA) detected the higher the level of expression of the corresponding gene. Thus, gene expression, either absolute or relative, is determined by the relative expression of the RNAs encoded by such genes.

RNA may be isolated from samples in a variety of ways, including lysis and denaturation with a phenolic solution containing a chaotropic agent (e.g., trizol) followed by isopropanol precipitation, ethanol wash, and resuspension in aqueous solution; or lysis and denaturation followed by isolation on solid support, such as a Qiagen resin and reconstitution in aqueous solution; or lysis and denaturation in non-phenolic, aqueous solutions followed by enzymatic conversion of RNA to DNA template copies.

Normally, prior to applying the processes of the invention, steady state RNA expression levels for the genes, and sets of genes, disclosed herein will have been obtained. It is the steady state level of such expression that is affected by potential anti-neoplastic agents as determined herein. Such steady state levels of expression are easily determined by any methods that are sensitive, specific and accurate. Such methods include, but are in no way limited to, real time quantitative polymerase chain reaction (PCR), for example, using a Perkin-Elmer 7700 sequence detection system with gene specific primer probe combinations as designed using any of several commercially available software packages, such as Primer Express software., solid support based hybridization array technology using appropriate internal controls for quantitation, including filter, bead, or microchip based arrays, solid support based hybridization arrays using, for example, chemiluminescent, fluorescent, or electrochemical reaction based detection systems.

The gene expression indicative of a cancerous state need not be characteristic of every cell of a given tissue. Thus, the methods disclosed herein are useful for detecting the presence of a cancerous condition within a tissue where less than all cells exhibit the complete pattern. Thus, for example, a selected gene corresponding to the sequence of SEQ ID NO: 1, may be found, using appropriate probes, either DNA or RNA, to be present in as little as 60% of cells derived from a sample of tumorous, or malignant, tissue. In a highly preferred embodiment, such gene pattern is found to be

present in at least 100% of cells drawn from a cancerous tissue and absent from at least 100% of a corresponding normal, non-cancerous, tissue sample.

Expression of a gene may be related to copy number, and changes in expression may be measured by determining copy number. Such change in gene copy number may be determined by determining a change in expression of messenger RNA encoded by a particular gene sequence, especially that of SEQ ID NO: 1-7, 14-20 and 27-33. Also in accordance with the present invention, said gene may be a cancer initiating or facilitating gene. In carrying out the methods of the present invention, a cancer facilitating gene is a gene that, while not directly initiating tumor formation or growth, acts, such as through the actions of its expression product, to direct, enhance, or otherwise facilitate the progress of the cancerous condition, including where such gene acts against genes, or gene expression products, that would otherwise have the effect of decreasing tumor formation and/or growth.

Although the expression of a gene corresponding to a sequence of SEQ ID NO: 1-7, 14-20 and 27-33 may be indicative of a cancerous status for a given cell, the mere presence of such a gene may not alone be sufficient to achieve a malignant condition and thus the level of expression of such gene may also be a significant factor in determining the attainment of a cancerous state. Thus, it becomes essential to also determine the level of expression of a gene as disclosed herein, including substantially similar sequences, as a separate means of diagnosing the presence of a cancerous status for a given cell, groups of cells, or tissues, either in culture or *in situ*.

The level of expression of the polypeptides disclosed herein is also a measure of gene expression, such as polypeptides having sequence identical, or similar to, any polypeptide encoded by a sequence of SEQ ID NO: 1-7, 14-20 and 27-33, especially a polypeptide whose amino acid sequence is the sequence of SEQ ID NO: 8-13, 21-26 and 34-39.

In accordance with the foregoing, the present invention specifically contemplates a method for determining the cancerous status of a cell to be tested, comprising determining the level of expression in said cell of a gene that includes one of the nucleotide sequences selected from the sequences of SEQ ID NO: 1-7, 14-20 and 27-33, including sequences substantially identical to said sequences, or characteristic fragments thereof, or the complements of any of the foregoing and then comparing said expression to that of a cell known to be non-cancerous whereby the difference in said expression indicates that said cell to be tested is cancerous.

In accordance with the invention, although gene expression for a gene that includes as a portion thereof one of the sequences of SEQ ID NO: 1-7, 14-20 and 27-33, is preferably determined by use of a probe that is a fragment of such nucleotide sequence, it is to be understood that the probe may be formed from a different portion of the gene. Expression of the gene may be determined by use of a nucleotide probe that hybridizes to messenger RNA (mRNA) transcribed from a portion of the gene other than the specific nucleotide sequence disclosed herein.

It should be noted that there are a variety of different contexts in which genes have been evaluated as being involved in the cancerous process. Thus, some genes may be oncogenes and encode proteins that are directly involved in the cancerous process and thereby promote the occurrence of cancer in an animal. In addition, other genes may serve to suppress the cancerous state in a given cell or cell type and thereby work against a cancerous condition forming in an animal. Other genes may simply be involved either directly or indirectly in the cancerous process or condition and may serve in an ancillary capacity with respect to the cancerous state. All such types of genes are deemed with those to be determined in accordance with the invention as disclosed herein. Thus, the gene determined by said process of the invention may be an oncogene, or the gene determined by said process may be a cancer facilitating gene, the latter including a gene that

directly or indirectly affects the cancerous process, either in the promotion of a cancerous condition or in facilitating the progress of cancerous growth or otherwise modulating the growth of cancer cells, either *in vivo* or *ex vivo*. In addition, the gene determined by said process may be a cancer suppressor gene, which gene works either directly or indirectly to suppress the initiation or progress of a cancerous condition. Such genes may work indirectly where their expression alters the activity of some other gene or gene expression product that is itself directly involved in initiating or facilitating the progress of a cancerous condition. For example, a gene that encodes a polypeptide, either wild or mutant in type, which polypeptide acts to suppress of tumor suppressor gene, or its expression product, will thereby act indirectly to promote tumor growth.

As noted previously, polynucleotides encoding the same proteins as any of SEQ ID NO: 1-7, 14-20 and 27-33, regardless of the percent identity of such sequences, are also specifically contemplated by any of the methods of the present invention that rely on any or all of said sequences, regardless of how they are otherwise described or limited. Thus, any such sequences are available for use in carrying out any of the methods disclosed according to the invention. Such sequences also include any open reading frames, as defined herein, present within the sequence of SEQ ID NO: 1-7, 14-20 and 27-33.

Because a gene disclosed according to the invention "corresponds to" a polynucleotide having a sequence of SEQ ID NO: 1-7, 14-20 and 27-33, said gene encodes an RNA (processed or unprocessed, including naturally occurring splice variants and alleles) that is at least 90% identical, preferably at least 95% identical, most preferably at least 98% identical to, and especially identical to, an RNA that would be encoded by, or be complementary to, such as by hybridization with, a polynucleotide having the indicated sequence. In addition, genes including sequences at least 90% identical to a sequence selected from SEQ ID NO: 1-7, 14-20 and 27-33, preferably at least about 95% identical to such a sequence, more preferably at

least about 98% identical to such sequence and most preferably comprising such sequence are specifically contemplated by all of the processes of the present invention. Sequences encoding the same proteins as any of these sequences, regardless of the percent identity of such sequences, are also specifically contemplated by any of the methods of the present invention that rely on any or all of said sequences, regardless of how they are otherwise described or limited. The polynucleotide sequences of the invention also include any open reading frames, as defined herein, present within any of the sequences of SEQ ID NO: 1-7, 14-20 and 27-33.

The sequences disclosed herein may be genomic in nature and thus represent the sequence of an actual gene, such as a human gene, or may be a cDNA sequence derived from a messenger RNA (mRNA) and thus represent contiguous exonic sequences derived from a corresponding genomic sequence, or they may be wholly synthetic in origin for purposes of practicing the processes of the invention. Because of the processing that may take place in transforming the initial RNA transcript into the final mRNA, the sequences disclosed herein may represent less than the full genomic sequence. They may also represent sequences derived from ribosomal and transfer RNAs. Consequently, the gene as present in the cell (and representing the genomic sequence) and the polynucleotide transcripts disclosed herein, including cDNA sequences, may be identical or may be such that the cDNAs contain less than the full genomic sequence. Such genes and cDNA sequences are still considered "corresponding sequences" (as defined elsewhere herein) because they both encode the same or related RNA sequences (i.e., related in the sense of being splice variants or RNAs at different stages of processing). Thus, by way of non-limiting example only, a gene that encodes an RNA transcript, which is then processed into a shorter mRNA, is deemed to encode both such RNAs and therefore encodes an RNA complementary to (using the usual Watson-Crick complementarity rules), or that would otherwise be encoded by, a cDNA (for example, a sequence as disclosed herein). Thus, the sequences disclosed herein correspond to genes

contained in the cancerous cells (here, kidney cancer) and are used to determine gene activity or expression because they represent the same sequence or are complementary to RNAs encoded by the gene. Such a gene also includes different alleles and splice variants that may occur in the cells used in the methods of the invention, such as where recombinant cells are used to assay for anti-neoplastic agents and such cells have been engineered to express a polynucleotide as disclosed herein, including cells that have been engineered to express such polynucleotides at a higher level than is found in non-engineered cancerous cells or where such recombinant cells express such polynucleotides only after having been engineered to do so. Such engineering includes genetic engineering, such as where one or more of the polynucleotides disclosed herein has been inserted into the genome of such cell or is present in a vector.

Such cells, especially mammalian cells, may also be engineered to express on their surfaces one or more of the polypeptides of the invention for testing with antibodies or other agents capable of masking such polypeptides and thereby removing the cancerous nature of the cell. Such engineering includes both genetic engineering, where the genetic complement of the cells is engineered to express the polypeptide, as well as non-genetic engineering, whereby the cell has been physically manipulated to incorporate a polypeptide of the invention in its plasma membrane, such as by direct insertion using chemical and/or other agents to achieve this result.

In accordance with the foregoing, the present invention includes anti-cancer agents that are themselves either polypeptides, or small chemical entities, that affect the cancerous process, including initiation, suppression or facilitation of tumor growth, either *in vivo* or *ex vivo.* Said cancer modulating agent will have the effect of decreasing gene expression.

The present invention thus also relates to a method for treating cancer comprising contacting a cancerous cell with an agent having activity against

an expression product encoded by a gene or polynucleotide sequence as disclosed herein, such as one having, or corresponding to, the nucleotide sequence of SEQ ID NO: 1-7, 14-20 and 27-33. The present invention also relates to a process for treating cancer comprising contacting a cancerous cell with an agent having activity against an expression product encoded by a gene or polynucleotide sequence corresponding to a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33. In one such embodiment, the cancerous cell is contacted *in vivo.* In another such embodiment, said agent has affinity for said expression product. In a preferred embodiment, such agent is an antibody disclosed herein, such as an antibody that is specific or selective for, or otherwise reacts with, a polypeptide of the invention. In a preferred embodiment, the expression product is a polypeptide incorporating an amino acid sequence selected from SEQ ID NO: 8-13, 21-26 and 34-39.

The present invention is also directed to such uses of the compositions of polypeptides and antibodies disclosed herein. Such uses include a process for treating cancer in an animal afflicted therewith comprising administering to said animal an amount of an immunogenic composition of one or more of the polypeptides disclosed herein where such amount if an amount sufficient to elicit the production of cytotoxic T lymphocytes specific for a polypeptide of the invention, preferably a polypeptide incorporating a sequence of SEQ ID NO: 8-13, 21-26 and 34-39. In a preferred embodiment, the animal to be so treated is a human patient.

The proteins encoded by the genes disclosed herein due to their expression, or elevated expression, in cancer cells, represent highly useful therapeutic targets for "targeted therapies" utilizing such affinity structures as, for example, antibodies coupled to some cytotoxic agent. In such methodology, it is advantageous that nothing need be known about the endogenous ligands or binding partners for such cell surface molecules. Rather, an antibody or equivalent molecule that can specifically recognize the

cell surface molecule (which could include an artificial peptide, a surrogate ligand, and the like) that is coupled to some agent that can induce cell death or a block in cell cycling offers therapeutic promise against these proteins. Thus, such approaches include the use of so-called suicide "bullets" against intracellular proteins. For example, monoclonal antibodies may readily by produced by methods well known in the art, for example, the method of Kohler and Milstein (see: *Nature*, **256**:495 (1975).

With the advent of methods of molecular biology and recombinant technology, it is now possible to produce antibody molecules by recombinant means and thereby generate gene sequences that code for specific amino acid sequences found in the polypeptide structure of the antibodies. Such antibodies can be produced by either cloning the gene sequences encoding the polypeptide chains of said antibodies or by direct synthesis of said polypeptide chains, with *in vitro* assembly of the synthesized chains to form active tetrameric ($H_2L_2$) structures with affinity for specific epitopes and antigenic determinants. This has permitted the ready production of antibodies having sequences characteristic of neutralizing antibodies from different species and sources.

Regardless of the source of the antibodies, or how they are recombinantly constructed, or how they are synthesized, *in vitro* or *in vivo*, using transgenic animals, such as cows, goats and sheep, using large cell cultures of laboratory or commercial size, in bioreactors or by direct chemical synthesis employing no living organisms at any stage of the process, all antibodies have a similar overall 3 dimensional structure. This structure is often given as $H_2L_2$ and refers to the fact that antibodies commonly comprise 2 light (L) amino acid chains and 2 heavy (H) amino acid chains. Both chains have regions capable of interacting with a structurally complementary antigenic target. The regions interacting with the target are referred to as "variable" or "V" regions and are characterized by differences in amino acid sequence from antibodies of different antigenic specificity.

The variable regions of either H or L chains contains the amino acid sequences capable of specifically binding to antigenic targets. Within these sequences are smaller sequences dubbed "hypervariable" because of their extreme variability between antibodies of differing specificity. Such hypervariable regions are also referred to as "complementarity determining regions" or "CDR" regions. These CDR regions account for the basic specificity of the antibody for a particular antigenic determinant structure.

The CDRs represent non-contiguous stretches of amino acids within the variable regions but, regardless of species, the positional locations of these critical amino acid sequences within the variable heavy and light chain regions have been found to have similar locations within the amino acid sequences of the variable chains. The variable heavy and light chains of all antibodies each have 3 CDR regions, each non-contiguous with the others (termed L1, L2, L3, H1, H2, H3) for the respective light (L) and heavy (H) chains. The accepted CDR regions have been described by Kabat et al., *J. Biol. Chem.* **252**:6609-6616 (1977).

In all mammalian species, antibody polypeptides contain constant (i.e., highly conserved) and variable regions, and, within the latter, there are the CDRs and the so-called "framework regions" made up of amino acid sequences within the variable region of the heavy or light chain but outside the CDRs.

The antibodies disclosed according to the invention may also be wholly synthetic, wherein the polypeptide chains of the antibodies are synthesized and, possibly, optimized for binding to the polypeptides disclosed herein as being receptors. Such antibodies may be chimeric or humanized antibodies and may be fully tetrameric in structure, or may be dimeric and comprise only a single heavy and a single light chain. Such antibodies may also include fragments, such as Fab and $F(ab_2)'$ fragments, capable of reacting with and binding to any of the polypeptides disclosed herein as being receptors.

In one aspect, the present invention relates to immunoglobulins, or antibodies, as described herein, that react with, especially where they are specific for, the polypeptides having amino acid sequences as disclosed herein, preferably those having an amino acid sequence of one of SEQ ID NO: 8-13, 21-26 and 34-39. Such antibodies may commonly be in the form of a composition, especially a pharmaceutical composition. Such antibodies, by themselves, may have therapeutic value in that they are able to bind to, and thereby tie up, surface sites on cancerous cells. Where such sites have some type of function to perform (i.e., where they are surface enzymes, or channel structures, or structures that otherwise facilitate, actively or passively, the transport of nutrients and other vital materials to the cell. Such nutrients serve to facilitate the growth and replication of the cell and molecules that bind to such sites and thereby interfere with such activities can prove to have a therapeutic effect in that the result of such binding is to remove sources of nutrients from such cells, thereby interfering with growth and replication. In like manner, such binding may serve to remove vital enzyme activities from the cell's functional repertoire, thereby also interfering with viability and/or the ability of the cell to multiply or metastasize. In addition, by binding to such surface sites, the antibodies may serve to prevent the cells from reacting to environmental agents, such as cytokines and the like, that may facilitate growth, replication and metastasis, thereby further reducing the cancerous status of such cell and ameliorating the cancerous condition in a patient, even without proving fatal to the cell or cells so affected.

The methods of the present invention also include processes wherein the cancer cell is contacted *in vivo* as well as *ex vivo* with an agent that comprises a portion, or is part of an overall molecular structure, having affinity for an expression product of a gene corresponding to a polynucleotide sequence as disclosed herein, preferably where the expression product is a cell surface structure, most preferably a polypeptide as disclosed herein, such as one that comprises an amino acid sequence of SEQ ID NO: 8-13, 21-26 and 34-39. In one such embodiment, said portion having affinity for said

expression product is an antibody, especially where said expression product is a polypeptide or oligopeptide or comprises an oligopeptide portion, or comprises a polypeptide.

In another aspect, the present invention also relates to an antibody that reacts with a polypeptide as disclosed herein, preferably a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39. Such an antibody may be polyclonal, monoclonal, recombinant or synthetic in origin. In one such embodiment, said antibody is associated, either covalently or non-covalently, with a cytotoxic agent, for example, an apoptotic agent. It is thus contemplated that the antibody acts a targeted vector for guiding an associated therapeutic agent to a cancerous cell, such as a cell expressing a polypeptide homologous to, if not identical to, a polypeptide as disclosed herein.

Where the cytotoxic agent is itself a polypeptide, said may be linked directly to an antibody specific for a surface target on a cancer cell, such as where the polypeptide represents an extension of the amino acid chain of the antibody. In alternative embodiments, such molecules may be covalently linked through a linker sequence of long or short duration, such as an amino acid sequence of 5 to 10 residues in length. Where the cytotoxic agents is some small organic molecule, such as a small organic compound, or some type of apoptotic agent, this may be covalently bonded to the antibody molecule or may be attached by some other type of non-covalent linkage, including hydrophobic and electrostatic linkages. Methods for forming such linkages, especially covalent linkages, are well known to those skilled in the art.

The antibodies disclosed herein may also serve as targeting vectors for much larger structures, such as liposomes. In one such embodiment, an antibody is part of, or otherwise linked to, or associated with, a membranous structure, preferably a liposome or possibly some type of cellular organelle,

which acts as a reservoir for a cytotoxic agent, such as ricin. The antibody then acts to target said liposome to a cancerous tissue in an animal, whereupon the liposome provides a source of cytotoxic agents for localized treatment of a solid tumor or other type of neoplasm.

The present invention further encompasses an immunogenic composition comprising a polypeptide disclosed herein, as well as compositions formed using antibodies specific for these polypeptides.

Methods well known in the art for making formulations are found in, for example, *Remington: The Science and Practice of Pharmacy*, (19th ed.) Ed. A.R. Gennaro, 1995, Mack Publishing Company, Easton, PA. Formulations for parenteral administration may, for example, contain excipients, sterile water, or saline, polyalkylene glycols such as polyethylene glycol, oils of vegetable origin, or hydrogenated napthalenes. Biocompatible, biodegradable lactide polymer, lactide/glycolide copolymer, or polyoxyethylene-polyoxypropylene copolymers may be used to control the release of the compounds. Other potentially useful parenteral delivery systems for agonists of the invention include ethylenevinyl acetate copolymer particles, osmotic pumps, implantable infusion systems, and liposomes. Formulations for inhalation may contain excipients, or example, lactose, or may be aqueous solutions containing, for example, polyoxyethylene-9-lauryl ether, glycocholate and deoxycholate, or may be oily solutions for administration in the form of nasal drops, or as a gel. It should be noted that, where the therapeutic agent to be administered is an immunoconjugate, these sometimes contain chemical linkages that are somewhat labile in aqueous media and therefor must be stored prior to administration is a more stable environment, such as in the form of a lyophilized powder.

Such an agent can be a single molecular structure, comprising both affinity portion and anti-cancer activity portions, wherein said portions are derived from separate molecules, or molecular structures, possessing such

activity when separated and wherein such agent has been formed by combining said portions into one larger molecular structure, such as where said portions are combined into the form of an adduct. Said anti-cancer and affinity portions may be joined covalently, such as in the form of a single polypeptide, or polypeptide-like, structure or may be joined non-covalently, such as by hydrophobic or electrostatic interactions, such structures having been formed by means well known in the chemical arts. Alternatively, the anti-cancer and affinity portions may be formed from separate domains of a single molecule that exhibits, as part of the same chemical structure, more than one activity wherein one of the activities is against cancer cells, or tumor formation or growth, and the other activity is affinity for an expression product produced by expression of genes related to the cancerous process or condition.

In one embodiment of the present invention, a chemical agent, such as a protein or other polypeptide, is joined to an agent, such as an antibody, having affinity for an expression product of a cancerous cell, such as a polypeptide or protein encoded by a gene related to the cancerous process, preferably a gene as disclosed herein according to the present invention, most preferably a polypeptide sequence disclosed herein. Thus, where the presence of said expression product is essential to tumor initiation and/or growth, binding of said agent to said expression product will have the effect of negating said tumor promoting activity. In one such embodiment, said agent is an apoptosis-inducing agent that induces cell suicide, thereby killing the cancer cell and halting tumor growth.

Other genes within the cancer cell that are regulated in a manner similar to that of the genes disclosed herein and thus change their expression in a coordinated way in response to chemical compounds represent genes that are located within a common metabolic, signaling, physiological, or functional pathway so that by analyzing and identifying such commonly regulated groups of genes (groups that include the gene, or similar sequences, disclosed according to the invention, one can (a) assign known genes and novel genes to specific pathways and (b) identify specific functions

and functional roles for novel genes that are grouped into pathways with genes for which their functions are already characterized or described. For example, one might identify a group of 10 genes, at least one of which is the gene as disclosed herein, that change expression in a coordinated fashion and for which the function of one, such as the polypeptide encoded by the sequence disclosed herein, is known then the other genes are thereby implicated in a similar function or pathway and may thus play a role in the cancer-initiating or cancer-facilitating process. In the same way, if a gene were found in normal cells but not in cancer cells, or happens to be expressed at a higher level in normal as opposed to cancer cells, then a similar conclusion may be drawn as to its involvement in cancer, or other diseases. Therefore, the processes disclosed according to the present invention at once provide a novel means of assigning function to genes, i.e. a novel method of functional genomics, and a means for identifying chemical compounds that have potential therapeutic effects on specific cellular pathways. Such chemical compounds may have therapeutic relevance to a variety of diseases outside of cancer as well, in cases where such diseases are known or are demonstrated to involve the specific cellular pathway that is affected.

The polypeptides disclosed herein, preferably those of SEQ ID NO: 8-13, 21-26 and 34-39, also find use as vaccines in that, where the polypeptide represents a surface protein present on a cancer cell, such polypeptide may be administered to an animal, especially a human being, for purposes of activating cytotoxic T lymphocytes (CTLs) that will be specific for, and act to lyze, cancer cells in said animal. Where used as vaccines, such polypeptides are present in the form of a pharmaceutical composition. The present invention may also employ polypeptides that have the same, or similar, immunogenic character as the polypeptides of SEQ ID NO: 8-13, 21-26 and 34-39 and thereby elicit the same, or similar, immunogenic response after administration to an animal, such as an animal at risk of developing cancer, or afflicted therewith. Thus, the polypeptides disclosed according to the invention will commonly find use as immunogenic compositions.

Expression of a gene corresponding to a polynucleotide disclosed herein, when in normal tissues, may indicate a predisposition towards development of kidney cancer. The encoded polypeptide might then present a potentially useful cell surface target for therapeutic molecules such as cytolytic antibodies, or antibodies attached to cytotoxic, or cytolytic, agents. .

The present invention specifically contemplates use of antibodies against the polypeptides encoded by the polynucleotides corresponding to the genes disclosed herein, whereby said antibodies are conjugates to one or more cytotoxic agents so that the antibodies serve to target the conjugated immunotoxins to a region of cancerous activity, such as a solid tumor. For many known cytotoxic agents, lack of selectivity has presented a drawback to their use as therapeutic agents in the treatment of malignancies. For example, the class of two-chain toxins, consisting of a binding subunit (or B-chain) linked to a toxic subunit (A-chain) are extremely cytotoxic. Thus, such agents as ricin, a protein isolated from castor beans, kills cells at very low concentrations (even less than $10^{-11}$ M) by inactivating ribosomes in said cells (see, for example, Lord et al., Ricin: structure, mode of action, and some current applications. *Faseb J,* **8***:* 201-208 (1994), and Blättler et al., Realizing the full potential of immunotoxins. *Cancer Cells*, **1***:* 50-55 (1989)). While isolated A-chains of protein toxins that functionally resemble ricin A-chain are only weakly cytotoxic for intact cells (in the concentration range of $10^{-7}$ to $10^{-6}$ M), they are very potent cytotoxic agents inside the cells. Thus, a single molecule of the A-subunit of diphtheria toxin can kill a cell once inside (see: Yamaizumi et al., One molecule of diphtheria toxin fragment A introduced into a cell can kill the cell. Cell, *15:* 245-250, 1978).

The present invention solves this selectivity problem by using antibodies specific for antigens present on cancer cells to target the cytotoxins to said cells. In addition, use of antibodies decreases toxicity because the antibodies are non-toxic until they reach the tumor and, because the cytotoxin

is bound to the antibody, it is presented with less opportunity to cause damage to non-targeted tissues.

In addition, use of such antibodies alone can provide therapeutic effects on the tumor through the antibody-dependent cellular cytotoxic response (ADCC) and complement-mediated cell lysis mechanisms.

A number of recombinant immunotoxins (for example, consisting of Fv regions of cancer specific antibodies fused to truncated bacterial toxins) are well known (see, for example, Smyth et al., Specific targeting of chlorambucil to tumors with the use of monoclonal antibodies, *J. Natl. Cancer Inst.*, **76**(3):503-510 (1986); Cho et al., Single-chain Fv/folate conjugates mediate efficient lysis of folate-receptor-positive tumor cells, *Bioconjug. Chem.*, **8**(3):338-346 (1997)). As noted in the literature, these may contain, for example, a truncated version of *Pseudomonas* exotoxin as a toxic moiety but the toxin is modified in such a manner that by itself it does not bind to normal human cells, but it retains all other functions of cytotoxicity. Here, recombinant antibody fragments target the modified toxin to cancer cells which are killed, such as by direct inhibition of protein synthesis, or by concomitant induction of apoptosis. Cells that are not recognized by the antibody fragment, because they do not carry the cancer antigen, are not affected. Good activity and specificity has been observed for many recombinant immunotoxins in *in vitro* assays using cultured cancer cells as well as in animal tumor models. Ongoing clinical trials provide examples where the promising pre-clinical data correlate with successful results in experimental cancer therapy. (see, for example, Brinkmann U., Recombinant antibody fragments and immunotoxin fusions for cancer therapy, *In Vivo* (2000) **14**:21-27).

While the safety of employing immunoconjugates in humans has been established, *in vivo* therapeutic results have been less impressive. Because clinical use of mouse MAbs in humans is limited by the development of a foreign anti-globulin immune response by the human host, genetically

engineered chimeric human-mouse MAbs have been developed by replacing the mouse Fc region with the human constant region. In other cases, the mouse antibodies have been "humanized" by replacing the framework regions of variable domains of rodent antibodies by their human equivalents. Such humanized and engineered antibodies can even be structurally arranged to have specificities and effector functions determined by design and which characteristics do not appear in nature. The development of bispecific antibodies, having different binding ends so that more than one antigenic site can be bound, have proven useful in targeting cancer cells. Thus, such antibody specificity has been improved by chemical coupling to various agents such as bacterial or plant toxins, radionuclides or cytotoxic drugs and other agents. (see, for example, Bodey, B. et al). Genetically engineered monoclonal antibodies for direct anti-neoplastic treatment and cancer cell specific delivery of chemotherapeutic agents. *Curr Pharm Des* (2000) Feb;6(3):261-76). See also, Garnett, M. C., Targeted drug conjugates: principles and progress. *Adv. Drug Deliv. Rev.* (2001 Dec 17) **53**(2):171-216; Brinkmann et al., Recombinant immunotoxins for cancer therapy. *Expert Opin Biol Ther.* (2001) **1**(4):693-702.

Among the cytotoxic agents specifically contemplated for use as immunoconjugates according to the present invention are Calicheamicin, a highly toxic enediyne antibiotic isolated from *Micromonospora echinospora ssp. Calichensis*, and which binds to the minor groove of DNA to induce double strand breaks and cell death (see: Lee et al., Calicheamicins, a novel family of antitumor antibiotics. 1. Chemistry and partial structure of calichemicin $g_1$. *J Am Chem Soc,* **109***:* 3464-3466 (1987); Zein et al., Calicheamicin gamma 1I: an antitumor antibiotic that cleaves double-stranded DNA site specifically, *Science*, **240***:* 1198-1201 (1988)). Useful derivatives of the calicheamicins include mylotarg and 138H11-Camθ. Mylotarg is an immunoconjugate of a humanized anti-CD33 antibody (CD33 being found in leukemic cells of most patients with acute myeloid leukemia) and N-acetyl gamma colicheamicin dimethyl hydrazide, the latter of which is

readily coupled to an antibody of the present invention (in place of the anti-CD33 but which can also be humanized by substitution of human framework regions into the antibody during production as described elsewhere herein) to form an immunoconjugate of the invention. (see: Hamann et al. Gemtuzumab Ozogamicin, A Potent and Selective Anti-CD33 Antibody-Calicheamicin Conjugate for Treatment of Acute Myeloid Leukemia, *Bioconjug. Chem.* **13**, 47-58 (2002)) For use with 138H11-Camθ, 138H11 is an anti-γ-glutamyl transferase antibody coupled to theta calicheamicin through a disulfide linkage and found useful *in vitro* against cultured renal cell carcinoma cells. (see: Knoll et al., Targeted therapy of experimental renal cell carcinoma with a novel conjugate of monoclonal antibody 138H11 and calicheamicin $\theta_1^I$, *Cancer Res*, **60***:* 6089-6094 (2000) The same linkage may be utilized to link this cytotoxic agent to an antibody of the present invention, thereby forming a targeting structure for kidney cancer cells.

Also useful in forming the immunoconjugates of the invention is DC1, a disulfide-containing analog of adozelesin, that kills cells by binding to the minor groove of DNA, followed by alkylation of adenine bases. Adozelesin is a structural analog of CC-1065, an anti-tumor antibiotic isolated from microbial fermentation of *Streptomyces zelensis*, and is about 1,000 fold more toxic to cultured cell lines that other DNA interacting agents, such as cis-platin and doxorubicin. This agent is readily linked to antibodies through the disulfide bond of adozelesin. (see: Chari et al., Enhancement of the selectivity and antitumor efficacy of a CC-1065 analogue through immunoconjugate formation, *Cancer Res*, **55***:* 4079-4084 (1995)).

Maytansine, a highly cytotoxic microtubular inhibitor isolated from the shrub *Maytenus serrata* found to have little value in human clinical trials, is much more effective in its derivatized form, denoted DM1, containing a disulfide bond to facilitate linkage to antibodies, is up to 10-fold more cytotoxic (see: Chari et al., Immunoconjugates containing novel maytansinoids: promising anticancer drugs, *Cancer Res*, **52***:* 127-131 (1992)). These same *in vitro* studies showed that up to four DM1 molecules could be linked to a single

immunoglobulin without destroying the binding affinity. Such conjugates have been used against breast cancer antigens, such as the *neu*/HER2/*erbB-2* antigen. (see: Goldmacher et al., Immunogen, Inc., (2002) *in press*); also see Liu, C. et al., Eradication of large colon tumor xenografts by targeted delivery of maytansinoids, *Proc. Natl. Acad. Sci. USA*, **93**, 8618-8623 (1996)). For example, Liu et al. (1996) describes formation of an immunoconjugate of the maytansinoid cytotoxin DM1 and C242 antibody, a murine IgG1 immunoglobulin, available from Pharmacia and which has affinity for a mucin-like glycoprotein variably expressed by human colorectal cancers. The latter immunoconjugate was prepared according to Chari et al., *Cancer Res.*, **52**:127-131 (1992) and was found to be highly cytotoxic against cultured colon cancer cells as well as showing anti-tumor effects *in vivo* in mice bearing subcutaneous COLO 205 human colon tumor xenografts using doses well below the maximum tolerated dose.

In addition, there are a variety of protein toxins (cytotoxic proteins), which include a number of different classes, such as those that inhibit protein synthesis: ribosome-inactivating proteins of plant origin, such as ricin, abrin, gelonin, and a number of others, and bacterial toxins such as pseudomonas exotoxin and diphtheria toxin.

Another useful class is the one including taxol, taxotere, and taxoids. Specific examples include paclitaxel (taxol), its analog docetaxel (taxotere), and derivatives thereof. The first two are clinical drugs used in treating a number of tumors while the taxoids act to induce cell death by inhibiting the de-polymerization of tubulin. Such agents are readily linked to antibodies through disulfide bonds without disadvantageous effects on binding specificity.

In one instance, a truncated Pseudomonas exotoxin was fused to an anti-CD22 variable fragment and used successfully to treat patients with chemotherapy-resistant hairy-cell leukemia. (see: Kreitman et al., Efficacy of

the anti-CD22 recombinant immunotoxin BL22 in chemotherapy-resistant hairy-cell leukemia, *N Engl J Med*, **345**: 241-247 (2001)) Conversely, the cancer-linked peptides of the present invention offer the opportunity to prepare antibodies, recombinant or otherwise, against the appropriate antigens to target solid tumors, preferably those of malignancies of kidney tissue, using the same or similar cytotoxic conjugates. Thus, many of the previously used immunoconjugates have been formed using antibodies against general antigenic sites linked to cancers whereas the antibodies formed using the peptides disclosed herein are more specific and target the antibody-cytotoxic agent to a particular tissue or organ, thus further reducing toxicity and other undesirable side effects.

In addition, the immunoconjugates formed using the antibodies prepared against the cancer-linked antigens disclosed herein can be formed by any type of chemical coupling. Thus, the cytotoxic agent of choice, along with the immunoglobulin, can be coupled by any type of chemical linkage, covalent or non-covalent, including electrostatic linkage, to form the immunoconjugates of the present invention.

When used as immunoconjugates, the antitumor agents of the present invention represent a class of pro-drugs that are relatively non-toxic when first administered to an animal (due mostly to the stability of the immunoconjugate), such as a human patient, but which are targeted by the conjugated immunoglobulin to a cancer cell where they then exhibit good toxicity. The tumor-related, associated, or linked, antigens, preferably those presented herein, serve as targets for the antibodies (monoclonal, recombinant, and the like) specific for said antigens. The end result is the release of active cytotoxic agent inside the cell after binding of the immunoglobulin portion of the immunoconjugate.

The cited references describe a number of useful procedures for the chemical linkage of cytotoxic agents to immunoglobulins and the disclosures

of all such references cited herein are hereby incorporated by reference in their entirety. For other reviews see Ghetie et al., Immunotoxins in the therapy of cancer: from bench to clinic, *Pharmacol Ther*, **63**: 209-234 (1994), Pietersz et al. The use of monoclonal antibody immunoconjugates in cancer therapy, *Adv Exp Med Biol*, **353**:169-179 (1994), and Pietersz, G. A. The linkage of cytotoxic drugs to monoclonal antibodies for the treatment of cancer, *Bioconjug Chem*, **1**:89-95 (1990).

Thus, the present invention provides highly useful cancer-associated antigens for generation of antibodies for linkage to a number of different cytotoxic agents which are already known to have some *in vitro* toxicity and possess chemical groups available for linkage to antibodies.

The present invention also relates to a process that comprises a method for producing a product, such as test data, comprising identifying an agent according to one of the disclosed processes for identifying such an agent (i.e., the therapeutic agents identified according to the assay procedures disclosed herein) wherein said product is the data collected with respect to said agent as a result of said identification process, or assay, and wherein said data is sufficient to convey the chemical character and/or structure and/or properties of said agent. For example, the present invention specifically contemplates a situation whereby a user of an assay of the invention may use the assay to screen for compounds having the desired enzyme modulating activity and, having identified the compound, then conveys that information (i.e., information as to structure, dosage, etc) to another user who then utilizes the information to reproduce the agent and administer it for therapeutic or research purposes according to the invention. For example, the user of the assay (user 1) may screen a number of test compounds without knowing the structure or identity of the compounds (such as where a number of code numbers are used the first user is simply given samples labeled with said code numbers) and, after performing the screening process, using one or more assay processes of the present invention, then

imparts to a second user (user 2), verbally or in writing or some equivalent fashion, sufficient information to identify the compounds having a particular modulating activity (for example, the code number with the corresponding results). This transmission of information from user 1 to user 2 is specifically contemplated by the present invention.

It should be cautioned that, in carrying out the procedures of the present invention as disclosed herein, whether to form immunoconjugates or screen for other antitumor agents using the genes and polypeptides disclosed herein, any reference to particular buffers, media, reagents, cells, culture conditions and the like are not intended to be limiting, but are to be read so as to include all related materials that one of ordinary skill in the art would recognize as being of interest or value in the particular context in which that discussion is presented. For example, it is often possible to substitute one buffer system or culture medium for another and still achieve similar, if not identical, results. Those of skill in the art will have sufficient knowledge of such systems and methodologies so as to be able, without undue experimentation, to make such substitutions as will optimally serve their purposes in using the methods and procedures disclosed herein.

The present invention will now be further described by way of the following non-limiting example. In applying the disclosure of the example, it should be kept clearly in mind that other and different embodiments of the methods disclosed according to the present invention will no doubt suggest themselves to those of skill in the relevant art. The following example shows how a potential anti-neoplastic agent may be identified using one or more of the genes disclosed herein.

## EXAMPLE

### Determination of Gene Inhibitory Activity of an Anti-neoplastic Agent

SW480 cells are grown to a density of $10^5$ cells/$cm^2$ in Leibovitz's L-15 medium supplemented with 2 mM L-glutamine (90%) and 10% fetal bovine serum. The cells are collected after treatment with 0.25% trypsin, 0.02% EDTA at 37°C for 2 to 5 minutes. The trypsinized cells are then diluted with 30 ml growth medium and plated at a density of 50,000 cells per well in a 96 well plate (100 μl/well). The following day, cells are treated with either compound buffer alone, or compound buffer containing a chemical agent to be tested, for 24 hours. The media is then removed, the cells lysed and the RNA recovered using the RNAeasy reagents and protocol obtained from Qiagen. RNA is quantitated and 10 ng of sample in 1 μl are added to 24 μl of Taqman reaction mix containing 1X PCR buffer, RNAsin, reverse transcriptase, nucleoside triphosphates, amplitaq gold, tween 20, glycerol, bovine serum albumin (BSA) and specific PCR primers and probes for a reference gene (18S RNA) and a test gene (Gene X). Reverse transcription is then carried out at 48°C for 30 minutes. The sample is then applied to a Perlin Elmer 7700 sequence detector and heat denatured for 10 minutes at 95°C. Amplification is performed through 40 cycles using 15 seconds annealing at 60°C followed by a 60 second extension at 72°C and 30 second denaturation at 95°C. Data files are then captured and the data analyzed with the appropriate baseline windows and thresholds.

The quantitative difference between the target and reference gene is then calculated and a relative expression value determined for all of the samples used. In this way, the ability of a chemotherapeutic agent to effectively and selectively reduce the activity of a cancer-specific gene is readily ascertained. The overall expression of the cancer-specific gene, as modulated by one chemical agent relative to another, is also determined.

Chemical agents having the most effect in reducing gene activity are thereby identified as the most anti-neoplastic.

References:

Walter A. Blättler and Ravi Chari: Drugs to enhance the therapeutic potency of anti-cancer antibodies: antibody-drug conjugates as tumor-activated prodrugs. In *Anticancer Agents - Frontiers in Cancer Chemotherapy* (Iwao Ojima, Gregory D. Vite, Karl-Heinz Altmann, Eds.), American Chemical Society, pp. 317-338 (2001).

Dan L. Longo, Patricia L. Duffey, John G. Gribben, Elaine S. Jaffe, Brendan D. Curti, Barry L. Gause, John E. Janik, Virginia M. Braman, Dixie Esseltine, Wyndham H. Wilson, Dwight Kaufman, Robert E. Wittes, Lee M. Nadler, and Walter J. Urba: Combination chemotherapy followed by an Immunotoxin (Anti-B4-blocked Ricin) in patients with indolent lymphoma: results of a Phase II study. *Cancer J. 6,* 146-150 (2000).

Walter A. Blättler and John M. Lambert: Preclinical immunotoxin development. In *Monoclonal Antibody-Based Therapy of Cancer* (M. Grossbard, Ed.), Marcel Dekker, Inc. NY, NY, pp. 1-22 (1998).

Ravi V. J. Chari: Targeted delivery of chemotherapeutics: tumor-activated prodrug therapy. In *Advanced Drug Delivery Reviews,* Elsevier Science B.V., pp. 89-104 (1998).

David T. Scadden, David P. Schenkein, Zale Bernstein, Barry Luskey, John Doweiko, Anil Tulpule, and Alexandra M. Levine: Immunotoxin combined with chemotherapy for patients with AIDS-related Non-Hodgkin's Lymphoma. *Cancer 83,* 2580-2587 (1998).

Changnian Liu and Ravi VJ Chari: The development of antibody delivery systems to target cancer with highly potent maytansinoids. *Exp. Opi. Invest. Drugs 6*, 169-172 (1997).

A. C. Goulet, Viktor S. Goldmacher, John M. Lambert, C. Baron, Dennis C. Roy and E. Kouassi: Conjugation of blocked ricin to an anti-CD19 monoclonal antibody increases antibody-induced cell calcium mobilization and CD19 internalization. *Blood 90,* 2364-2375 (1997).

Changnian Liu, John M. Lambert, Beverly A. Teicher, Walter A. Blättler, and Rosemary O'Connor: Cure of multidrug-resistant human B-cell lymphoma xenografts by combinations of anti-B4-blocked ricin and chemotherapeutic drugs. *Blood 87,* 3892-3898 (1996).

Rajeeva Singh, Lana Kats, Walter A. Blättler, and John M. Lambert: Formation of N-Substituted 2-Iminothiolanes when amino groups in proteins and peptides are modified by 2-Iminothiolane. *Anal. Biochem. 236,* 114-125 (1996).

Changnian Liu, B. Mitra Tadayoni, Lizabeth A. Bourret, Kristin M. Mattocks, Susan M. Derr, Wayne C. Widdison, Nancy L. Kedersha, Pamela D. Ariniello, Victor S. Goldmacher, John M. Lambert, Walter A. Blättler, and Ravi V.J. Chari: Eradication of large colon tumor xenografts by targeted delivery of maytansinoids. *Proc. Natl. Acad. Sci. USA 93,* 8618-8623 (1996).

Denis C. Roy, Sophie Ouellet, Christiane Le Houiller, Pamela D. Ariniello, Claude Perreault and John M. Lambert: Elimination of neuroblastoma and small-cell lung cancer cells with an anti-neural cell adhesion molecule immunotoxin. *J. Natl. Cancer Inst. 88,* 1136-1145 (1996).

Walter A. Blättler, Ravi V.J. Chari and John M. Lambert: Immunoconjugates. In Cancer Therapeutics: Experimental and Clinical Agents. (B. Teicher, Ed.), Humana Press, Totowa, NJ, pp. 371-394 (1996).

Michael L Grossbard, John M. Lambert, Victor S. Goldmacher, Arnold S. Freedman, Jeanne Kinsella, Danny P. Ducello, Susan N. Rabinowe, Laura Elisea, Felice Carol, James A. Taylor, Walter A. Blättler, Carol L. Epstein, and Lee M. Nadler: Anti-B4-blocked Ricin: A phase I trial of 7 day continuous infusion in patients with B-cell neoplasms. *J. Clin. Oncol. 11,* 726-737 (1993).

Michael L. Grossbard, John G. Gribben, Arnold S. Freedman, John M. Lambert, Jeanne Kinsella, Susan N. Rabinowe, Laura Eliseo, James A. Taylor, Walter A. Blättler, Carol L. Epstein, and Lee M. Nadler: Adjuvant immunotoxin therapy with anti-B4-blocked ricin following autologous bone marrow transplantation for patients with B-cell Non-Hodgkin's lymphoma. *Blood 81,* 2263-2271 (1993).

Sudhir A. Shah, Patricia M. Halloran, Cynthia A. Ferris, Beth A. Levine, Lizabeth A. Bourret, Victor S. Goldmacher, and Walter A. Blättler: Anti-B4-blocked Ricin immunotoxin shows therapeutic efficacy in four different SCID mouse tumor models. *Cancer Res. 53,* 1360-1367 (1993).

Ravi V.J. Chari, Bridget A. Martell, Jonathan L. Gross, Sherilyn B. Cook, Sudhir A. Shah, Walter A. Blättler, Sara J. McKenzie, and Victor S. Goldmacher: Immunoconjugates containing novel maytansinoids: promising anti-cancer drugs. *Cancer Res. 52,* 127-131 (1992).

John M. Lambert, Peter D. Senter, Annie Yau-Young, Walter A. Blättler, and Victor S. Goldmacher: Purified immunotoxins that are reactive with human lymphoid cells. *J. Biol. Chem. 250,* 12035-12041 (1985).

WHAT IS CLAIMED IS:

1. A process for identifying an agent that modulates the activity of a cancer-related gene comprising:

(a) contacting a compound with a cell containing a gene that corresponds to a polynucleotide having a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33 and under conditions promoting the expression of said gene; and

(b) detecting a difference in expression of said gene relative to when said compound is not present

thereby identifying an agent that modulates the activity of a cancer-related gene.

2 The process of claim 1 wherein said gene has a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33.

3. The process of claim 2 wherein the cell is a cancer cell and the difference in expression is a decrease in expression.

4. The process of claim 3 wherein said cancer cell is a kidney cancer cell.

5. A process for identifying an anti-neoplastic agent comprising contacting a cell exhibiting neoplastic activity with a compound first identified as a cancer related gene modulator using the process of one of claim 1 and detecting a decrease in said neoplastic activity after said contacting compared to when said contacting does not occur.

6. The process of claim 5 wherein said neoplastic activity is accelerated cellular replication.

7. The process of claim 5 wherein said decrease in neoplastic activity results from the death of the cell.

8. A process for identifying an anti-neoplastic agent comprising administering to an animal exhibiting a cancer condition an effective amount of an agent first identified according to the process of one of claim 1 and detecting a decrease in said cancerous condition.

9. A process for determining the cancerous status of a cell, comprising determining an increase in the level of expression in said cell of a gene that corresponds to a polynucleotide having a sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33 wherein an elevated expression relative to a known non-cancerous cell indicates a cancerous state or potentially cancerous state.

10. The process of claim 9 wherein said elevated expression is due to an increased copy number.

11. An isolated polypeptide comprising an amino acid sequence homologous to an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39 wherein any difference between said amino acid sequence and the sequence of SEQ ID NO: 8-13, 21-26 and 34-39 is due solely to conservative amino acid substitutions and wherein said isolated polypeptide comprises at least one immunogenic fragment.

12. An isolated polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39.

13. An antibody that reacts with a polypeptide comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 8-13, 21-26 and 34-39.

14. The antibody of claim 13 wherein said antibody is a recombinant antibody.

15. The antibody of claim 13 wherein said antibody is a synthetic antibody.

16. The antibody of claim 13 wherein said antibody is a humanized antibody.

17. An immunoconjugate comprising the antibody of claim 13 and a cytotoxic agent.

18. The antibody of claim 17 wherein said cytotoxic agent is a member selected from the group consisting of a calicheamicin, a maytansinoid, an adozelesin, a cytotoxic protein, a taxol, a taxotere, a taxoid and DC1.

19. The immunoconjugate of claim 18 wherein said calicheamicin is calicheamicin $\gamma_1^I$, N-acetyl gamma calicheamicin dimethyl hydrazide or calicheamicin $\theta_1^I$.

20. The immunoconjugate of claim 18 wherein said maytansinoid is DM1.

21. The immunoconjugate of claim 18 wherein said cytotoxic protein is ricin, abrin, gelonin, pseudomonas exotoxin or diphtheria toxin.

22. The immunoconjugate of claim 18 wherein said taxol is paclitaxel.

23. The immunoconjugate of claim 18 wherein said taxotere is docetaxel.

24. A process for treating cancer comprising contacting a cancerous cell *in vivo* with an agent having activity against an expression product encoded by a gene sequence selected from the group consisting of SEQ ID NO: 1-7, 14-20 and 27-33.

25. The process of claim 24 wherein said agent is the antibody of claim 13.

26. The process of claim 24 wherein said agent is an immunoconjugate of claim 17.

27. An immunogenic composition comprising a polypeptide of claim 11.

28. An immunogenic composition comprising a polypeptide of claim 12.

29. The process of claim 24 wherein said cancer is kidney cancer.

30. A process for treating cancer in an animal afflicted therewith comprising administering to said animal an amount of an immunogenic composition of claim 27 sufficient to elicit the production of cytotoxic T lymphocytes specific for the polypeptide of claim 11.

31. A process for treating cancer in an animal afflicted therewith comprising administering to said animal an amount of an immunogenic composition of claim 28 sufficient to elicit the production of cytotoxic T lymphocytes specific for the polypeptide of claim 12.

32. A process for treating a cancerous condition in an animal afflicted therewith comprising administering to said animal a therapeutically effective

amount of an agent first identified as having anti-neoplastic activity using the process of claim 8.

33. A process for protecting an animal against cancer comprising administering to an animal at risk of developing cancer a therapeutically effective amount of an agent first identified as having anti-neoplastic activity using the process of claim 8.

34. The process of claim 30 wherein said animal is a human being.

35. The process of claim 30 wherein said cancer is kidney cancer.

36. A method for producing test data with respect to the gene modulating activity of a compound comprising:

(a) contacting a compound with one or more cells containing a polynucleotide comprising a nucleotide sequence corresponding to a gene whose expression is increased in a cancerous cell over that in a non-cancerous cell or a gene whose expression is elevated in a non-cancerous cell over that in a cancerous cell under conditions wherein said polynucleotide is being expressed, and

(b) determining a change in expression of more than one of said polynucleotides, and

(c) producing test data with respect to the gene modulating activity of said compound based on an increase in the expression of the determined genes whose expression is otherwise elevated in a non-cancerous cell over that in a cancerous cell and a decrease in the expression of the determined genes whose expression is otherwise increased in a cancerous cell over that in a non-cancerous cell indicating gene modulating activity.

SEQUENCE LISTING

<110> Avalon Pharmaceuticals

<120> Cancer-Linked Gene as Target for Chemotherapy

<130> 689290-165

<140>
<141>

<150> US/60/385,505
<151> 2002-06-04

<160> 39

<170> PatentIn version 3.0

<210> 1
<211> 4567
<212> DNA
<213> Homo sapiens

<400> 1
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc 300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg 360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt 420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa 480
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg 540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc 600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc 660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg 720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc 780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc 840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca 900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg 960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca 1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac 1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt ggggggccga 1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg 1200
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac 1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc 1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc 1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct 1440
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc 1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg 1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc 1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc 1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc 1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg 1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg 1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc 1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat 1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca 2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt 2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg 2160

tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga 2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag 2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc 2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc 2400
gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc 2460
tgccccgaag tgcggcgcaa cacccccctgg acgccgtggc tgcccgtgaa cgtgacgcag 2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgccccct tgcagacccg 2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc 2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg 2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc 2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc 2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt 2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt 2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt 3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg 3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt 3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc 3180
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt 3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg 3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc 3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag 3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat 3480
gacagagcca acttctaccc attgcagcag accaatgtgt acacgactac ttactaccca 3540
agccccctga acaaacacag cttccggccc gaggcctcac ctggacaacg gtgcttcccc 3600
aacagctgat accgccgtcc tggggacttg ggcttcttgc cttcataagg cacagagcag 3660
atggagatgg gacagtggag ccagtttggt tttctccctc tgcactaggc caagaacttg 3720
ctgccttgcc tgtgggggggt cccatccggc ttcagagagc tctggctggc attgaccatg 3780
ggggaaaggg ctggtttcag gctgacatat ggccgcaggt ccagttcagc ccaggtctmt 3840
catggttatc ttccaaccca ctgtcacgct gacactatgc tgccatgcct gggctgtgga 3900
cctactgggc atttgaggaa ytggagaatg gagatggcaa gagggcaggc ttttaagttt 3960
gggttggaga caacttcctg tggcccccac aagctgagtc tggccttctc cagctggccc 4020
caaaaaaggc ctttgctaca tcctgattat ctctgaaagt aatcaatcaa gtggctccag 4080
tagctctgga ttttctgcca gggctgggcc attgtggtgc tgccccagta tgacatggga 4140
ccaaggccag cgcaggttat ccacctctgc ctggaagtct atactctacc cagggcatcc 4200
ctctggtcag aggcagtgag tactgggaac tggaggctga cctgtgctta gaagtccttt 4260
aatctgggct ggtacaggcc tcagccttgc cctcaatgca cgaaaggtgg cccaggagag 4320
aggatcaatg ccataggagg cagaagtctg gcctctgtgc ctctatggag actatcttcc 4380
agttgctgct caacagagtt gttggctgag acctgcttgg gagtctctgc tggcccttca 4440
tctgttcagg aacacacaca cacacacact cacacacgca cacacaatca caatttgcta 4500
cagcaacaaa aaagacattg ggctgtggca ttattaatta aagatgatat ccagtcaaaa 4560
aaaaact 4567

<210> 2
<211> 453
<212> DNA
<213> Homo sapiens

<400> 2
agtaatcagc tcggtaccgg catgtgctgt agccagcgca ggttatccac ctctgcctgg 60
aagtctatac tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga 120
ggctgacctg tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc 180
aatgcacgaa aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct 240
ctgtgcctct atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct 300
gcttgggagt ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca 360
cacgcacaca caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat 420
taattaaaga tgatatccag tcaaaaaaaa act 453

<210> 3
<211> 4675

<212> DNA
<213> Homo sapiens

<400> 3
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggca ggaagccaag tggagacttt 300
gaatggaggc aaggatggag gggacctggg gaagaggact ggcctgaatc accttcccca 360
aaggtcctca tggactgagc tggaggtctt ctcccctgtc tgcctcccgg agctaggact 420
gcagaggggc ctatcatggt gcttgcaggc cccctggctg tctcgctgtt gctgcccagc 480
ctcacactgc tggtgtccca cctctccagc tcccaggatg tctccagtga gcccagcagt 540
gagcagcagc tgtgcgccct tagcaagcac cccaccgtgg cctttgaaga cctgcagccg 600
tgggtctcta acttcaccta ccctggagcc cgggatttct cccagctggc tttggacccc 660
tccgggaacc agctcatcgt gggagccagg aactacctct tcagactcag ccttgccaat 720
gtctctcttc ttcaggccac agagtgggcc tccagtgagg acacgcgccg ctcctgccaa 780
agcaaaggga agactgagga ggagtgtcag aactacgtgc gagtcctgat cgtcgccggc 840
cggaaggtgt tcatgtgtgg aaccaatgcc ttttccccca tgtgcaccag cagacaggtg 900
gggaacctca gccggactac tgagaagatc aatggtgtgg cccgctgccc ctatgaccca 960
cgccacaact ccacagctgt catctcctcc caggggagc tctatgcagc cacggtcatc 1020
gacttctcag gtcgggaccc tgccatctac cgcagcctgg gcagtgggcc accgcttcgc 1080
actgcccaat ataactccaa gtggcttaat gagccaaact tcgtggcagc ctatgatatt 1140
gggctgtttg catacttctt cctgcgggag aacgcagtgg agcacgactg tggacgcacc 1200
gtgtactctc gcgtggcccg cgtgtgcaag aatgacgtgg ggggccgatt cctgctggag 1260
gacacatgga ccacattcat gaaggcccgg ctcaactgct cccgcccggg cgaggtcccc 1320
ttctactata acgagctgca gagtgccttc cacttgccag agcaggacct catctatgga 1380
gttttcacaa ccaacgtaaa cagcatcgcg gcttctgctg tctgcgcctt caacctcagt 1440
gctatctccc aggctttcaa tggcccattt cgctaccagg agaaccccag ggctgcctgg 1500
ctccccatag ccaaccccat ccccaatttc cagtgtggca ccctgcctga gaccggtccc 1560
aacgagaacc tgacggagcg cagcctgcag gacgcgcagc gcctcttcct gatgagcgag 1620
gccgtgcagc cggtgacacc cgagccctgt gtcacccagg acagcgtgcg cttctcacac 1680
ctcgtggtgg acctggtgca ggctaaagac acgctctacc atgtactcta cattggcacc 1740
gagtcgggca ccatcctgaa ggcgctgtcc acggcgagcc gcagcctcca cggctgctac 1800
ctggaggagc tgcacgtgct gccccccggg cgccgcgagc ccctgcgcag cctgcgcatc 1860
ctgcacagcg cccgcgcgct cttcgtgggg ctgagagacg gcgtcctgcg ggtcccactg 1920
gagaggtgcg ccgcctaccg cagccagggg gcatgcctgg gggcccggga cccgtactgt 1980
ggctgggacg ggaagcagca acgttgcagc acactcgagg acagctccaa catgagcctc 2040
tggacccaga acatcaccgc ctgtcctgtg cggaatgtga cacgggatgg gggcttcggc 2100
ccatggtcac catggcaacc atgtgagcac ttggatgggg acaactcagg ctcttgcctg 2160
tgtcgagctc gatcctgtga ttcccctcga ccccgctgtg ggggccttga ctgcctgggg 2220
ccagccatcc acatcgccaa ctgctccagg aatggggcgt ggaccccgtg gtcatcgtgg 2280
gcgctgtgca gcacgtcctg tggcatcggc ttccaggtcc gccagcgaag ttgcagcaac 2340
cctgctcccc gccacggggg ccgcatctgc gtgggcaaga gccgggagga acggttctgt 2400
aatgagaaca cgccttgccc ggtgcccatc ttctgggctt cctggggctc ctggagcaag 2460
tgcagcagca actgtggagg gggcatgcag tcgcggcgtc gggcctgcga gaacggcaac 2520
tcctgcctgg gctgcggcgt ggagttcaag acgtgcaacc ccgagggctg ccccgaagtg 2580
cggcgcaaca ccccctggac gccgtggctg cccgtgaacg tgacgcaggg cggggcacgg 2640
caggagcagc ggttccgctt cacctgccgc gcgccccttg cagacccgca cggcctgcag 2700
ttcggcagga gaaggaccga gacgaggacc tgtcccgcgg acggctccgg ctcctgcgac 2760
accgacgccc tggtggaggt cctcctgcgc agcgggagca cctccccgca cacggtgagc 2820
gggggctggg ccgcctgggg cccgtggtcg tcctgctccc gggactgcga gctgggcttc 2880
cgcgtccgca agagaacgtg cactaacccg gagccccgca acgggggcct gccctgcgtg 2940
ggcgatgctg ccgagtacca ggactgcaac ccccaggctt gcccagttcg gggtgcttgg 3000
tcctgctgga cctcatggtc tccatgctca gcttcctgtg gtgggggtca ctatcaacgc 3060
acccgttcct gcaccagccc cgcaccctcc ccaggtgagg acatctgtct cgggctgcac 3120
acggaggagg cactatgtgc cacacaggcc tgcccagaag gctggtcgcc ctggtctgag 3180
tggagtaagt gcactgacga cggagcccag agccgaagcc ggcactgtga ggagctcctc 3240
ccagggtcca gcgcmtgtgc tggaaacagc agccagagcc gcccctgccc ctacagcgag 3300
attcscgtca tcctgccagc ctccagcatg gaggaggcca ccgactgtgc agggttcaat 3360
ctcatccact tggtggccac gggcatctcc tgcttcttgg gctctgggct cctgacccta 3420

gcagtgtacc tgtcttgcca gcactgccag cgtcagtccc aggagtccac actggtccat 3480
cctgccaccc ccaaccattt gcactacaag ggcggaggca ccccgaagaa tgaaaagtac 3540
acacccatgg aattcaagac cctgaacaag aataacttga tccctgatga cagagccaac 3600
ttctacccat tgcagcagac caatgtgtac acgactactt actacccaag ccccctgaac 3660
aaacacagct tccggcccga ggcctcacct ggacaacggt gcttccccaa cagctgatac 3720
cgccgtcctg gggacttggg cttcttgcct tcataaggca cagagcagat ggagatggga 3780
cagtggagcc agtttggttt tctccctctg cactaggcca agaacttgct gccttgcctg 3840
tgggggtcc catccggctt cagagagctc tggctggcat tgaccatggg ggaaagggct 3900
ggtttcaggc tgacatatgg ccgcaggtcc agttcagccc aggtctmtca tggttatctt 3960
ccaacccact gtcacgctga cactatgctg ccatgcctgg gctgtggacc tactgggcat 4020
ttgaggaayt ggagaatgga gatggcaaga gggcaggctt ttaagtttgg gttggagaca 4080
acttcctgtg gcccccacaa gctgagtctg gccttctcca gctggcccca aaaaaggcct 4140
ttgctacatc ctgattatct ctgaaagtaa tcaatcaagt ggctccagta gctctggatt 4200
ttctgccagg gctgggccat tgtggtgctg ccccagtatg acatgggacc aaggccagcg 4260
caggttatcc acctctgcct ggaagtctat actctaccca gggcatccct ctggtcagag 4320
gcagtgagta ctgggaactg gaggctgacc tgtgcttaga agtcctttaa tctgggctgg 4380
tacaggcctc agccttgccc tcaatgcacg aaaggtggcc caggagagag gatcaatgcc 4440
ataggaggca gaagtctggc ctctgtgcct ctatggagac tatcttccag ttgctgctca 4500
acagagttgt tggctgagac ctgcttggga gtctctgctg gcccttcatc tgttcaggaa 4560
cacacacaca cacacactca cacacgcaca cacaatcaca atttgctaca gcaacaaaaa 4620
agacattggg ctgtggcatt attaattaaa gatgatatcc agtcaaaaaa aaact 4675

<210> 4
<211> 4731
<212> DNA
<213> Homo sapiens

<400> 4
attggagatg ctcgggggca ggctgccgcg ttgtgtcctg cttttctgcg gccagaccaa 60
gccgtctgga gctgctggtc aggttttctt gctgacctca cctgaccaca gtggcctggg 120
tggactctac agggaaatgt tgttttctcc ctgggagcag tagcagcagt cctggctccc 180
ctggactgag aactcctcat cagccccagg aagcccggac ccccttttcag ggatctggaa 240
ccggtgtgcc tgtggcccca ggtctgctcc caggcgtggg ctgaagtcct gacttctgtc 300
gctgggggca aggagtggga gagcccagct gctgcctggg ctttggcaga cagcaggctg 360
atggtgctgg cttccccgag actgcttctc ctgcctgctg tctgatttcc ctgcatggtg 420
cccgcagctg agctgctacg ggtcttctcc cctgtctgcc tcccggagct aggactgcag 480
aggggcctat catggtgctt gcaggccccc tggctgtctc gctgttgctg cccagcctca 540
cactgctggt gtcccacctc tccagctccc aggatgtctc cagtgagccc agcagtgagc 600
agcagctgtg cgcccttagc aagcacccca ccgtggcctt tgaagacctg cagccgtggg 660
tctctaactt cacctaccct ggagcccggg atttctccca gctggctttg gacccctccg 720
ggaaccagct catcgtggga gccaggaact acctcttcag actcagcctt gccaatgtct 780
ctcttcttca ggccacagag tgggcctcca gtgaggacac gcgccgctcc tgccaaagca 840
aagggaagac tgaggaggag tgtcagaact acgtgcgagt cctgatcgtc gccggccgga 900
aggtgttcat gtgtggaacc aatgcctttt cccccatgtg caccagcaga caggtgggga 960
acctcagccg gactactgag aagatcaatg gtgtggcccg ctgcccctat gacccacgcc 1020
acaactccac agctgtcatc tcctcccagg gggagctcta tgcagccacg gtcatcgact 1080
tctcaggtcg ggaccctgcc atctaccgca gcctgggcag tgggccaccg cttcgcactg 1140
cccaatataa ctccaagtgg cttaatgagc caaacttcgt ggcagcctat gatattgggc 1200
tgtttgcata cttcttcctg cgggagaacg cagtggagca cgactgtgga cgcaccgtgt 1260
actctcgcgt ggcccgcgtg tgcaagaatg acgtgggggg ccgattcctg ctggaggaca 1320
catggaccac attcatgaag gcccggctca actgctcccg cccgggcgag gtccccttct 1380
actataacga gctgcagagt gccttccact tgccagagca ggacctcatc tatggagttt 1440
tcacaaccaa cgtaaacagc atcgcggctt ctgctgtctg cgccttcaac ctcagtgcta 1500
tctcccaggc tttcaatggc ccatttcgct accaggagaa ccccagggct gcctggctcc 1560
ccatagccaa ccccatcccc aatttccagt gtggcaccct gcctgagacc ggtcccaacg 1620
agaacctgac ggagcgcagc ctgcaggacg cgcagcgcct cttcctgatg agcgaggccg 1680
tgcagccggt gacacccgag ccctgtgtca cccaggacag cgtgcgcttc tcacacctcg 1740
tggtggacct ggtgcaggct aaagacacgc tctaccatgt actctacatt ggcaccgagt 1800
cgggcaccat cctgaaggcg ctgtccacgg cgagccgcag cctccacggc tgctacctgg 1860
aggagctgca cgtgctgccc cccgggcgcc gcgagcccct gcgcagcctg cgcatcctgc 1920

acagcgcccg cgcgctcttc gtggggctga gagacggcgt cctgcgggtc ccactggaga 1980
ggtgcgccgc ctaccgcagc caggggggcat gcctgggggc ccgggacccg tactgtggct 2040
gggacgggaa gcagcaacgt tgcagcacac tcgaggacag ctccaacatg agcctctgga 2100
cccagaacat caccgcctgt cctgtgcgga atgtgacacg ggatgggggc ttcggcccat 2160
ggtcaccatg gcaaccatgt gagcacttgg atggggacaa ctcaggctct tgcctgtgtc 2220
gagctcgatc ctgtgattcc cctcgacccc gctgtggggg ccttgactgc ctggggccag 2280
ccatccacat cgccaactgc tccaggaatg gggcgtggac cccgtggtca tcgtgggcgc 2340
tgtgcagcac gtcctgtggc atcggcttcc aggtccgcca gcgaagttgc agcaaccctg 2400
ctccccgcca cgggggccgc atctgcgtgg gcaagagccg ggaggaacgg ttctgtaatg 2460
agaacacgcc ttgcccggtg cccatcttct gggcttcctg gggctcctgg agcaagtgca 2520
gcagcaactg tggaggggggc atgcagtcgc ggcgtcgggc ctgcgagaac ggcaactcct 2580
gcctgggctg cggcgtggag ttcaagacgt gcaaccccga gggctgcccc gaagtgcggc 2640
gcaacacccc ctggacgccg tggctgcccg tgaacgtgac gcagggcggg gcacggcagg 2700
agcagcggtt ccgcttcacc tgccgcgcgc cccttgcaga cccgcacggc ctgcagttcg 2760
gcaggagaag gaccgagacg aggacctgtc ccgcggacgg ctccggctcc tgcgacaccg 2820
acgccctggt ggaggtcctc ctgcgcagcg ggagcacctc cccgcacacg gtgagcgggg 2880
gctgggccgc ctggggcccg tggtcgtcct gctcccggga ctgcgagctg ggcttccgcg 2940
tccgcaagag aacgtgcact aacccggagc cccgcaacgg gggcctgccc tgcgtgggcg 3000
atgctgccga gtaccaggac tgcaaccccc aggcttgccc agttcggggt gcttggtcct 3060
gctggacctc atggtctcca tgctcagctt cctgtggtgg gggtcactat caacgcaccc 3120
gttcctgcac cagccccgca ccctccccag gtgaggacat ctgtctcggg ctgcacacgg 3180
aggaggcact atgtgccaca caggcctgcc cagaaggctg gtcgccctgg tctgagtgga 3240
gtaagtgcac tgacgacgga gcccagagcc gaagccggca ctgtgaggag ctcctcccag 3300
ggtccagcgc mtgtgctgga aacagcagcc agagccgccc ctgcccctac agcgagattc 3360
scgtcatcct gccagcctcc agcatggagg aggccaccga ctgtgcaggg ttcaatctca 3420
tccacttggt ggccacgggc atctcctgct tcttgggctc tgggctcctg accctagcag 3480
tgtacctgtc ttgccagcac tgccagcgtc agtcccagga gtccacactg gtccatcctg 3540
ccacccccaa ccatttgcac tacaagggcg gaggcacccc gaagaatgaa aagtacacac 3600
ccatggaatt caagaccctg aacaagaata acttgatccc tgatgacaga gccaacttct 3660
acccattgca gcagaccaat gtgtacacga ctacttacta cccaagcccc ctgaacaaac 3720
acagcttccg gcccgaggcc tcacctggac aacggtgctt ccccaacagc tgataccgcc 3780
gtcctgggga cttgggcttc ttgccttcat aaggcacaga gcagatggag atgggacagt 3840
ggagccagtt tggttttctc cctctgcact aggccaagaa cttgctgcct tgcctgtggg 3900
gggtcccatc cggcttcaga gagctctggc tggcattgac catgggggaa agggctggtt 3960
tcaggctgac atatggccgc aggtccagtt cagcccaggt ctmtcatggt tatcttccaa 4020
cccactgtca cgctgacact atgctgccat gcctgggctg tggacctact gggcatttga 4080
ggaaytggag aatggagatg gcaagagggc aggcttttaa gtttgggttg gagacaactt 4140
cctgtggccc ccacaagctg agtctggcct tctccagctg gccccaaaaa aggcctttgc 4200
tacatcctga ttatctctga aagtaatcaa tcaagtggct ccagtagctc tggattttct 4260
gccagggctg ggccattgtg gtgctgcccc agtatgacat gggaccaagg ccagcgcagg 4320
ttatccacct ctgcctggaa gtctatactc tacccagggc atccctctgg tcagaggcag 4380
tgagtactgg gaactggagg ctgacctgtg cttagaagtc ctttaatctg ggctggtaca 4440
ggcctcagcc ttgccctcaa tgcacgaaag gtggcccagg agagaggatc aatgccatag 4500
gaggcagaag tctggcctct gtgcctctat ggagactatc ttccagttgc tgctcaacag 4560
agttgttggc tgagacctgc ttgggagtct ctgctggccc ttcatctgtt caggaacaca 4620
cacacacaca cactcacaca cgcacacaca atcacaattt gctacagcaa caaaaaagac 4680
attgggctgt ggcattatta attaaagatg atatccagtc aaaaaaaaac t 4731

<210> 5
<211> 4703
<212> DNA
<213> Homo sapiens

<400> 5
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagccagc 120
ttgggtccgg gttgcactgg gccctgccag ggctgtggtc ggcgcatctg ggctgcagcg 180
gcgatgggga cccgggaccc aggcctggag aaggagacgg acgagtgagg ctgagggacg 240
gagggacaga gtgagtggtt ccagctggtg cctggcctgt gtctcttgga tgccctgtgg 300
cttcagtccg tctcctgttg cccaccacct cgtccctggg ccgcctgata ccccagccca 360

acagctaagg tgtggatgga cagtaggggg ctggcttctc tcactggtca ggggtcttct 420
cccctgtctg cctcccggag ctaggactgc agaggggcct atcatggtgc ttgcaggccc 480
cctggctgtc tcgctgttgc tgcccagcct cacactgctg gtgtcccacc tctccagctc 540
ccaggatgtc tccagtgagc ccagcagtga gcagcagctg tgcgcccttа gcaagcaccc 600
caccgtggcc tttgaagacc tgcagccgtg ggtctctaac ttcacctacc ctggagcccg 660
ggatttctcc cagctggctt tggacccctc cgggaaccag ctcatcgtgg gagccaggaa 720
ctacctcttc agactcagcc ttgccaatgt ctctcttctt caggccacag agtgggcctc 780
cagtgaggac acgcgccgct cctgccaaag caaagggaag actgaggagg agtgtcagaa 840
ctacgtgcga gtcctgatcg tcgccggccg gaaggtgttc atgtgtggaa ccaatgcctt 900
ttcccccatg tgcaccagca gacaggtggg gaacctcagc cggactactg agaagatcaa 960
tggtgtggcc cgctgccccт atgacccacg ccacaactcc acagctgtca tctcctccca 1020
gggggagctc tatgcagcca cggtcatcga cttctcaggt cgggaccctg ccatctaccg 1080
cagcctgggc agtgggccac cgcttcgcac tgcccaatat aactccaagt ggcttaatga 1140
gccaaacttc gtggcagcct atgatattgg gctgtttgca tacttcttcc tgcgggagaa 1200
cgcagtggag cacgactgtg gacgcaccgt gtactctcgc gtggcccgcg tgtgcaagaa 1260
tgacgtgggg ggccgattcc tgctggagga cacatggacc acattcatga aggcccggct 1320
caactgctcc cgcccgggcg aggtcccctt ctactataac gagctgcaga gtgccttcca 1380
cttgccagag caggacctca tctatggagt tttcacaacc aacgtaaaca gcatcgcggc 1440
ttctgctgtc tgcgccttca acctcagtgc tatctcccag gctttcaatg gcccatttcg 1500
ctaccaggag aaccccaggg ctgcctggct ccccatagcc aaccccatcc ccaatttcca 1560
gtgtggcacc ctgcctgaga ccggtcccaa cgagaacctg acggagcgca gcctgcagga 1620
cgcgcagcgc ctcttcctga tgagcgaggc cgtgcagccg gtgacacccg agccctgtgt 1680
cacccaggac agcgtgcgct tctcacacct cgtggtggac ctggtgcagg ctaaagacac 1740
gctctaccat gtactctaca ttggcaccga gtcgggcacc atcctgaagg cgctgtccac 1800
ggcgagccgc agcctccacg gctgctacct ggaggagctg cacgtgctgc cccccgggcg 1860
ccgcgagccc ctgcgcagcc tgcgcatcct gcacagcgcc cgcgcgctct tcgtggggct 1920
gagagacggc gtcctgcggg tcccactgga gaggtgcgcc gcctaccgca gccagggggc 1980
atgcctgggg gcccgggacc cgtactgtgg ctgggacggg aagcagcaac gttgcagcac 2040
actcgaggac agctccaaca tgagcctctg gacccagaac atcaccgcct gtcctgtgcg 2100
gaatgtgaca cgggatgggg gcttcggccc atggtcacca tggcaaccat gtgagcactt 2160
ggatggggac aactcaggct cttgcctgtg tcgagctcga tcctgtgatt cccctcgacc 2220
ccgctgtggg ggccttgact gcctggggcc agccatccac atcgccaact gctccaggaa 2280
tggggcgtgg accccgtggt catcgtgggc gctgtgcagc acgtcctgtg gcatcggctt 2340
ccaggtccgc cagcgaagtt gcagcaaccc tgctccccgc cacgggggcc gcatctgcgt 2400
gggcaagagc cgggaggaac ggttctgtaa tgagaacacg ccttgcccgg tgcccatctt 2460
ctgggcttcc tggggctcct ggagcaagtg cagcagcaac tgtggagggg gcatgcagtc 2520
gcggcgtcgg gcctgcgaga acggcaactc ctgcctgggc tgcggcgtgg agttcaagac 2580
gtgcaacccc gagggctgcc ccgaagtgcg gcgcaacacc ccctggacgc cgtggctgcc 2640
cgtgaacgtg acgcagggcg gggcacggca ggagcagcgg ttccgcttca cctgccgcgc 2700
gcccccttgca gacccgcacg gcctgcagtt cggcaggaga aggaccgaga cgaggacctg 2760
tcccgcggac ggctccggct cctgcgacac cgacgccctg gtggaggtcc tcctgcgcag 2820
cgggagcacc tccccgcaca cggtgagcgg gggctgggcc gcctggggcc cgtggtcgtc 2880
ctgctcccgg gactgcgagc tgggcttccg cgtccgcaag agaacgtgca ctaacccgga 2940
gccccgcaac gggggcctgc cctgcgtggg cgatgctgcc gagtaccagg actgcaaccc 3000
ccaggcttgc ccagttcggg gtgcttggtc ctgctggacc tcatggtctc catgctcagc 3060
ttcctgtggt gggggtcact atcaacgcac ccgttcctgc accagccccg caccctcccc 3120
aggtgaggac atctgtctcg ggctgcacac ggaggaggca ctatgtgcca cacaggcctg 3180
cccagaaggc tggtcgccct ggtctgagtg gagtaagtgc actgacgacg gagcccagag 3240
ccgaagccgg cactgtgagg agctcctccc agggtccagc gcmtgtgctg gaaacagcag 3300
ccagagccgc ccctgcccct acagcgagat tcscgtcatc ctgccagcct ccagcatgga 3360
ggaggccacc gactgtgcag ggttcaatct catccacttg gtggccacgg gcatctcctg 3420
cttcttgggc tctgggctcc tgaccctagc agtgtacctg tcttgccagc actgccagcg 3480
tcagtcccag gagtccacac tggtccatcc tgccaccccc aaccatttgc actacaaggg 3540
cggaggcacc ccgaagaatg aaaagtacac acccatggaa ttcaagaccc tgaacaagaa 3600
taacttgatc cctgatgaca gagccaactt ctacccattg cagcagacca atgtgtacac 3660
gactacttac tacccaagcc ccctgaacaa acacagcttc cggcccgagg cctcacctgg 3720
acaacggtgc ttccccaaca gctgataccg ccgtcctggg gacttgggct tcttgccttc 3780
ataaggcaca gagcagatgg agatgggaca gtggagccag tttggttttc tccctctgca 3840
ctaggccaag aacttgctgc cttgcctgtg gggggtccca tccggcttca gagagctctg 3900
gctggcattg accatggggg aaagggctgg tttcaggctg acatatggcc gcaggtccag 3960
ttcagcccag gtctmtcatg gttatcttcc aacccactgt cacgctgaca ctatgctgcc 4020

atgcctgggc tgtggaccta ctgggcattt gaggaaytgg agaatggaga tggcaagagg 4080
gcaggctttt aagtttgggt tggagacaac ttcctgtggc ccccacaagc tgagtctggc 4140
cttctccagc tggccccaaa aaaggccttt gctacatcct gattatctct gaaagtaatc 4200
aatcaagtgg ctccagtagc tctggatttt ctgccagggc tgggccattg tggtgctgcc 4260
ccagtatgac atgggaccaa ggccagcgca ggttatccac ctctgcctgg aagtctatac 4320
tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga ggctgacctg 4380
tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc aatgcacgaa 4440
aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct ctgtgcctct 4500
atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct gcttgggagt 4560
ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca cacgcacaca 4620
caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat taattaaaga 4680
tgatatccag tcaaaaaaaa act 4703

<210> 6
<211> 4405
<212> DNA
<213> Homo sapiens

<400> 6
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggaggtctt 120
ctcccctgtc tgcctcccgg agctaggact gcagaggggc ctatcatggt gcttgcaggc 180
cccctggctg tctcgctgtt gctgcccagc ctcacactgc tggtgtccca cctctccagc 240
tcccaggatg tctccagtga gcccagcagt gagcagcagc tgtgcgccct tagcaagcac 300
cccaccgtgg cctttgaaga cctgcagccg tgggtctcta acttcaccta ccctggagcc 360
cgggatttct cccagctggc tttggacccc tccgggaacc agctcatcgt gggagccagg 420
aactacctct tcagactcag ccttgccaat gtctctcttc ttcaggccac agagtgggcc 480
tccagtgagg acacgcgccg ctcctgccaa agcaaaggga agactgagga ggagtgtcag 540
aactacgtgc gagtcctgat cgtcgccggc cggaaggtgt tcatgtgtgg aaccaatgcc 600
ttttccccca tgtgcaccag cagacaggtg gggaacctca gccggactac tgagaagatc 660
aatggtgtgg cccgctgccc ctatgaccca cgccacaact ccacagctgt catctcctcc 720
caggggggagc tctatgcagc cacggtcatc gacttctcag gtcgggaccc tgccatctac 780
cgcagcctgg gcagtgggcc accgcttcgc actgcccaat ataactccaa gtggcttaat 840
gagccaaact tcgtggcagc ctatgatatt gggctgtttg catacttctt cctgcgggag 900
aacgcagtgg agcacgactg tggacgcacc gtgtactctc gcgtggcccg cgtgtgcaag 960
aatgacgtgg ggggccgatt cctgctggag gacacatgga ccacattcat gaaggcccgg 1020
ctcaactgct cccgcccggg cgaggtcccc ttctactata acgagctgca gagtgccttc 1080
cacttgccag agcaggacct catctatgga gttttcacaa ccaacgtaaa cagcatcgcg 1140
gcttctgctg tctgcgcctt caacctcagt gctatctccc aggctttcaa tggcccattt 1200
cgctaccagg agaaccccag ggctgcctgg ctccccatag ccaaccccat ccccaatttc 1260
cagtgtggca ccctgcctga gaccggtccc aacgagaacc tgacggagcg cagcctgcag 1320
gacgcgcagc gcctcttcct gatgagcgag gccgtgcagc cggtgacacc cgagccctgt 1380
gtcacccagg acagcgtgcg cttctcacac ctcgtggtgg acctggtgca ggctaaagac 1440
acgctctacc atgtactcta cattggcacc gagtcgggca ccatcctgaa ggcgctgtcc 1500
acggcgagcc gcagcctcca cggctgctac ctggaggagc tgcacgtgct gccccccggg 1560
cgccgcgagc ccctgcgcag cctgcgcatc ctgcacagcg cccgcgcgct cttcgtgggg 1620
ctgagagacg gcgtcctgcg ggtcccactg gagaggtgcg ccgcctaccg cagccagggg 1680
gcatgcctgg gggcccggga cccgtactgt ggctgggacg ggaagcagca acgttgcagc 1740
acactcgagg acagctccaa catgagcctc tggacccaga acatcaccgc ctgtcctgtg 1800
cggaatgtga cacgggatgg gggcttcggc ccatggtcac catggcaacc atgtgagcac 1860
ttggatgggg acaactcagg ctcttgcctg tgtcgagctc gatcctgtga ttcccctcga 1920
ccccgctgtg ggggccttga ctgcctgggg ccagccatcc acatcgccaa ctgctccagg 1980
aatggggcgt ggaccccgtg gtcatcgtgg gcgctgtgca gcacgtcctg tggcatcggc 2040
ttccaggtcc gccagcgaag ttgcagcaac cctgctcccc gccacggggg ccgcatctgc 2100
gtgggcaaga gccgggagga acggttctgt aatgagaaca cgccttgccc ggtgcccatc 2160
ttctgggctt cctggggctc ctggagcaag tgcagcagca actgtggagg gggcatgcag 2220
tcgcggcgtc gggcctgcga gaacggcaac tcctgcctgg gctgcggcgt ggagttcaag 2280
acgtgcaacc ccgagggctg ccccgaagtg cggcgcaaca ccccctggac gccgtggctg 2340
cccgtgaacg tgacgcaggg cggggcacgg caggagcagc ggttccgctt cacctgccgc 2400
gcgccccttg cagacccgca cggcctgcag ttcggcagga gaaggaccga gacgaggacc 2460

tgtcccgcgg acggctccgg ctcctgcgac accgacgccc tggtggaggt cctcctgcgc 2520
agcgggagca cctccccgca cacggtgagc gggggctggg ccgcctgggg cccgtggtcg 2580
tcctgctccc gggactgcga gctgggcttc cgcgtccgca agagaacgtg cactaacccg 2640
gagccccgca acgggggcct gccctgcgtg ggcgatgctg ccgagtacca ggactgcaac 2700
ccccaggctt gcccagttcg gggtgcttgg tcctgctgga cctcatggtc tccatgctca 2760
gcttcctgtg gtgggggtca ctatcaacgc acccgttcct gcaccagccc cgcaccctcc 2820
ccaggtgagg acatctgtct cgggctgcac acggaggagg cactatgtgc cacacaggcc 2880
tgcccagaag gctggtcgcc ctggtctgag tggagtaagt gcactgacga cggagcccag 2940
agccgaagcc ggcactgtga ggagctcctc ccagggtcca gcgcmtgtgc tggaaacagc 3000
agccagagcc gcccctgccc ctacagcgag attcscgtca tcctgccagc ctccagcatg 3060
gaggaggcca ccgactgtgc agggttcaat ctcatccact tggtggccac gggcatctcc 3120
tgcttcttgg gctctgggct cctgacccta gcagtgtacc tgtcttgcca gcactgccag 3180
cgtcagtccc aggagtccac actggtccat cctgccaccc ccaaccattt gcactacaag 3240
ggcggaggca ccccgaagaa tgaaaagtac acacccatgg aattcaagac cctgaacaag 3300
aataacttga tccctgatga cagagccaac ttctacccat tgcagcagac caatgtgtac 3360
acgactactt actacccaag ccccctgaac aaacacagct tccggcccga ggcctcacct 3420
ggacaacggt gcttccccaa cagctgatac cgccgtcctg gggacttggg cttcttgcct 3480
tcataaggca cagagcagat ggagatggga cagtggagcc agtttggttt tctccctctg 3540
cactaggcca agaacttgct gccttgcctg tgggggggtcc catccggctt cagagagctc 3600
tggctggcat tgaccatggg ggaaaggggct ggtttcaggc tgacatatgg ccgcaggtcc 3660
agttcagccc aggtctmtca tggttatctt ccaacccact gtcacgctga cactatgctg 3720
ccatgcctgg gctgtggacc tactgggcat ttgaggaayt ggagaatgga gatggcaaga 3780
gggcaggctt ttaagtttgg gttggagaca acttcctgtg gcccccacaa gctgagtctg 3840
gccttctcca gctggcccca aaaaaggcct ttgctacatc ctgattatct ctgaaagtaa 3900
tcaatcaagt ggctccagta gctctggatt ttctgccagg gctgggccat tgtggtgctg 3960
ccccagtatg acatgggacc aaggccagcg caggttatcc acctctgcct ggaagtctat 4020
actctaccca gggcatccct ctggtcagag gcagtgagta ctgggaactg gaggctgacc 4080
tgtgcttaga agtcctttaa tctgggctgg tacaggcctc agccttgccc tcaatgcacg 4140
aaaggtggcc caggagagag gatcaatgcc ataggaggca gaagtctggc ctctgtgcct 4200
ctatggagac tatcttccag ttgctgctca acagagttgt tggctgagac ctgcttggga 4260
gtctctgctg gcccttcatc tgttcaggaa cacacacaca cacacactca cacacgcaca 4320
cacaatcaca atttgctaca gcaacaaaaa agacattggg ctgtggcatt attaattaaa 4380
gatgatatcc agtcaaaaaa aaact 4405

<210> 7
<211> 3938
<212> DNA
<213> Homo sapiens

<400> 7
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc 300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg 360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt 420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa 480
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg 540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc 600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc 660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg 720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc 780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc 840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca 900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg 960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca 1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac 1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt gggggggccga 1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg 1200

```
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac   1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc   1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc   1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct   1440
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc   1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg   1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc   1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc   1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc   1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg   1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg   1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc   1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat   1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca   2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt   2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg   2160
tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga   2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag   2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc   2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc   2400
gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc   2460
tgccccgaag tgcggcgcaa cacccccctgg acgccgtggc tgcccgtgaa cgtgacgcag   2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgccccct tgcagacccg   2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc   2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg   2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc   2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc   2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt   2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt   2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt   3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg   3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt   3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc   3180
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt   3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg   3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc   3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag   3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat   3480
gacagagcca acttctaccc attgcagcag accaatgcca gcgcaggtta tccacctctg   3540
cctggaagtc tatactctac ccagggcatc cctctggtca gaggcagtga gtactgggaa   3600
ctggaggctg acctgtgctt agaagtcctt taatctgggc tggtacaggc ctcagccttg   3660
ccctcaatgc acgaaaggtg gcccaggaga gaggatcaat gccataggag gcagaagtct   3720
ggcctctgtg cctctatgga gactatcttc cagttgctgc tcaacagagt tgttggctga   3780
gacctgcttg ggagtctctg ctggcccttc atctgttcag gaacacacac acacacacac   3840
tcacacacgc acacacaatc acaatttgct acagcaacaa aaaagacatt gggctgtggc   3900
attattaatt aaagatgata tccagtcaaa aaaaaact                           3938
```

<210> 8
<211> 1095
<212> PRT
<213> Homo sapiens

<400> 8

```
Met Val Leu Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu
1               5                   10                  15

Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu
            20                  25                  30

Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val
```

```
        35                  40                  45

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

Ala Arg Asp Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu
65                  70                  75                  80

Ile Val Gly Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val
                85                  90                  95

Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg
            100                 105                 110

Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val
        115                 120                 125

Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn
    130                 135                 140

Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg
145                 150                 155                 160

Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg
                165                 170                 175

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

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

Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu
    210                 215                 220

Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr
225                 230                 235                 240

Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val
                245                 250                 255

Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe
            260                 265                 270

Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys
        275                 280                 285

Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala
    290                 295                 300

Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn
305                 310                 315                 320

Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala
                325                 330                 335

Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg
            340                 345                 350

Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly
        355                 360                 365
```

```
Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu
    370                 375                 380

Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val
385                 390                 395                 400

Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu
                405                 410                 415

Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr
            420                 425                 430

Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser
        435                 440                 445

Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro
    450                 455                 460

Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg
465                 470                 475                 480

Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu
                485                 490                 495

Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp
            500                 505                 510

Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu
        515                 520                 525

Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro
    530                 535                 540

Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp
545                 550                 555                 560

Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys
                565                 570                 575

Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp
            580                 585                 590

Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Gly
        595                 600                 605

Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro Val
    610                 615                 620

Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro
625                 630                 635                 640

Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe
                645                 650                 655

Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp
            660                 665                 670

Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser
        675                 680                 685
```

```
Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val
    690                 695                 700

Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn
705                 710                 715                 720

Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala
                725                 730                 735

Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp
            740                 745                 750

Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys
        755                 760                 765

Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val
    770                 775                 780

Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp
785                 790                 795                 800

Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly
                805                 810                 815

Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly
            820                 825                 830

Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro
        835                 840                 845

Gln Ala Cys Pro Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser
    850                 855                 860

Pro Cys Ser Ala Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser
865                 870                 875                 880

Cys Thr Ser Pro Ala Pro Ser Pro Gly Glu Asp Ile Cys Leu Gly Leu
                885                 890                 895

His Thr Glu Glu Ala Leu Cys Ala Thr Gln Ala Cys Pro Glu Gly Trp
            900                 905                 910

Ser Pro Trp Ser Glu Trp Ser Lys Cys Thr Asp Asp Gly Ala Gln Ser
        915                 920                 925

Arg Ser Arg His Cys Glu Glu Leu Leu Pro Gly Ser Ser Ala Cys Ala
    930                 935                 940

Gly Asn Ser Ser Gln Ser Arg Pro Cys Pro Tyr Ser Glu Ile Arg Val
945                 950                 955                 960

Ile Leu Pro Ala Ser Ser Met Glu Glu Ala Thr Asp Cys Ala Gly Phe
                965                 970                 975

Asn Leu Ile His Leu Val Ala Thr Gly Ile Ser Cys Phe Leu Gly Ser
            980                 985                 990

Gly Leu Leu Thr Leu Ala Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg
        995                 1000                1005

Gln Ser  Gln Glu Ser Thr Leu  Val His Pro Ala Thr  Pro Asn His
```

1010 1015 1020

Leu His Tyr Lys Gly Gly Gly Thr Pro Lys Asn Glu Lys Tyr Thr
1025 1030 1035

Pro Met Glu Phe Lys Thr Leu Asn Lys Asn Asn Leu Ile Pro Asp
1040 1045 1050

Asp Arg Ala Asn Phe Tyr Pro Leu Gln Gln Thr Asn Val Tyr Thr
1055 1060 1065

Thr Thr Tyr Tyr Pro Ser Pro Leu Asn Lys His Ser Phe Arg Pro
1070 1075 1080

Glu Ala Ser Pro Gly Gln Arg Cys Phe Pro Asn Ser
1085 1090 1095

<210> 9
<211> 1248
<212> PRT
<213> Homo sapiens

<400> 9
Arg Pro Pro His Ser Gln Thr Gly Arg Gln Pro Ile Trp Leu Ala Pro
1 5 10 15

Ala Ala Pro Arg Arg Pro Gly Val Gly Ser Arg Gly Glu Pro Gly Thr
20 25 30

Cys Thr Arg Leu Trp Glu Pro Ala Trp Val Arg Val Ala Leu Gly Pro
35 40 45

Ala Arg Ala Val Val Gly Ala Ser Gly Leu Gln Arg Arg Trp Gly Pro
50 55 60

Gly Thr Gln Ala Trp Arg Arg Arg Arg Thr Ser Glu Ala Glu Gly Arg
65 70 75 80

Arg Asp Arg Val Ser Gly Ser Ser Trp Cys Leu Ala Cys Val Ser Trp
85 90 95

Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His Leu Val Pro
100 105 110

Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly Trp Thr Val
115 120 125

Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro Cys Leu Pro
130 135 140

Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro
145 150 155 160

Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His
165 170 175

Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln
180 185 190

Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln

```
        195                 200                 205

Pro Trp Val Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln
    210                 215                 220

Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn
225                 230                 235                 240

Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr
                245                 250                 255

Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly
            260                 265                 270

Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala
        275                 280                 285

Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys
    290                 295                 300

Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu Lys Ile Asn
305                 310                 315                 320

Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val
                325                 330                 335

Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser
            340                 345                 350

Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu
        355                 360                 365

Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val
    370                 375                 380

Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn
385                 390                 395                 400

Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg
                405                 410                 415

Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp
            420                 425                 430

Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val
        435                 440                 445

Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln
    450                 455                 460

Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala
465                 470                 475                 480

Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn
                485                 490                 495

Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile
            500                 505                 510

Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly
        515                 520                 525
```

```
Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu
    530                 535                 540

Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val
545                 550                 555                 560

Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln
                565                 570                 575

Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly
            580                 585                 590

Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys
        595                 600                 605

Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu
    610                 615                 620

Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu
625                 630                 635                 640

Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg
                645                 650                 655

Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp
            660                 665                 670

Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser
        675                 680                 685

Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg
    690                 695                 700

Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu
705                 710                 715                 720

Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp
                725                 730                 735

Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile
            740                 745                 750

His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val Ile
        755                 760                 765

Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe Gln Val Arg
    770                 775                 780

Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly Arg Ile Cys
785                 790                 795                 800

Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys
                805                 810                 815

Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser
            820                 825                 830

Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn
        835                 840                 845
```

```
Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro
    850                 855                 860

Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu
865                 870                 875                 880

Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg
                885                 890                 895

Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly
            900                 905                 910

Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser
        915                 920                 925

Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr
    930                 935                 940

Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser
945                 950                 955                 960

Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr
                965                 970                 975

Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp
            980                 985                 990

Ala Ala Glu Tyr Gln Asp Cys Asn  Pro Gln Ala Cys Pro  Val Arg Gly
        995                 1000                1005

Ala Trp  Ser Cys Trp Thr Ser  Trp Ser Pro Cys Ser  Ala Ser Cys
    1010                1015                1020

Gly Gly  Gly His Tyr Gln Arg  Thr Arg Ser Cys Thr  Ser Pro Ala
    1025                1030                1035

Pro Ser  Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
    1040                1045                1050

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1055                1060                1065

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1070                1075                1080

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1085                1090                1095

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1100                1105                1110

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1115                1120                1125

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1130                1135                1140

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1145                1150                1155

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
```

1160 1165 1170

Pro Asn His Leu His Tyr Lys Gly Gly Gly Thr Pro Lys Asn Glu
1175 1180 1185

Lys Tyr Thr Pro Met Glu Phe Lys Thr Leu Asn Lys Asn Asn Leu
1190 1195 1200

Ile Pro Asp Asp Arg Ala Asn Phe Tyr Pro Leu Gln Gln Thr Asn
1205 1210 1215

Val Tyr Thr Thr Thr Tyr Tyr Pro Ser Pro Leu Asn Lys His Ser
1220 1225 1230

Phe Arg Pro Glu Ala Ser Pro Gly Gln Arg Cys Phe Pro Asn Ser
1235 1240 1245

<210> 10
<211> 1150
<212> PRT
<213> Homo sapiens

<400> 10
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1 5 10 15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
20 25 30

Leu His Pro Pro Leu Gly Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala
35 40 45

Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro Leu Ala Val
50 55 60

Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His Leu Ser Ser
65 70 75 80

Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala
85 90 95

Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln Pro Trp Val
100 105 110

Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln Leu Ala Leu
115 120 125

Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn Tyr Leu Phe
130 135 140

Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr Glu Trp Ala
145 150 155 160

Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu
165 170 175

Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala Gly Arg Lys
180 185 190

Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys Thr Ser Arg

195 200 205

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

Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val Ile Ser Ser
225 230 235 240

Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser Gly Arg Asp
245 250 255

Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala
260 265 270

Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr
275 280 285

Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu
290 295 300

His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg Val Cys Lys
305 310 315 320

Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe
325 330 335

Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr
340 345 350

Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln Asp Leu Ile
355 360 365

Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala Ser Ala Val
370 375 380

Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe
385 390 395 400

Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro
405 410 415

Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu
420 425 430

Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu Phe Leu Met
435 440 445

Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val Thr Gln Asp
450 455 460

Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln Ala Lys Asp
465 470 475 480

Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu
485 490 495

Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys Tyr Leu Glu
500 505 510

Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu
515 520 525

```
Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu Arg Asp Gly
    530                 535                 540

Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly
545                 550                 555                 560

Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln
                565                 570                 575

Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser Leu Trp Thr
            580                 585                 590

Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg Asp Gly Gly
        595                 600                 605

Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu Asp Gly Asp
    610                 615                 620

Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg
625                 630                 635                 640

Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile His Ile Ala
                645                 650                 655

Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val His Arg Gly Pro
            660                 665                 670

Leu Cys Ser His Val Leu Trp His Ala Ala Ser Arg Ser Ala Ser Glu
        675                 680                 685

Val Ala Ala Thr Leu Leu Pro Ala Thr Gly Ala Ala Ser Ala Trp Ala
    690                 695                 700

Arg Ala Trp Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys Pro Val
705                 710                 715                 720

Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn
                725                 730                 735

Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn
            740                 745                 750

Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro Glu Gly
        755                 760                 765

Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu Pro Val
    770                 775                 780

Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr
785                 790                 795                 800

Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly Arg Arg
                805                 810                 815

Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp
            820                 825                 830

Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr Ser Pro
        835                 840                 845
```

```
His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys
    850                 855                 860

Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr Cys Thr
865                 870                 875                 880

Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp Ala Ala
                885                 890                 895

Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro Val Arg Gly Ala Trp
            900                 905                 910

Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala Ser Cys Gly Gly Gly
        915                 920                 925

His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro Ala Pro Ser Pro Gly
    930                 935                 940

Glu Asp Ile Cys Leu Gly Leu His Thr Glu Glu Ala Leu Cys Ala Thr
945                 950                 955                 960

Gln Ala Cys Pro Glu Gly Trp Ser Pro Trp Ser Glu Trp Ser Lys Cys
                965                 970                 975

Thr Asp Asp Gly Ala Gln Ser Arg Ser Arg His Cys Glu Glu Leu Leu
            980                 985                 990

Pro Gly Ser Ser Ala Cys Ala Gly  Asn Ser Ser Gln Ser  Arg Pro Cys
        995                 1000                1005

Pro Tyr  Ser Glu Ile Arg Val  Ile Leu Pro Ala Ser  Ser Met Glu
    1010                1015                1020

Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1025                1030                1035

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1040                1045                1050

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1055                1060                1065

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1070                1075                1080

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
    1085                1090                1095

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1100                1105                1110

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1115                1120                1125

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1130                1135                1140

Gln Arg  Cys Phe Pro Asn Ser
    1145                1150
```

<210> 11
<211> 1211
<212> PRT
<213> Homo sapiens

<400> 11

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140

Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

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

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
```

290 295 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
305 310 315 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
325 330 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
340 345 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
355 360 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
370 375 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385 390 395 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
405 410 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
420 425 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
435 440 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
450 455 460

Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465 470 475 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
485 490 495

Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
500 505 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
515 520 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
530 535 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545 550 555 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
565 570 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
580 585 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
595 600 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
610 615 620

```
Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625                 630                 635                 640

Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
                645                 650                 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
            660                 665                 670

Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
        675                 680                 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
    690                 695                 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705                 710                 715                 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
                725                 730                 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
            740                 745                 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
        755                 760                 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
    770                 775                 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
785                 790                 795                 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
                805                 810                 815

Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
            820                 825                 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
        835                 840                 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
    850                 855                 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865                 870                 875                 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
                885                 890                 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
            900                 905                 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
        915                 920                 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
    930                 935                 940
```

```
Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945                 950                 955                 960

Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
                965                 970                 975

Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
            980                 985                 990

Ala Pro Ser Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
        995                 1000                1005

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1010                1015                1020

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1025                1030                1035

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1040                1045                1050

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1055                1060                1065

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1070                1075                1080

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1085                1090                1095

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1100                1105                1110

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                1120                1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1130                1135                1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                1150                1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                1165                1170

Ala Ser  Ala Gly Tyr Pro Pro  Leu Pro Gly Ser Leu  Tyr Ser Thr
    1175                1180                1185

Gln Gly  Ile Pro Leu Val Arg  Gly Ser Glu Tyr Trp  Glu Leu Glu
    1190                1195                1200

Ala Asp  Leu Cys Leu Glu Val  Leu
    1205                1210
```

<210> 12
<211> 1203
<212> PRT
<213> Homo sapiens

<400> 12

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140

Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

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

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
    290                 295                 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
305                 310                 315                 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
```

325 330 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
340 345 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
355 360 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
370 375 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385 390 395 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
405 410 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
420 425 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
435 440 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
450 455 460

Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465 470 475 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
485 490 495

Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
500 505 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
515 520 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
530 535 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545 550 555 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
565 570 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
580 585 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
595 600 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
610 615 620

Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625 630 635 640

Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
645 650 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
660 665 670

Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
675 680 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
690 695 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705 710 715 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
725 730 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
740 745 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
755 760 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
770 775 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
785 790 795 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
805 810 815

Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
820 825 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
835 840 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
850 855 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865 870 875 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
885 890 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
900 905 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
915 920 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
930 935 940

Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945 950 955 960

Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
965 970 975

```
Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
            980                 985                 990

Ala Pro Ser Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
        995                 1000                1005

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1010                1015                1020

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1025                1030                1035

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1040                1045                1050

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1055                1060                1065

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1070                1075                1080

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1085                1090                1095

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1100                1105                1110

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                1120                1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1130                1135                1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                1150                1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                1165                1170

Val Tyr  Thr Thr Thr Tyr Tyr  Pro Ser Pro Leu Asn  Lys His Ser
    1175                1180                1185

Phe Arg  Pro Glu Ala Ser Pro  Gly Gln Arg Cys Phe  Pro Asn Ser
    1190                1195                1200
```

<210> 13
<211> 1240
<212> PRT
<213> Homo sapiens

<400> 13

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                   10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45
```

```
Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Arg Lys Pro
                85                  90                  95

Ser Gly Asp Phe Glu Trp Arg Gln Gly Trp Arg Gly Pro Gly Glu Glu
            100                 105                 110

Asp Trp Pro Glu Ser Pro Ser Pro Lys Val Leu Met Asp Ser Ala Gly
        115                 120                 125

Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro
    130                 135                 140

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

Leu Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser
                165                 170                 175

Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr
            180                 185                 190

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

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

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

Val Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg
                245                 250                 255

Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr
            260                 265                 270

Val Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr
        275                 280                 285

Asn Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser
    290                 295                 300

Arg Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro
305                 310                 315                 320

Arg His Asn Ser Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala
                325                 330                 335

Ala Thr Val Ile Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser
            340                 345                 350

Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp
        355                 360                 365
```

```
Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala
    370                 375                 380

Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr
385                 390                 395                 400

Val Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg
                405                 410                 415

Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn
            420                 425                 430

Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser
        435                 440                 445

Ala Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr
    450                 455                 460

Asn Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser
465                 470                 475                 480

Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro
                485                 490                 495

Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys
            500                 505                 510

Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser
        515                 520                 525

Leu Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro
    530                 535                 540

Val Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His
545                 550                 555                 560

Leu Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu
                565                 570                 575

Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala
            580                 585                 590

Ser Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro
        595                 600                 605

Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala
    610                 615                 620

Arg Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu
625                 630                 635                 640

Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg
                645                 650                 655

Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu
            660                 665                 670

Glu Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys
        675                 680                 685

Pro Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro
```

690 695 700

Trp Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu
705 710 715 720

Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu
725 730 735

Asp Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly
740 745 750

Gly Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro
755 760 765

Val Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala
770 775 780

Pro Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg
785 790 795 800

Phe Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser
805 810 815

Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln
820 825 830

Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly
835 840 845

Val Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg
850 855 860

Asn Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly
865 870 875 880

Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala
885 890 895

Asp Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr
900 905 910

Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu
915 920 925

Val Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly
930 935 940

Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu
945 950 955 960

Gly Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn
965 970 975

Gly Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn
980 985 990

Pro Gln Ala Cys Pro Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp
995 1000 1005

Ser Pro Cys Ser Ala Ser Cys Gly Gly Gly His Tyr Gln Arg Thr
1010 1015 1020

```
Arg Ser  Cys Thr Ser Pro Ala  Pro Ser Pro Gly Glu  Asp Ile Cys
    1025                 1030                 1035

Leu Gly  Leu His Thr Glu Glu  Ala Leu Cys Ala Thr  Gln Ala Cys
    1040                 1045                 1050

Pro Glu  Gly Trp Ser Pro Trp  Ser Glu Trp Ser Lys  Cys Thr Asp
    1055                 1060                 1065

Asp Gly  Ala Gln Ser Arg Ser  Arg His Cys Glu Glu  Leu Leu Pro
    1070                 1075                 1080

Gly Ser  Ser Ala Cys Ala Gly  Asn Ser Ser Gln Ser  Arg Pro Cys
    1085                 1090                 1095

Pro Tyr  Ser Glu Ile Arg Val  Ile Leu Pro Ala Ser  Ser Met Glu
    1100                 1105                 1110

Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1115                 1120                 1125

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1130                 1135                 1140

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1145                 1150                 1155

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1160                 1165                 1170

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
    1175                 1180                 1185

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1190                 1195                 1200

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1205                 1210                 1215

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1220                 1225                 1230

Gln Arg  Cys Phe Pro Asn Ser
    1235                 1240
```

<210> 14
<211> 4567
<212> DNA
<213> Homo sapiens

<400> 14

```
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg     60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt    120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct    180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga    240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc    300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg    360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt    420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa    480
```

```
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg    540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc    600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc    660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg    720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc    780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc    840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca    900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg    960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca   1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac   1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt ggggggccga   1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg   1200
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac   1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc   1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc   1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct   1440
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc   1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg   1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc   1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc   1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc   1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg   1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg   1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc   1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat   1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca   2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt   2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg   2160
tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga   2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag   2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc   2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc   2400
gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc   2460
tgccccgaag tgcggcgcaa cacccctgg acgccgtggc tgcccgtgaa cgtgacgcag   2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgcccct tgcagacccg   2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc   2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg   2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc   2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc   2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt   2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt   2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt   3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg   3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt   3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc   3180
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt   3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg   3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc   3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag   3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat   3480
gacagagcca acttctaccc attgcagcag accaatgtgt acacgactac ttactaccca   3540
agccccctga acaaacacag cttccggccc gaggcctcac ctggacaacg gtgcttcccc   3600
aacagctgat accgccgtcc tggggacttg ggcttcttgc cttcataagg cacagagcag   3660
atggagatgg gacagtggag ccagtttggt tttctccctc tgcactaggc caagaacttg   3720
ctgccttgcc tgtgggggggt cccatccggc ttcagagagc tctggctggc attgaccatg   3780
ggggaaaggg ctggtttcag gctgacatat ggccgcaggt ccagttcagc ccaggtctmt   3840
catggttatc ttccaaccca ctgtcacgct gacactatgc tgccatgcct gggctgtgga   3900
cctactgggc atttgaggaa ytggagaatg gagatggcaa gagggcaggc ttttaagttt   3960
gggttggaga caacttcctg tggcccccac aagctgagtc tggccttctc cagctggccc   4020
caaaaaaggc ctttgctaca tcctgattat ctctgaaagt aatcaatcaa gtggctccag   4080
tagctctgga ttttctgcca gggctgggcc attgtggtgc tgccccagta tgacatggga   4140
```

ccaaggccag cgcaggttat ccacctctgc ctggaagtct atactctacc cagggcatcc 4200
ctctggtcag aggcagtgag tactgggaac tggaggctga cctgtgctta gaagtccttt 4260
aatctgggct ggtacaggcc tcagccttgc cctcaatgca cgaaaggtgg cccaggagag 4320
aggatcaatg ccataggagg cagaagtctg gcctctgtgc ctctatggag actatcttcc 4380
agttgctgct caacagagtt gttggctgag acctgcttgg gagtctctgc tggcccttca 4440
tctgttcagg aacacacaca cacacacact cacacacgca cacacaatca caatttgcta 4500
cagcaacaaa aaagacattg ggctgtggca ttattaatta aagatgatat ccagtcaaaa 4560
aaaaact 4567

<210> 15
<211> 453
<212> DNA
<213> Homo sapiens

<400> 15
agtaatcagc tcggtaccgg catgtgctgt agccagcgca ggttatccac ctctgcctgg 60
aagtctatac tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga 120
ggctgacctg tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc 180
aatgcacgaa aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct 240
ctgtgcctct atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct 300
gcttgggagt ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca 360
cacgcacaca caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat 420
taattaaaga tgatatccag tcaaaaaaaa act 453

<210> 16
<211> 4675
<212> DNA
<213> Homo sapiens

<400> 16
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggca ggaagccaag tggagacttt 300
gaatggaggc aaggatggag gggacctggg gaagaggact ggcctgaatc accttcccca 360
aaggtcctca tggactgagc tggaggtctt ctcccctgtc tgcctcccgg agctaggact 420
gcagaggggc ctatcatggt gcttgcaggc cccctggctg tctcgctgtt gctgcccagc 480
ctcacactgc tggtgtccca cctctccagc tcccaggatg tctccagtga gcccagcagt 540
gagcagcagc tgtgcgccct tagcaagcac cccaccgtgg cctttgaaga cctgcagccg 600
tgggtctcta acttcaccta ccctggagcc cgggatttct cccagctggc tttggacccc 660
tccgggaacc agctcatcgt gggagccagg aactacctct tcagactcag ccttgccaat 720
gtctctcttc ttcaggccac agagtgggcc tccagtgagg acacgcgccg ctcctgccaa 780
agcaaaggga agactgagga ggagtgtcag aactacgtgc gagtcctgat cgtcgccggc 840
cggaaggtgt tcatgtgtgg aaccaatgcc ttttccccca tgtgcaccag cagacaggtg 900
gggaacctca gccggactac tgagaagatc aatggtgtgg cccgctgccc ctatgaccca 960
cgccacaact ccacagctgt catctcctcc caggggggagc tctatgcagc cacggtcatc 1020
gacttctcag gtcgggaccc tgccatctac cgcagcctgg gcagtgggcc accgcttcgc 1080
actgcccaat ataactccaa gtggcttaat gagccaaact tcgtggcagc ctatgatatt 1140
gggctgtttg catacttctt cctgcgggag aacgcagtgg agcacgactg tggacgcacc 1200
gtgtactctc gcgtggcccg cgtgtgcaag aatgacgtgg ggggccgatt cctgctggag 1260
gacacatgga ccacattcat gaaggcccgg ctcaactgct cccgcccggg cgaggtcccc 1320
ttctactata acgagctgca gagtgccttc cacttgccag agcaggacct catctatgga 1380
gttttcacaa ccaacgtaaa cagcatcgcg gcttctgctg tctgcgcctt caacctcagt 1440
gctatctccc aggctttcaa tggcccattt cgctaccagg agaaccccag ggctgcctgg 1500
ctccccatag ccaaccccat ccccaatttc cagtgtggca ccctgcctga gaccggtccc 1560
aacgagaacc tgacggagcg cagcctgcag gacgcgcagc gcctcttcct gatgagcgag 1620
gccgtgcagc cggtgacacc cgagccctgt gtcacccagg acagcgtgcg cttctcacac 1680
ctcgtggtgg acctggtgca ggctaaagac acgctctacc atgtactcta cattggcacc 1740

gagtcgggca ccatcctgaa ggcgctgtcc acggcgagcc gcagcctcca cggctgctac 1800
ctggaggagc tgcacgtgct gccccccggg cgccgcgagc ccctgcgcag cctgcgcatc 1860
ctgcacagcg cccgcgcgct cttcgtgggg ctgagagacg gcgtcctgcg ggtcccactg 1920
gagaggtgcg ccgcctaccg cagccagggg gcatgcctgg gggcccggga cccgtactgt 1980
ggctgggacg ggaagcagca acgttgcagc acactcgagg acagctccaa catgagcctc 2040
tggacccaga acatcaccgc ctgtcctgtg cggaatgtga cacgggatgg gggcttcggc 2100
ccatggtcac catggcaacc atgtgagcac ttggatgggg acaactcagg ctcttgcctg 2160
tgtcgagctc gatcctgtga ttcccctcga ccccgctgtg ggggccttga ctgcctgggg 2220
ccagccatcc acatcgccaa ctgctccagg aatggggcgt ggaccccgtg gtcatcgtgg 2280
gcgctgtgca gcacgtcctg tggcatcggc ttccaggtcc gccagcgaag ttgcagcaac 2340
cctgctcccc gccacggggg ccgcatctgc gtgggcaaga gccgggagga acggttctgt 2400
aatgagaaca cgccttgccc ggtgcccatc ttctgggctt cctggggctc ctggagcaag 2460
tgcagcagca actgtggagg gggcatgcag tcgcggcgtc gggcctgcga gaacggcaac 2520
tcctgcctgg gctgcggcgt ggagttcaag acgtgcaacc ccgagggctg ccccgaagtg 2580
cggcgcaaca ccccctggac gccgtggctg cccgtgaacg tgacgcaggg cggggcacgg 2640
caggagcagc ggttccgctt cacctgccgc gcgccccttg cagacccgca cggcctgcag 2700
ttcggcagga gaaggaccga gacgaggacc tgtcccgcgg acggctccgg ctcctgcgac 2760
accgacgccc tggtggaggt cctcctgcgc agcgggagca cctccccgca cacggtgagc 2820
gggggctggg ccgcctgggg cccgtggtcg tcctgctccc gggactgcga gctgggcttc 2880
cgcgtccgca agagaacgtg cactaacccg gagccccgca acgggggcct gccctgcgtg 2940
ggcgatgctg ccgagtacca ggactgcaac ccccaggctt gcccagttcg gggtgcttgg 3000
tcctgctgga cctcatggtc tccatgctca gcttcctgtg gtgggggtca ctatcaacgc 3060
acccgttcct gcaccagccc cgcaccctcc ccaggtgagg acatctgtct cgggctgcac 3120
acggaggagg cactatgtgc cacacaggcc tgcccagaag gctggtcgcc ctggtctgag 3180
tggagtaagt gcactgacga cggagcccag agccgaagcc ggcactgtga ggagctcctc 3240
ccagggtcca gcgcmtgtgc tggaaacagc agccagagcc gcccctgccc ctacagcgag 3300
attcscgtca tcctgccagc ctccagcatg gaggaggcca ccgactgtgc agggttcaat 3360
ctcatccact tggtggccac gggcatctcc tgcttcttgg gctctgggct cctgacccta 3420
gcagtgtacc tgtcttgcca gcactgccag cgtcagtccc aggagtccac actggtccat 3480
cctgccaccc ccaaccattt gcactacaag ggcggaggca ccccgaagaa tgaaaagtac 3540
acacccatgg aattcaagac cctgaacaag aataacttga tccctgatga cagagccaac 3600
ttctacccat tgcagcagac caatgtgtac acgactactt actacccaag ccccctgaac 3660
aaacacagct tccggcccga ggcctcacct ggacaacggt gcttccccaa cagctgatac 3720
cgccgtcctg gggacttggg cttcttgcct tcataaggca cagagcagat ggagatggga 3780
cagtggagcc agtttggttt tctccctctg cactaggcca agaacttgct gccttgcctg 3840
tgggggtcc catccggctt cagagagctc tggctggcat tgaccatggg ggaaagggct 3900
ggtttcaggc tgacatatgg ccgcaggtcc agttcagccc aggtctmtca tggttatctt 3960
ccaacccact gtcacgctga cactatgctg ccatgcctgg gctgtggacc tactgggcat 4020
ttgaggaayt ggagaatgga gatggcaaga gggcaggctt ttaagtttgg gttggagaca 4080
acttcctgtg gcccccacaa gctgagtctg gccttctcca gctggcccca aaaaaggcct 4140
ttgctacatc ctgattatct ctgaaagtaa tcaatcaagt ggctccagta gctctggatt 4200
ttctgccagg gctgggccat tgtggtgctg ccccagtatg acatgggacc aaggccagcg 4260
caggttatcc acctctgcct ggaagtctat actctaccca gggcatccct ctggtcagag 4320
gcagtgagta ctgggaactg gaggctgacc tgtgcttaga agtcctttaa tctgggctgg 4380
tacaggcctc agccttgccc tcaatgcacg aaaggtggcc caggagagag gatcaatgcc 4440
ataggaggca gaagtctggc ctctgtgcct ctatggagac tatcttccag ttgctgctca 4500
acagagttgt tggctgagac ctgcttggga gtctctgctg gcccttcatc tgttcaggaa 4560
cacacacaca cacacactca cacacgcaca cacaatcaca atttgctaca gcaacaaaaa 4620
agacattggg ctgtggcatt attaattaaa gatgatatcc agtcaaaaaa aaact 4675

<210> 17
<211> 4731
<212> DNA
<213> Homo sapiens

<400> 17
attggagatg ctcgggggca ggctgccgcg ttgtgtcctg cttttctgcg gccagaccaa 60
gccgtctgga gctgctggtc aggttttctt gctgacctca cctgaccaca gtggcctggg 120
tggactctac agggaaatgt tgttttctcc ctgggagcag tagcagcagt cctggctccc 180
ctggactgag aactcctcat cagccccagg aagcccggac ccccttttcag ggatctggaa 240

```
ccggtgtgcc tgtggcccca ggtctgctcc caggcgtggg ctgaagtcct gacttctgtc    300
gctgggggca aggagtggga gagcccagct gctgcctggg ctttggcaga cagcaggctg    360
atggtgctgg cttccccgag actgcttctc ctgcctgctg tctgatttcc ctgcatggtg    420
cccgcagctg agctgctacg ggtcttctcc cctgtctgcc tcccggagct aggactgcag    480
aggggcctat catggtgctt gcaggccccc tggctgtctc gctgttgctg cccagcctca    540
cactgctggt gtcccacctc tccagctccc aggatgtctc cagtgagccc agcagtgagc    600
agcagctgtg cgcccttagc aagcacccca ccgtggcctt tgaagacctg cagccgtggg    660
tctctaactt cacctaccct ggagcccggg atttctccca gctggctttg gacccctccg    720
ggaaccagct catcgtggga gccaggaact acctcttcag actcagcctt gccaatgtct    780
ctcttcttca ggccacagag tgggcctcca gtgaggacac gcgccgctcc tgccaaagca    840
aagggaagac tgaggaggag tgtcagaact acgtgcgagt cctgatcgtc gccggccgga    900
aggtgttcat gtgtggaacc aatgcctttt cccccatgtg caccagcaga caggtgggga    960
acctcagccg gactactgag aagatcaatg gtgtggcccg ctgcccctat gacccacgcc   1020
acaactccac agctgtcatc tcctcccagg gggagctcta tgcagccacg gtcatcgact   1080
tctcaggtcg ggaccctgcc atctaccgca gcctgggcag tgggccaccg cttcgcactg   1140
cccaatataa ctccaagtgg cttaatgagc caaacttcgt ggcagcctat gatattgggc   1200
tgtttgcata cttcttcctg cgggagaacg cagtggagca cgactgtgga cgcaccgtgt   1260
actctcgcgt ggcccgcgtg tgcaagaatg acgtgggggg ccgattcctg ctggaggaca   1320
catggaccac attcatgaag gcccggctca actgctcccg cccgggcgag gtccccttct   1380
actataacga gctgcagagt gccttccact tgccagagca ggacctcatc tatggagttt   1440
tcacaaccaa cgtaaacagc atcgcggctt ctgctgtctg cgccttcaac ctcagtgcta   1500
tctcccaggc tttcaatggc ccatttcgct accaggagaa ccccagggct gcctggctcc   1560
ccatagccaa ccccatcccc aatttccagt gtggcaccct gcctgagacc ggtcccaacg   1620
agaacctgac ggagcgcagc ctgcaggacg cgcagcgcct cttcctgatg agcgaggccg   1680
tgcagccggt gacacccgag ccctgtgtca cccaggacag cgtgcgcttc tcacacctcg   1740
tggtggacct ggtgcaggct aaagacacgc tctaccatgt actctacatt ggcaccgagt   1800
cgggcaccat cctgaaggcg ctgtccacgg cgagccgcag cctccacggc tgctacctgg   1860
aggagctgca cgtgctgccc cccgggcgcc gcgagcccct gcgcagcctg cgcatcctgc   1920
acagcgcccg cgcgctcttc gtggggctga gagacggcgt cctgcgggtc ccactggaga   1980
ggtgcgccgc ctaccgcagc caggggggcat gcctgggggc ccgggacccg tactgtggct   2040
gggacgggaa gcagcaacgt tgcagcacac tcgaggacag ctccaacatg agcctctgga   2100
cccagaacat caccgcctgt cctgtgcgga atgtgacacg ggatgggggc ttcggcccat   2160
ggtcaccatg gcaaccatgt gagcacttgg atggggacaa ctcaggctct tgcctgtgtc   2220
gagctcgatc ctgtgattcc cctcgacccc gctgtggggg ccttgactgc ctggggccag   2280
ccatccacat cgccaactgc tccaggaatg gggcgtggac cccgtggtca tcgtgggcgc   2340
tgtgcagcac gtcctgtggc atcggcttcc aggtccgcca gcgaagttgc agcaaccctg   2400
ctccccgcca cgggggccgc atctgcgtgg gcaagagccg ggaggaacgg ttctgtaatg   2460
agaacacgcc ttgcccggtg cccatcttct gggcttcctg gggctcctgg agcaagtgca   2520
gcagcaactg tggaggggggc atgcagtcgc ggcgtcgggc ctgcgagaac ggcaactcct   2580
gcctgggctg cggcgtggag ttcaagacgt gcaaccccga gggctgcccc gaagtgcggc   2640
gcaacacccc ctggacgccg tggctgcccg tgaacgtgac gcagggcggg gcacggcagg   2700
agcagcggtt ccgcttcacc tgccgcgcgc cccttgcaga cccgcacggc ctgcagttcg   2760
gcaggagaag gaccgagacg aggacctgtc ccgcggacgg ctccggctcc tgcgacaccg   2820
acgccctggt ggaggtcctc ctgcgcagcg ggagcacctc cccgcacacg gtgagcgggg   2880
gctgggccgc ctggggcccg tggtcgtcct gctcccggga ctgcgagctg ggcttccgcg   2940
tccgcaagag aacgtgcact aacccggagc cccgcaacgg gggcctgccc tgcgtgggcg   3000
atgctgccga gtaccaggac tgcaaccccc aggcttgccc agttcggggt gcttggtcct   3060
gctggacctc atggtctcca tgctcagctt cctgtggtgg gggtcactat caacgcaccc   3120
gttcctgcac cagccccgca ccctccccag gtgaggacat ctgtctcggg ctgcacacgg   3180
aggaggcact atgtgccaca caggcctgcc cagaaggctg gtcgccctgg tctgagtgga   3240
gtaagtgcac tgacgacgga gcccagagcc gaagccggca ctgtgaggag ctcctcccag   3300
ggtccagcgc mtgtgctgga aacagcagcc agagccgccc ctgcccctac agcgagattc   3360
scgtcatcct gccagcctcc agcatggagg aggccaccga ctgtgcaggg ttcaatctca   3420
tccacttggt ggccacgggc atctcctgct tcttgggctc tgggctcctg accctagcag   3480
tgtacctgtc ttgccagcac tgccagcgtc agtcccagga gtccacactg gtccatcctg   3540
ccacccccaa ccatttgcac tacaagggcg gaggcacccc gaagaatgaa aagtacacac   3600
ccatggaatt caagaccctg aacaagaata acttgatccc tgatgacaga gccaacttct   3660
acccattgca gcagaccaat gtgtacacga ctacttacta cccaagcccc ctgaacaaac   3720
acagcttccg gcccgaggcc tcacctggac aacggtgctt ccccaacagc tgataccgcc   3780
gtcctgggga cttgggcttc ttgccttcat aaggcacaga gcagatggag atgggacagt   3840
ggagccagtt tggttttctc cctctgcact aggccaagaa cttgctgcct tgcctgtggg   3900
```

gggtcccatc cggcttcaga gagctctggc tggcattgac catggggaa agggctggtt 3960
tcaggctgac atatggccgc aggtccagtt cagcccaggt ctmtcatggt tatcttccaa 4020
cccactgtca cgctgacact atgctgccat gcctgggctg tggacctact gggcatttga 4080
ggaaytggag aatggagatg gcaagagggc aggcttttaa gtttgggttg gagacaactt 4140
cctgtggccc ccacaagctg agtctggcct tctccagctg gccccaaaaa aggcctttgc 4200
tacatcctga ttatctctga aagtaatcaa tcaagtggct ccagtagctc tggattttct 4260
gccagggctg ggccattgtg gtgctgcccc agtatgacat gggaccaagg ccagcgcagg 4320
ttatccacct ctgcctggaa gtctatactc tacccagggc atccctctgg tcagaggcag 4380
tgagtactgg gaactggagg ctgacctgtg cttagaagtc ctttaatctg ggctggtaca 4440
ggcctcagcc ttgccctcaa tgcacgaaag gtggcccagg agagaggatc aatgccatag 4500
gaggcagaag tctggcctct gtgcctctat ggagactatc ttccagttgc tgctcaacag 4560
agttgttggc tgagacctgc ttgggagtct ctgctggccc ttcatctgtt caggaacaca 4620
cacacacaca cactcacaca cgcacacaca atcacaattt gctacagcaa caaaaaagac 4680
attgggctgt ggcattatta attaaagatg atatccagtc aaaaaaaaac t 4731

<210> 18
<211> 4703
<212> DNA
<213> Homo sapiens

<400> 18
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagccagc 120
ttgggtccgg gttgcactgg gccctgccag ggctgtggtc ggcgcatctg ggctgcagcg 180
gcgatgggga cccgggaccc aggcctggag aaggagacgg acgagtgagg ctgagggacg 240
gagggacaga gtgagtggtt ccagctggtg cctggcctgt gtctcttgga tgccctgtgg 300
cttcagtccg tctcctgttg cccaccacct cgtccctggg ccgcctgata ccccagccca 360
acagctaagg tgtggatgga cagtaggggg ctggcttctc tcactggtca ggggtcttct 420
cccctgtctg cctcccggag ctaggactgc agaggggcct atcatggtgc ttgcaggccc 480
cctggctgtc tcgctgttgc tgcccagcct cacactgctg gtgtcccacc tctccagctc 540
ccaggatgtc tccagtgagc ccagcagtga gcagcagctg tgcgccctta gcaagcaccc 600
caccgtggcc tttgaagacc tgcagccgtg ggtctctaac ttcacctacc ctggagcccg 660
ggatttctcc cagctggctt tggacccctc cgggaaccag ctcatcgtgg gagccaggaa 720
ctacctcttc agactcagcc ttgccaatgt ctctcttctt caggccacag agtgggcctc 780
cagtgaggac acgcgccgct cctgccaaag caaagggaag actgaggagg agtgtcagaa 840
ctacgtgcga gtcctgatcg tcgccggccg gaaggtgttc atgtgtggaa ccaatgcctt 900
ttcccccatg tgcaccagca gacaggtggg gaacctcagc cggactactg agaagatcaa 960
tggtgtggcc cgctgccccc atgacccacg ccacaactcc acagctgtca tctcctccca 1020
gggggagctc tatgcagcca cggtcatcga cttctcaggt cgggaccctg ccatctaccg 1080
cagcctgggc agtgggccac cgcttcgcac tgcccaatat aactccaagt ggcttaatga 1140
gccaaacttc gtggcagcct atgatattgg gctgtttgca tacttcttcc tgcgggagaa 1200
cgcagtggag cacgactgtg gacgcaccgt gtactctcgc gtggcccgcg tgtgcaagaa 1260
tgacgtgggg ggccgattcc tgctggagga cacatggacc acattcatga aggcccggct 1320
caactgctcc cgcccgggcg aggtcccctt ctactataac gagctgcaga gtgccttcca 1380
cttgccagag caggacctca tctatggagt tttcacaacc aacgtaaaca gcatcgcggc 1440
ttctgctgtc tgcgccttca acctcagtgc tatctcccag gctttcaatg gcccatttcg 1500
ctaccaggag aaccccaggg ctgcctggct ccccatagcc aaccccatcc ccaatttcca 1560
gtgtggcacc ctgcctgaga ccggtcccaa cgagaacctg acggagcgca gcctgcagga 1620
cgcgcagcgc ctcttcctga tgagcgaggc cgtgcagccg gtgacacccg agccctgtgt 1680
cacccaggac agcgtgcgct tctcacacct cgtggtggac ctggtgcagg ctaaagacac 1740
gctctaccat gtactctaca ttggcaccga gtcgggcacc atcctgaagg cgctgtccac 1800
ggcgagccgc agcctccacg gctgctacct ggaggagctg cacgtgctgc cccccgggcg 1860
ccgcgagccc ctgcgcagcc tgcgcatcct gcacagcgcc cgcgcgctct tcgtggggct 1920
gagagacggc gtcctgcggg tcccactgga gaggtgcgcc gcctaccgca gccaggggc 1980
atgcctgggg gcccgggacc cgtactgtgg ctgggacggg aagcagcaac gttgcagcac 2040
actcgaggac agctccaaca tgagcctctg gacccagaac atcaccgcct gtcctgtgcg 2100
gaatgtgaca cgggatgggg gcttcggccc atggtcacca tggcaaccat gtgagcactt 2160
ggatggggac aactcaggct cttgcctgtg tcgagctcga tcctgtgatt cccctcgacc 2220
ccgctgtggg ggccttgact gcctggggcc agccatccac atcgccaact gctccaggaa 2280
tggggcgtgg accccgtggt catcgtgggc gctgtgcagc acgtcctgtg gcatcggctt 2340

ccaggtccgc cagcgaagtt gcagcaaccc tgctccccgc cacgggggcc gcatctgcgt 2400
gggcaagagc cgggaggaac ggttctgtaa tgagaacacg ccttgcccgg tgcccatctt 2460
ctgggcttcc tggggctcct ggagcaagtg cagcagcaac tgtggagggg gcatgcagtc 2520
gcggcgtcgg gcctgcgaga acggcaactc ctgcctgggc tgcggcgtgg agttcaagac 2580
gtgcaacccc gagggctgcc ccgaagtgcg gcgcaacacc ccctggacgc cgtggctgcc 2640
cgtgaacgtg acgcagggcg gggcacggca ggagcagcgg ttccgcttca cctgccgcgc 2700
gcccccttgca gacccgcacg gcctgcagtt cggcaggaga aggaccgaga cgaggacctg 2760
tcccgcggac ggctccggct cctgcgacac cgacgccctg gtggaggtcc tcctgcgcag 2820
cgggagcacc tccccgcaca cggtgagcgg gggctgggcc gcctggggcc cgtggtcgtc 2880
ctgctcccgg gactgcgagc tgggcttccg cgtccgcaag agaacgtgca ctaacccgga 2940
gccccgcaac gggggcctgc cctgcgtggg cgatgctgcc gagtaccagg actgcaaccc 3000
ccaggcttgc ccagttcggg gtgcttggtc ctgctggacc tcatggtctc catgctcagc 3060
ttcctgtggt gggggtcact atcaacgcac ccgttcctgc accagccccg caccctcccc 3120
aggtgaggac atctgtctcg ggctgcacac ggaggaggca ctatgtgcca cacaggcctg 3180
cccagaaggc tggtcgccct ggtctgagtg gagtaagtgc actgacgacg gagcccagag 3240
ccgaagccgg cactgtgagg agctcctccc agggtccagc gcmtgtgctg gaaacagcag 3300
ccagagccgc ccctgcccct acagcgagat tcscgtcatc ctgccagcct ccagcatgga 3360
ggaggccacc gactgtgcag ggttcaatct catccacttg gtggccacgg gcatctcctg 3420
cttcttgggc tctgggctcc tgaccctagc agtgtacctg tcttgccagc actgccagcg 3480
tcagtcccag gagtccacac tggtccatcc tgccaccccc aaccatttgc actacaaggg 3540
cggaggcacc ccgaagaatg aaaagtacac acccatggaa ttcaagaccc tgaacaagaa 3600
taacttgatc cctgatgaca gagccaactt ctacccattg cagcagacca atgtgtacac 3660
gactacttac tacccaagcc ccctgaacaa acacagcttc cggcccgagg cctcacctgg 3720
acaacggtgc ttccccaaca gctgataccg ccgtcctggg gacttgggct tcttgccttc 3780
ataaggcaca gagcagatgg agatgggaca gtggagccag tttggttttc tccctctgca 3840
ctaggccaag aacttgctgc cttgcctgtg gggggtccca tccggcttca gagagctctg 3900
gctggcattg accatggggg aaagggctgg tttcaggctg acatatggcc gcaggtccag 3960
ttcagcccag gtctmtcatg gttatcttcc aacccactgt cacgctgaca ctatgctgcc 4020
atgcctgggc tgtggaccta ctgggcattt gaggaaytgg agaatggaga tggcaagagg 4080
gcaggctttt aagtttgggt tggagacaac ttcctgtggc ccccacaagc tgagtctggc 4140
cttctccagc tggccccaaa aaaggccttt gctacatcct gattatctct gaaagtaatc 4200
aatcaagtgg ctccagtagc tctggatttt ctgccagggc tgggccattg tggtgctgcc 4260
ccagtatgac atgggaccaa ggccagcgca ggttatccac ctctgcctgg aagtctatac 4320
tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga ggctgacctg 4380
tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc aatgcacgaa 4440
aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct ctgtgcctct 4500
atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct gcttgggagt 4560
ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca cacgcacaca 4620
caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat taattaaaga 4680
tgatatccag tcaaaaaaaa act 4703

<210> 19
<211> 4405
<212> DNA
<213> Homo sapiens

<400> 19
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggaggtctt 120
ctcccctgtc tgcctcccgg agctaggact gcagaggggc ctatcatggt gcttgcaggc 180
cccctggctg tctcgctgtt gctgcccagc ctcacactgc tggtgtccca cctctccagc 240
tcccaggatg tctccagtga gcccagcagt gagcagcagc tgtgcgccct tagcaagcac 300
cccaccgtgg cctttgaaga cctgcagccg tgggtctcta acttcaccta ccctggagcc 360
cgggatttct cccagctggc tttggacccc tccgggaacc agctcatcgt gggagccagg 420
aactacctct tcagactcag ccttgccaat gtctctcttc ttcaggccac agagtgggcc 480
tccagtgagg acacgcgccg ctcctgccaa agcaaaggga agactgagga ggagtgtcag 540
aactacgtgc gagtcctgat cgtcgccggc cggaaggtgt tcatgtgtgg aaccaatgcc 600
ttttccccca tgtgcaccag cagacaggtg gggaacctca gccggactac tgagaagatc 660
aatggtgtgg cccgctgccc ctatgaccca cgccacaact ccacagctgt catctcctcc 720
caggggggagc tctatgcagc cacggtcatc gacttctcag gtcgggaccc tgccatctac 780

```
cgcagcctgg gcagtgggcc accgcttcgc actgcccaat ataactccaa gtggcttaat    840
gagccaaact tcgtggcagc ctatgatatt gggctgtttg catacttctt cctgcgggag    900
aacgcagtgg agcacgactg tggacgcacc gtgtactctc gcgtggcccg cgtgtgcaag    960
aatgacgtgg ggggccgatt cctgctggag gacacatgga ccacattcat gaaggcccgg   1020
ctcaactgct cccgcccggg cgaggtcccc ttctactata acgagctgca gagtgccttc   1080
cacttgccag agcaggacct catctatgga gttttcacaa ccaacgtaaa cagcatcgcg   1140
gcttctgctg tctgcgcctt caacctcagt gctatctccc aggctttcaa tggcccattt   1200
cgctaccagg agaaccccag ggctgcctgg ctccccatag ccaaccccat ccccaatttc   1260
cagtgtggca ccctgcctga gaccggtccc aacgagaacc tgacggagcg cagcctgcag   1320
gacgcgcagc gcctcttcct gatgagcgag gccgtgcagc cggtgacacc cgagccctgt   1380
gtcacccagg acagcgtgcg cttctcacac ctcgtggtgg acctggtgca ggctaaagac   1440
acgctctacc atgtactcta cattggcacc gagtcgggca ccatcctgaa ggcgctgtcc   1500
acggcgagcc gcagcctcca cggctgctac ctggaggagc tgcacgtgct gccccccggg   1560
cgccgcgagc ccctgcgcag cctgcgcatc ctgcacagcg cccgcgcgct cttcgtgggg   1620
ctgagagacg gcgtcctgcg ggtcccactg gagaggtgcg ccgcctaccg cagccagggg   1680
gcatgcctgg gggcccggga cccgtactgt ggctgggacg ggaagcagca acgttgcagc   1740
acactcgagg acagctccaa catgagcctc tggacccaga acatcaccgc ctgtcctgtg   1800
cggaatgtga cacgggatgg gggcttcggc ccatggtcac catggcaacc atgtgagcac   1860
ttggatgggg acaactcagg ctcttgcctg tgtcgagctc gatcctgtga ttcccctcga   1920
ccccgctgtg ggggccttga ctgcctgggg ccagccatcc acatcgccaa ctgctccagg   1980
aatggggcgt ggaccccgtg gtcatcgtgg gcgctgtgca gcacgtcctg tggcatcggc   2040
ttccaggtcc gccagcgaag ttgcagcaac cctgctcccc gccacggggg ccgcatctgc   2100
gtgggcaaga gccgggagga acggttctgt aatgagaaca cgccttgccc ggtgcccatc   2160
ttctgggctt cctggggctc ctggagcaag tgcagcagca actgtggagg gggcatgcag   2220
tcgcggcgtc gggcctgcga gaacggcaac tcctgcctgg gctgcggcgt ggagttcaag   2280
acgtgcaacc ccgagggctg ccccgaagtg cggcgcaaca ccccctggac gccgtggctg   2340
cccgtgaacg tgacgcaggg cggggcacgg caggagcagc ggttccgctt cacctgccgc   2400
gcgccccttg cagacccgca cggcctgcag ttcggcagga gaaggaccga gacgaggacc   2460
tgtcccgcgg acggctccgg ctcctgcgac accgacgccc tggtggaggt cctcctgcgc   2520
agcgggagca cctccccgca cacggtgagc gggggctggg ccgcctgggg cccgtggtcg   2580
tcctgctccc gggactgcga gctgggcttc cgcgtccgca agagaacgtg cactaacccg   2640
gagccccgca acgggggcct gccctgcgtg ggcgatgctg ccgagtacca ggactgcaac   2700
ccccaggctt gcccagttcg gggtgcttgg tcctgctgga cctcatggtc tccatgctca   2760
gcttcctgtg gtgggggtca ctatcaacgc acccgttcct gcaccagccc cgcaccctcc   2820
ccaggtgagg acatctgtct cgggctgcac acggaggagg cactatgtgc cacacaggcc   2880
tgcccagaag gctggtcgcc ctggtctgag tggagtaagt gcactgacga cggagcccag   2940
agccgaagcc ggcactgtga ggagctcctc ccagggtcca gcgcmtgtgc tggaaacagc   3000
agccagagcc gcccctgccc ctacagcgag attcscgtca tcctgccagc ctccagcatg   3060
gaggaggcca ccgactgtgc agggttcaat ctcatccact tggtggccac gggcatctcc   3120
tgcttcttgg gctctgggct cctgacccta gcagtgtacc tgtcttgcca gcactgccag   3180
cgtcagtccc aggagtccac actggtccat cctgccaccc ccaaccattt gcactacaag   3240
ggcggaggca ccccgaagaa tgaaaagtac acacccatgg aattcaagac cctgaacaag   3300
aataacttga tccctgatga cagagccaac ttctacccat tgcagcagac caatgtgtac   3360
acgactactt actacccaag ccccctgaac aaacacagct tccggcccga ggcctcacct   3420
ggacaacggt gcttccccaa cagctgatac cgccgtcctg gggacttggg cttcttgcct   3480
tcataaggca cagagcagat ggagatggga cagtggagcc agtttggttt tctccctctg   3540
cactaggcca agaacttgct gccttgcctg tgggggggtcc catccggctt cagagagctc   3600
tggctggcat tgaccatggg ggaaagggct ggtttcaggc tgacatatgg ccgcaggtcc   3660
agttcagccc aggtctmtca tggttatctt ccaacccact gtcacgctga cactatgctg   3720
ccatgcctgg gctgtggacc tactgggcat ttgaggaayt ggagaatgga gatggcaaga   3780
gggcaggctt ttaagtttgg gttggagaca acttcctgtg gcccccacaa gctgagtctg   3840
gccttctcca gctggcccca aaaaaggcct ttgctacatc ctgattatct ctgaaagtaa   3900
tcaatcaagt ggctccagta gctctggatt ttctgccagg gctgggccat tgtggtgctg   3960
ccccagtatg acatgggacc aaggccagcg caggttatcc acctctgcct ggaagtctat   4020
actctaccca gggcatccct ctggtcagag gcagtgagta ctgggaactg gaggctgacc   4080
tgtgcttaga agtcctttaa tctgggctgg tacaggcctc agccttgccc tcaatgcacg   4140
aaaggtggcc caggagagag gatcaatgcc ataggaggca gaagtctggc ctctgtgcct   4200
ctatggagac tatcttccag ttgctgctca acagagttgt tggctgagac ctgcttggga   4260
gtctctgctg gcccttcatc tgttcaggaa cacacacaca cacacactca cacacgcaca   4320
cacaatcaca atttgctaca gcaacaaaaa agacattggg ctgtggcatt attaattaaa   4380
gatgatatcc agtcaaaaaa aaact                                         4405
```

<210> 20
<211> 3938
<212> DNA
<213> Homo sapiens

<400> 20
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc 300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg 360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt 420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa 480
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg 540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc 600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc 660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg 720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc 780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc 840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca 900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg 960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca 1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac 1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt gggggggccga 1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg 1200
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac 1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc 1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc 1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct 1440
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc 1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg 1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc 1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc 1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc 1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg 1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg 1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc 1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat 1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca 2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt 2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg 2160
tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga 2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag 2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc 2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc 2400
gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc 2460
tgccccgaag tgcggcgcaa cacccccctgg acgccgtggc tgcccgtgaa cgtgacgcag 2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgccccct tgcagacccg 2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc 2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg 2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc 2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc 2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt 2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt 2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt 3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg 3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt 3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc 3180

```
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt   3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg   3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc   3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag   3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat   3480
gacagagcca acttctaccc attgcagcag accaatgcca gcgcaggtta tccacctctg   3540
cctggaagtc tatactctac ccagggcatc cctctggtca gaggcagtga gtactgggaa   3600
ctggaggctg acctgtgctt agaagtcctt taatctgggc tggtacaggc ctcagccttg   3660
ccctcaatgc acgaaaggtg gcccaggaga gaggatcaat gccataggag gcagaagtct   3720
ggcctctgtg cctctatgga gactatcttc cagttgctgc tcaacagagt tgttggctga   3780
gacctgcttg ggagtctctg ctggcccttc atctgttcag gaacacacac acacacacac   3840
tcacacacgc acacacaatc acaatttgct acagcaacaa aaaagacatt gggctgtggc   3900
attattaatt aaagatgata tccagtcaaa aaaaaact                           3938
```

<210> 21
<211> 1095
<212> PRT
<213> Homo sapiens

<400> 21

```
Met Val Leu Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu
1               5                  10                  15

Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu
            20                  25                  30

Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val
        35                  40                  45

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

Ala Arg Asp Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu
65                  70                  75                  80

Ile Val Gly Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val
                85                  90                  95

Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg
            100                 105                 110

Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val
        115                 120                 125

Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn
    130                 135                 140

Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg
145                 150                 155                 160

Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg
                165                 170                 175

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

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

Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu
```

```
    210                 215                 220

Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr
225                 230                 235                 240

Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val
                245                 250                 255

Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe
            260                 265                 270

Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys
        275                 280                 285

Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala
    290                 295                 300

Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn
305                 310                 315                 320

Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala
                325                 330                 335

Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg
            340                 345                 350

Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly
        355                 360                 365

Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu
    370                 375                 380

Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val
385                 390                 395                 400

Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu
                405                 410                 415

Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr
            420                 425                 430

Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser
        435                 440                 445

Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro
    450                 455                 460

Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg
465                 470                 475                 480

Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu
                485                 490                 495

Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp
            500                 505                 510

Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu
        515                 520                 525

Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro
    530                 535                 540
```

```
Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp
545                 550                 555                 560

Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys
                565                 570                 575

Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp
            580                 585                 590

Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Gly
        595                 600                 605

Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro Val
    610                 615                 620

Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro
625                 630                 635                 640

Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe
                645                 650                 655

Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp
            660                 665                 670

Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser
        675                 680                 685

Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val
    690                 695                 700

Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn
705                 710                 715                 720

Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala
                725                 730                 735

Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp
            740                 745                 750

Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys
        755                 760                 765

Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val
    770                 775                 780

Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp
785                 790                 795                 800

Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly
                805                 810                 815

Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly
            820                 825                 830

Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro
        835                 840                 845

Gln Ala Cys Pro Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser
    850                 855                 860
```

```
Pro Cys Ser Ala Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser
865                 870                 875                 880

Cys Thr Ser Pro Ala Pro Ser Pro Gly Glu Asp Ile Cys Leu Gly Leu
                885                 890                 895

His Thr Glu Glu Ala Leu Cys Ala Thr Gln Ala Cys Pro Glu Gly Trp
            900                 905                 910

Ser Pro Trp Ser Glu Trp Ser Lys Cys Thr Asp Asp Gly Ala Gln Ser
        915                 920                 925

Arg Ser Arg His Cys Glu Glu Leu Leu Pro Gly Ser Ser Ala Cys Ala
    930                 935                 940

Gly Asn Ser Ser Gln Ser Arg Pro Cys Pro Tyr Ser Glu Ile Arg Val
945                 950                 955                 960

Ile Leu Pro Ala Ser Ser Met Glu Glu Ala Thr Asp Cys Ala Gly Phe
                965                 970                 975

Asn Leu Ile His Leu Val Ala Thr Gly Ile Ser Cys Phe Leu Gly Ser
            980                 985                 990

Gly Leu Leu Thr Leu Ala Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg
        995                 1000                1005

Gln Ser  Gln Glu Ser Thr Leu  Val His Pro Ala Thr  Pro Asn His
    1010                1015                1020

Leu His  Tyr Lys Gly Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr
    1025                1030                1035

Pro Met  Glu Phe Lys Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp
    1040                1045                1050

Asp Arg  Ala Asn Phe Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr
    1055                1060                1065

Thr Thr  Tyr Tyr Pro Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro
    1070                1075                1080

Glu Ala  Ser Pro Gly Gln Arg  Cys Phe Pro Asn Ser
    1085                1090                1095


<210>  22
<211>  1248
<212>  PRT
<213>  Homo sapiens

<400>  22
Arg Pro Pro His Ser Gln Thr Gly Arg Gln Pro Ile Trp Leu Ala Pro
1               5                   10                  15

Ala Ala Pro Arg Arg Pro Gly Val Gly Ser Arg Gly Glu Pro Gly Thr
            20                  25                  30

Cys Thr Arg Leu Trp Glu Pro Ala Trp Val Arg Val Ala Leu Gly Pro
        35                  40                  45
```

```
Ala Arg Ala Val Val Gly Ala Ser Gly Leu Gln Arg Arg Trp Gly Pro
    50                  55                  60

Gly Thr Gln Ala Trp Arg Arg Arg Arg Thr Ser Glu Ala Glu Gly Arg
65                  70                  75                  80

Arg Asp Arg Val Ser Gly Ser Ser Trp Cys Leu Ala Cys Val Ser Trp
                85                  90                  95

Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His Leu Val Pro
            100                 105                 110

Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly Trp Thr Val
        115                 120                 125

Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro Cys Leu Pro
    130                 135                 140

Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro
145                 150                 155                 160

Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His
                165                 170                 175

Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln
            180                 185                 190

Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln
        195                 200                 205

Pro Trp Val Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln
    210                 215                 220

Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn
225                 230                 235                 240

Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr
                245                 250                 255

Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly
            260                 265                 270

Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala
        275                 280                 285

Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys
    290                 295                 300

Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu Lys Ile Asn
305                 310                 315                 320

Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val
                325                 330                 335

Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser
            340                 345                 350

Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu
        355                 360                 365

Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val
```

370 375 380

Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn
385 390 395 400

Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg
405 410 415

Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp
420 425 430

Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val
435 440 445

Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln
450 455 460

Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala
465 470 475 480

Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn
485 490 495

Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile
500 505 510

Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly
515 520 525

Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu
530 535 540

Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val
545 550 555 560

Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln
565 570 575

Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly
580 585 590

Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys
595 600 605

Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu
610 615 620

Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu
625 630 635 640

Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg
645 650 655

Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp
660 665 670

Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser
675 680 685

Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg
690 695 700

```
Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu
705                 710                 715                 720

Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp
                725                 730                 735

Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile
            740                 745                 750

His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val Ile
        755                 760                 765

Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe Gln Val Arg
    770                 775                 780

Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly Arg Ile Cys
785                 790                 795                 800

Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys
                805                 810                 815

Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser
            820                 825                 830

Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn
        835                 840                 845

Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro
    850                 855                 860

Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu
865                 870                 875                 880

Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg
                885                 890                 895

Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly
            900                 905                 910

Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser
        915                 920                 925

Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr
    930                 935                 940

Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser
945                 950                 955                 960

Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr
                965                 970                 975

Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp
            980                 985                 990

Ala Ala Glu Tyr Gln Asp Cys Asn  Pro Gln Ala Cys Pro  Val Arg Gly
        995                 1000                1005

Ala Trp  Ser Cys Trp Thr Ser  Trp Ser Pro Cys Ser  Ala Ser Cys
    1010                1015                1020
```

```
Gly Gly  Gly His Tyr Gln Arg  Thr Arg Ser Cys Thr  Ser Pro Ala
    1025                 1030                 1035

Pro Ser  Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
    1040                 1045                 1050

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1055                 1060                 1065

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1070                 1075                 1080

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1085                 1090                 1095

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1100                 1105                 1110

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1115                 1120                 1125

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1130                 1135                 1140

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1145                 1150                 1155

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1160                 1165                 1170

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1175                 1180                 1185

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1190                 1195                 1200

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1205                 1210                 1215

Val Tyr  Thr Thr Thr Tyr Tyr  Pro Ser Pro Leu Asn  Lys His Ser
    1220                 1225                 1230

Phe Arg  Pro Glu Ala Ser Pro  Gly Gln Arg Cys Phe  Pro Asn Ser
    1235                 1240                 1245
```

<210> 23
<211> 1150
<212> PRT
<213> Homo sapiens

<400> 23

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala
        35                  40                  45
```

```
Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro Leu Ala Val
    50                  55                  60

Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His Leu Ser Ser
65                  70                  75                  80

Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala
                85                  90                  95

Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln Pro Trp Val
            100                 105                 110

Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln Leu Ala Leu
        115                 120                 125

Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn Tyr Leu Phe
    130                 135                 140

Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr Glu Trp Ala
145                 150                 155                 160

Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu
                165                 170                 175

Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala Gly Arg Lys
            180                 185                 190

Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys Thr Ser Arg
        195                 200                 205

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

Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val Ile Ser Ser
225                 230                 235                 240

Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser Gly Arg Asp
                245                 250                 255

Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala
            260                 265                 270

Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr
        275                 280                 285

Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu
    290                 295                 300

His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg Val Cys Lys
305                 310                 315                 320

Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe
                325                 330                 335

Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr
            340                 345                 350

Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln Asp Leu Ile
        355                 360                 365
```

```
Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala Ser Ala Val
    370                 375                 380

Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe
385                 390                 395                 400

Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro
                405                 410                 415

Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu
            420                 425                 430

Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu Phe Leu Met
        435                 440                 445

Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val Thr Gln Asp
    450                 455                 460

Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln Ala Lys Asp
465                 470                 475                 480

Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu
                485                 490                 495

Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys Tyr Leu Glu
            500                 505                 510

Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu
        515                 520                 525

Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu Arg Asp Gly
    530                 535                 540

Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly
545                 550                 555                 560

Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln
                565                 570                 575

Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser Leu Trp Thr
            580                 585                 590

Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg Asp Gly Gly
        595                 600                 605

Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu Asp Gly Asp
    610                 615                 620

Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg
625                 630                 635                 640

Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile His Ile Ala
                645                 650                 655

Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val His Arg Gly Pro
            660                 665                 670

Leu Cys Ser His Val Leu Trp His Ala Ala Ser Arg Ser Ala Ser Glu
        675                 680                 685

Val Ala Ala Thr Leu Leu Pro Ala Thr Gly Ala Ala Ser Ala Trp Ala
```

690 695 700

Arg Ala Trp Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys Pro Val
705 710 715 720

Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn
725 730 735

Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn
740 745 750

Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro Glu Gly
755 760 765

Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu Pro Val
770 775 780

Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr
785 790 795 800

Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly Arg Arg
805 810 815

Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp
820 825 830

Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr Ser Pro
835 840 845

His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys
850 855 860

Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr Cys Thr
865 870 875 880

Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp Ala Ala
885 890 895

Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro Val Arg Gly Ala Trp
900 905 910

Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala Ser Cys Gly Gly Gly
915 920 925

His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro Ala Pro Ser Pro Gly
930 935 940

Glu Asp Ile Cys Leu Gly Leu His Thr Glu Glu Ala Leu Cys Ala Thr
945 950 955 960

Gln Ala Cys Pro Glu Gly Trp Ser Pro Trp Ser Glu Trp Ser Lys Cys
965 970 975

Thr Asp Asp Gly Ala Gln Ser Arg Ser Arg His Cys Glu Glu Leu Leu
980 985 990

Pro Gly Ser Ser Ala Cys Ala Gly Asn Ser Ser Gln Ser Arg Pro Cys
995 1000 1005

Pro Tyr Ser Glu Ile Arg Val Ile Leu Pro Ala Ser Ser Met Glu
1010 1015 1020

```
Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1025                1030                1035

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1040                1045                1050

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1055                1060                1065

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1070                1075                1080

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
    1085                1090                1095

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1100                1105                1110

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1115                1120                1125

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1130                1135                1140

Gln Arg  Cys Phe Pro Asn Ser
    1145                1150
```

<210> 24
<211> 1211
<212> PRT
<213> Homo sapiens

<400> 24

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140
```

```
Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

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

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
    290                 295                 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
305                 310                 315                 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
                325                 330                 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
            340                 345                 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
        355                 360                 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
    370                 375                 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385                 390                 395                 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
                405                 410                 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
            420                 425                 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
        435                 440                 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
    450                 455                 460
```

```
Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465                 470                 475                 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
                485                 490                 495

Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
            500                 505                 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
        515                 520                 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
    530                 535                 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545                 550                 555                 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
                565                 570                 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
            580                 585                 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
        595                 600                 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
    610                 615                 620

Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625                 630                 635                 640

Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
                645                 650                 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
            660                 665                 670

Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
        675                 680                 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
    690                 695                 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705                 710                 715                 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
                725                 730                 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
            740                 745                 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
        755                 760                 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
    770                 775                 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
```

785 790 795 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
805 810 815

Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
820 825 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
835 840 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
850 855 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865 870 875 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
885 890 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
900 905 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
915 920 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
930 935 940

Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945 950 955 960

Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
965 970 975

Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
980 985 990

Ala Pro Ser Pro Gly Glu Asp Ile Cys Leu Gly Leu His Thr Glu Glu
995 1000 1005

Ala Leu Cys Ala Thr Gln Ala Cys Pro Glu Gly Trp Ser Pro Trp
1010 1015 1020

Ser Glu Trp Ser Lys Cys Thr Asp Asp Gly Ala Gln Ser Arg Ser
1025 1030 1035

Arg His Cys Glu Glu Leu Leu Pro Gly Ser Ser Ala Cys Ala Gly
1040 1045 1050

Asn Ser Ser Gln Ser Arg Pro Cys Pro Tyr Ser Glu Ile Arg Val
1055 1060 1065

Ile Leu Pro Ala Ser Ser Met Glu Glu Ala Thr Asp Cys Ala Gly
1070 1075 1080

Phe Asn Leu Ile His Leu Val Ala Thr Gly Ile Ser Cys Phe Leu
1085 1090 1095

Gly Ser Gly Leu Leu Thr Leu Ala Val Tyr Leu Ser Cys Gln His
1100 1105 1110

```
Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                 1120                 1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1130                 1135                 1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                 1150                 1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                 1165                 1170

Ala Ser  Ala Gly Tyr Pro Pro  Leu Pro Gly Ser Leu  Tyr Ser Thr
    1175                 1180                 1185

Gln Gly  Ile Pro Leu Val Arg  Gly Ser Glu Tyr Trp  Glu Leu Glu
    1190                 1195                 1200

Ala Asp  Leu Cys Leu Glu Val  Leu
    1205                 1210
```

<210> 25
<211> 1203
<212> PRT
<213> Homo sapiens

<400> 25

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140

Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

Asp Leu Gln Pro Trp Val Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp
```

```
                165                 170                 175

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
    290                 295                 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
305                 310                 315                 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
                325                 330                 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
            340                 345                 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
        355                 360                 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
    370                 375                 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385                 390                 395                 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
                405                 410                 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
            420                 425                 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
        435                 440                 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
    450                 455                 460

Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465                 470                 475                 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
                485                 490                 495
```

```
Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
            500                 505                 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
        515                 520                 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
    530                 535                 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545                 550                 555                 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
                565                 570                 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
            580                 585                 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
        595                 600                 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
    610                 615                 620

Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625                 630                 635                 640

Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
                645                 650                 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
            660                 665                 670

Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
        675                 680                 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
    690                 695                 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705                 710                 715                 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
                725                 730                 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
            740                 745                 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
        755                 760                 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
    770                 775                 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
785                 790                 795                 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
                805                 810                 815
```

```
Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
            820                 825                 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
        835                 840                 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
    850                 855                 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865                 870                 875                 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
                885                 890                 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
            900                 905                 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
        915                 920                 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
    930                 935                 940

Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945                 950                 955                 960

Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
                965                 970                 975

Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
            980                 985                 990

Ala Pro Ser Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
        995                 1000                1005

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1010                1015                1020

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1025                1030                1035

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1040                1045                1050

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1055                1060                1065

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1070                1075                1080

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1085                1090                1095

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1100                1105                1110

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                1120                1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
```

```
    1130                1135                1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                1150                1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                1165                1170

Val Tyr  Thr Thr Thr Tyr Tyr  Pro Ser Pro Leu Asn  Lys His Ser
    1175                1180                1185

Phe Arg  Pro Glu Ala Ser Pro  Gly Gln Arg Cys Phe  Pro Asn Ser
    1190                1195                1200


<210>  26
<211>  1240
<212>  PRT
<213>  Homo sapiens

<400>  26
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                   10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Arg Lys Pro
                85                  90                  95

Ser Gly Asp Phe Glu Trp Arg Gln Gly Trp Arg Gly Pro Gly Glu Glu
            100                 105                 110

Asp Trp Pro Glu Ser Pro Ser Pro Lys Val Leu Met Asp Ser Ala Gly
        115                 120                 125

Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro
    130                 135                 140

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

Leu Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser
                165                 170                 175

Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr
            180                 185                 190

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

Gly Ala Arg Asp Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln
```

```
    210                 215                 220

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

Val Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg
                245                 250                 255

Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr
            260                 265                 270

Val Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr
        275                 280                 285

Asn Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser
    290                 295                 300

Arg Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro
305                 310                 315                 320

Arg His Asn Ser Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala
                325                 330                 335

Ala Thr Val Ile Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser
            340                 345                 350

Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp
        355                 360                 365

Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala
    370                 375                 380

Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr
385                 390                 395                 400

Val Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg
                405                 410                 415

Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn
            420                 425                 430

Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser
        435                 440                 445

Ala Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr
    450                 455                 460

Asn Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser
465                 470                 475                 480

Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro
                485                 490                 495

Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys
            500                 505                 510

Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser
        515                 520                 525

Leu Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro
    530                 535                 540
```

```
Val Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His
545                 550                 555                 560

Leu Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu
                565                 570                 575

Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala
            580                 585                 590

Ser Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro
        595                 600                 605

Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala
    610                 615                 620

Arg Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu
625                 630                 635                 640

Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg
                645                 650                 655

Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu
            660                 665                 670

Glu Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys
        675                 680                 685

Pro Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro
    690                 695                 700

Trp Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu
705                 710                 715                 720

Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu
                725                 730                 735

Asp Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly
            740                 745                 750

Gly Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro
        755                 760                 765

Val Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala
    770                 775                 780

Pro Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg
785                 790                 795                 800

Phe Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser
                805                 810                 815

Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln
            820                 825                 830

Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly
        835                 840                 845

Val Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg
    850                 855                 860
```

```
Asn Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly
865                 870                 875                 880

Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala
                885                 890                 895

Asp Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr
            900                 905                 910

Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu
        915                 920                 925

Val Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly
    930                 935                 940

Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu
945                 950                 955                 960

Gly Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn
                965                 970                 975

Gly Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn
            980                 985                 990

Pro Gln Ala Cys Pro Val Arg Gly  Ala Trp Ser Cys Trp  Thr Ser Trp
        995                 1000                1005

Ser Pro  Cys Ser Ala Ser Cys  Gly Gly Gly His Tyr  Gln Arg Thr
    1010                1015                1020

Arg Ser  Cys Thr Ser Pro Ala  Pro Ser Pro Gly Glu  Asp Ile Cys
    1025                1030                1035

Leu Gly  Leu His Thr Glu Glu  Ala Leu Cys Ala Thr  Gln Ala Cys
    1040                1045                1050

Pro Glu  Gly Trp Ser Pro Trp  Ser Glu Trp Ser Lys  Cys Thr Asp
    1055                1060                1065

Asp Gly  Ala Gln Ser Arg Ser  Arg His Cys Glu Glu  Leu Leu Pro
    1070                1075                1080

Gly Ser  Ser Ala Cys Ala Gly  Asn Ser Ser Gln Ser  Arg Pro Cys
    1085                1090                1095

Pro Tyr  Ser Glu Ile Arg Val  Ile Leu Pro Ala Ser  Ser Met Glu
    1100                1105                1110

Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1115                1120                1125

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1130                1135                1140

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1145                1150                1155

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1160                1165                1170

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
```

```
    1175                1180                1185

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1190                1195                1200

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1205                1210                1215

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1220                1225                1230

Gln Arg  Cys Phe Pro Asn Ser
    1235                1240
```

<210> 27
<211> 4567
<212> DNA
<213> Homo sapiens

<400> 27

```
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg     60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt    120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct    180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga    240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc    300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg    360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt    420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa    480
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg    540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc    600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc    660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg    720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc    780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc    840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca    900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg    960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca   1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac   1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt ggggggccga   1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg   1200
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac   1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc   1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc   1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct   1440
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc   1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg   1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc   1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc   1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc   1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg   1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg   1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc   1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat   1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca   2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt   2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg   2160
tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga   2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag   2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc   2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc   2400
```

gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc 2460
tgccccgaag tgcggcgcaa caccccctgg acgccgtggc tgcccgtgaa cgtgacgcag 2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgccccc tgcagacccg 2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc 2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg 2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc 2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc 2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt 2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt 2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt 3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg 3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt 3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc 3180
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt 3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg 3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc 3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag 3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat 3480
gacagagcca acttctaccc attgcagcag accaatgtgt acacgactac ttactaccca 3540
agccccctga acaaacacag cttccggccc gaggcctcac ctggacaacg gtgcttcccc 3600
aacagctgat accgccgtcc tggggacttg ggcttcttgc cttcataagg cacagagcag 3660
atggagatgg gacagtggag ccagtttggt tttctccctc tgcactaggc caagaacttg 3720
ctgccttgcc tgtggggggt cccatccggc ttcagagagc tctggctggc attgaccatg 3780
ggggaaaggg ctggtttcag gctgacatat ggccgcaggt ccagttcagc ccaggtctmt 3840
catggttatc ttccaaccca ctgtcacgct gacactatgc tgccatgcct gggctgtgga 3900
cctactgggc atttgaggaa ytggagaatg gagatggcaa gagggcaggc ttttaagttt 3960
gggttggaga caacttcctg tggcccccac aagctgagtc tggccttctc cagctggccc 4020
caaaaaaggc ctttgctaca tcctgattat ctctgaaagt aatcaatcaa gtggctccag 4080
tagctctgga ttttctgcca gggctgggcc attgtggtgc tgccccagta tgacatggga 4140
ccaaggccag cgcaggttat ccacctctgc ctggaagtct atactctacc cagggcatcc 4200
ctctggtcag aggcagtgag tactgggaac tggaggctga cctgtgctta gaagtccttt 4260
aatctgggct ggtacaggcc tcagccttgc cctcaatgca cgaaaggtgg cccaggagag 4320
aggatcaatg ccataggagg cagaagtctg gcctctgtgc ctctatggag actatcttcc 4380
agttgctgct caacagagtt gttggctgag acctgcttgg gagtctctgc tggcccttca 4440
tctgttcagg aacacacaca cacacacact cacacacgca cacacaatca caatttgcta 4500
cagcaacaaa aaagacattg ggctgtggca ttattaatta aagatgatat ccagtcaaaa 4560
aaaaact 4567

<210> 28
<211> 453
<212> DNA
<213> Homo sapiens

<400> 28
agtaatcagc tcggtaccgg catgtgctgt agccagcgca ggttatccac ctctgcctgg 60
aagtctatac tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga 120
ggctgacctg tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc 180
aatgcacgaa aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct 240
ctgtgcctct atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct 300
gcttgggagt ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca 360
cacgcacaca caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat 420
taattaaaga tgatatccag tcaaaaaaaa act 453

<210> 29
<211> 4675
<212> DNA
<213> Homo sapiens

<400> 29

```
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg     60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt    120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct    180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga    240
tggacagtag ggggctggct tctctcactg gtcaggggca ggaagccaag tggagacttt    300
gaatggaggc aaggatggag gggacctggg gaagaggact ggcctgaatc accttcccca    360
aaggtcctca tggactgagc tggaggtctt ctcccctgtc tgcctcccgg agctaggact    420
gcagaggggc ctatcatggt gcttgcaggc cccctggctg tctcgctgtt gctgcccagc    480
ctcacactgc tggtgtccca cctctccagc tcccaggatg tctccagtga gcccagcagt    540
gagcagcagc tgtgcgccct tagcaagcac cccaccgtgg cctttgaaga cctgcagccg    600
tgggtctcta acttcaccta ccctggagcc cgggatttct cccagctggc tttggacccc    660
tccgggaacc agctcatcgt gggagccagg aactacctct tcagactcag ccttgccaat    720
gtctctcttc ttcaggccac agagtgggcc tccagtgagg acacgcgccg ctcctgccaa    780
agcaaaggga agactgagga ggagtgtcag aactacgtgc gagtcctgat cgtcgccggc    840
cggaaggtgt tcatgtgtgg aaccaatgcc ttttccccca tgtgcaccag cagacaggtg    900
gggaacctca gccggactac tgagaagatc aatggtgtgg cccgctgccc ctatgaccca    960
cgccacaact ccacagctgt catctcctcc caggggagc tctatgcagc cacggtcatc   1020
gacttctcag gtcgggaccc tgccatctac cgcagcctgg gcagtgggcc accgcttcgc   1080
actgcccaat ataactccaa gtggcttaat gagccaaact tcgtggcagc ctatgatatt   1140
gggctgtttg catacttctt cctgcgggag aacgcagtgg agcacgactg tggacgcacc   1200
gtgtactctc gcgtggcccg cgtgtgcaag aatgacgtgg ggggccgatt cctgctggag   1260
gacacatgga ccacattcat gaaggcccgg ctcaactgct cccgcccggg cgaggtcccc   1320
ttctactata acgagctgca gagtgccttc cacttgccag agcaggacct catctatgga   1380
gttttcacaa ccaacgtaaa cagcatcgcg gcttctgctg tctgcgcctt caacctcagt   1440
gctatctccc aggctttcaa tggcccattt cgctaccagg agaaccccag ggctgcctgg   1500
ctccccatag ccaaccccat ccccaatttc cagtgtggca ccctgcctga gaccggtccc   1560
aacgagaacc tgacggagcg cagcctgcag gacgcgcagc gcctcttcct gatgagcgag   1620
gccgtgcagc cggtgacacc cgagccctgt gtcacccagg acagcgtgcg cttctcacac   1680
ctcgtggtgg acctggtgca ggctaaagac acgctctacc atgtactcta cattggcacc   1740
gagtcgggca ccatcctgaa ggcgctgtcc acggcgagcc gcagcctcca cggctgctac   1800
ctggaggagc tgcacgtgct gccccccggg cgccgcgagc ccctgcgcag cctgcgcatc   1860
ctgcacagcg cccgcgcgct cttcgtgggg ctgagagacg gcgtcctgcg ggtcccactg   1920
gagaggtgcg ccgcctaccg cagccagggg gcatgcctgg gggcccggga cccgtactgt   1980
ggctgggacg ggaagcagca acgttgcagc acactcgagg acagctccaa catgagcctc   2040
tggacccaga acatcaccgc ctgtcctgtg cggaatgtga cacgggatgg gggcttcggc   2100
ccatggtcac catggcaacc atgtgagcac ttggatgggg acaactcagg ctcttgcctg   2160
tgtcgagctc gatcctgtga ttcccctcga ccccgctgtg ggggccttga ctgcctgggg   2220
ccagccatcc acatcgccaa ctgctccagg aatggggcgt ggaccccgtg gtcatcgtgg   2280
gcgctgtgca gcacgtcctg tggcatcggc ttccaggtcc gccagcgaag ttgcagcaac   2340
cctgctcccc gccacggggg ccgcatctgc gtgggcaaga gccgggagga acggttctgt   2400
aatgagaaca cgccttgccc ggtgcccatc ttctgggctt cctggggctc ctggagcaag   2460
tgcagcagca actgtggagg gggcatgcag tcgcggcgtc gggcctgcga gaacggcaac   2520
tcctgcctgg gctgcggcgt ggagttcaag acgtgcaacc ccgagggctg ccccgaagtg   2580
cggcgcaaca ccccctggac gccgtggctg cccgtgaacg tgacgcaggg cggggcacgg   2640
caggagcagc ggttccgctt cacctgccgc gcgccccttg cagacccgca cggcctgcag   2700
ttcggcagga gaaggaccga gacgaggacc tgtcccgcgg acggctccgg ctcctgcgac   2760
accgacgccc tggtggaggt cctcctgcgc agcgggagca cctccccgca cacggtgagc   2820
gggggctggg ccgcctgggg cccgtggtcg tcctgctccc gggactgcga gctgggcttc   2880
cgcgtccgca agagaacgtg cactaacccg gagccccgca acgggggcct gccctgcgtg   2940
ggcgatgctg ccgagtacca ggactgcaac ccccaggctt gcccagttcg gggtgcttgg   3000
tcctgctgga cctcatggtc tccatgctca gcttcctgtg gtgggggtca ctatcaacgc   3060
acccgttcct gcaccagccc cgcaccctcc ccaggtgagg acatctgtct cgggctgcac   3120
acggaggagg cactatgtgc cacacaggcc tgcccagaag gctggtcgcc ctggtctgag   3180
tggagtaagt gcactgacga cggagcccag agccgaagcc ggcactgtga ggagctcctc   3240
ccagggtcca gcgcmtgtgc tggaaacagc agccagagcc gcccctgccc ctacagcgag   3300
attcscgtca tcctgccagc ctccagcatg gaggaggcca ccgactgtgc agggttcaat   3360
ctcatccact tggtggccac gggcatctcc tgcttcttgg gctctgggct cctgacccta   3420
gcagtgtacc tgtcttgcca gcactgccag cgtcagtccc aggagtccac actggtccat   3480
cctgccaccc ccaaccattt gcactacaag ggcggaggca ccccgaagaa tgaaaagtac   3540
acacccatgg aattcaagac cctgaacaag aataacttga tccctgatga cagagccaac   3600
ttctacccat tgcagcagac caatgtgtac acgactactt actacccaag ccccctgaac   3660
```

aaacacagct tccggcccga ggcctcacct ggacaacggt gcttccccaa cagctgatac 3720
cgccgtcctg gggacttggg cttcttgcct tcataaggca cagagcagat ggagatggga 3780
cagtggagcc agtttggttt tctccctctg cactaggcca agaacttgct gccttgcctg 3840
tgggggtcc catccggctt cagagagctc tggctggcat tgaccatggg ggaaagggct 3900
ggtttcaggc tgacatatgg ccgcaggtcc agttcagccc aggtctmtca tggttatctt 3960
ccaacccact gtcacgctga cactatgctg ccatgcctgg gctgtggacc tactgggcat 4020
ttgaggaayt ggagaatgga gatggcaaga gggcaggctt ttaagtttgg gttggagaca 4080
acttcctgtg gcccccacaa gctgagtctg gccttctcca gctggcccca aaaaaggcct 4140
ttgctacatc ctgattatct ctgaaagtaa tcaatcaagt ggctccagta gctctggatt 4200
ttctgccagg gctgggccat tgtggtgctg ccccagtatg acatgggacc aaggccagcg 4260
caggttatcc acctctgcct ggaagtctat actctaccca gggcatccct ctggtcagag 4320
gcagtgagta ctgggaactg gaggctgacc tgtgcttaga agtcctttaa tctgggctgg 4380
tacaggcctc agccttgccc tcaatgcacg aaaggtggcc caggagagag gatcaatgcc 4440
ataggaggca gaagtctggc ctctgtgcct ctatggagac tatcttccag ttgctgctca 4500
acagagttgt tggctgagac ctgcttggga gtctctgctg gcccttcatc tgttcaggaa 4560
cacacacaca cacacactca cacacgcaca cacaatcaca atttgctaca gcaacaaaaa 4620
agacattggg ctgtggcatt attaattaaa gatgatatcc agtcaaaaaa aaact 4675

<210> 30
<211> 4731
<212> DNA
<213> Homo sapiens

<400> 30
attggagatg ctcgggggca ggctgccgcg ttgtgtcctg cttttctgcg gccagaccaa 60
gccgtctgga gctgctggtc aggttttctt gctgacctca cctgaccaca gtggcctggg 120
tggactctac agggaaatgt tgttttctcc ctgggagcag tagcagcagt cctggctccc 180
ctggactgag aactcctcat cagccccagg aagcccggac ccccttcag ggatctggaa 240
ccggtgtgcc tgtggcccca ggtctgctcc caggcgtggg ctgaagtcct gacttctgtc 300
gctgggggca aggagtggga gagcccagct gctgcctggg ctttggcaga cagcaggctg 360
atggtgctgg cttccccgag actgcttctc ctgcctgctg tctgatttcc ctgcatggtg 420
cccgcagctg agctgctacg ggtcttctcc cctgtctgcc tcccggagct aggactgcag 480
aggggcctat catggtgctt gcaggccccc tggctgtctc gctgttgctg cccagcctca 540
cactgctggt gtcccacctc tccagctccc aggatgtctc cagtgagccc agcagtgagc 600
agcagctgtg cgcccttagc aagcacccca ccgtggcctt tgaagacctg cagccgtggg 660
tctctaactt cacctaccct ggagcccggg atttctccca gctggctttg gacccctccg 720
ggaaccagct catcgtggga gccaggaact acctcttcag actcagcctt gccaatgtct 780
ctcttcttca ggccacagag tgggcctcca gtgaggacac gcgccgctcc tgccaaagca 840
aagggaagac tgaggaggag tgtcagaact acgtgcgagt cctgatcgtc gccggccgga 900
aggtgttcat gtgtggaacc aatgcctttt cccccatgtg caccagcaga caggtgggga 960
acctcagccg gactactgag aagatcaatg gtgtggcccg ctgcccctat gacccacgcc 1020
acaactccac agctgtcatc tcctcccagg gggagctcta tgcagccacg gtcatcgact 1080
tctcaggtcg ggaccctgcc atctaccgca gcctgggcag tgggccaccg cttcgcactg 1140
cccaatataa ctccaagtgg cttaatgagc caaacttcgt ggcagcctat gatattgggc 1200
tgtttgcata cttcttcctg cgggagaacg cagtggagca cgactgtgga cgcaccgtgt 1260
actctcgcgt ggcccgcgtg tgcaagaatg acgtgggggg ccgattcctg ctggaggaca 1320
catggaccac attcatgaag gcccggctca actgctcccg cccgggcgag gtccccttct 1380
actataacga gctgcagagt gccttccact tgccagagca ggacctcatc tatggagttt 1440
tcacaaccaa cgtaaacagc atcgcggctt ctgctgtctg cgccttcaac ctcagtgcta 1500
tctcccaggc tttcaatggc ccatttcgct accaggagaa ccccagggct gcctggctcc 1560
ccatagccaa ccccatcccc aatttccagt gtggcaccct gcctgagacc ggtcccaacg 1620
agaacctgac ggagcgcagc ctgcaggacg cgcagcgcct cttcctgatg agcgaggccg 1680
tgcagccggt gacacccgag ccctgtgtca cccaggacag cgtgcgcttc tcacacctcg 1740
tggtggacct ggtgcaggct aaagacacgc tctaccatgt actctacatt ggcaccgagt 1800
cgggcaccat cctgaaggcg ctgtccacgg cgagccgcag cctccacggc tgctacctgg 1860
aggagctgca cgtgctgccc cccgggcgcc gcgagcccct gcgcagcctg cgcatcctgc 1920
acagcgcccg cgcgctcttc gtggggctga gagacggcgt cctgcgggtc ccactggaga 1980
ggtgcgccgc ctaccgcagc caggggggcat gcctgggggc ccgggacccg tactgtggct 2040
gggacgggaa gcagcaacgt tgcagcacac tcgaggacag ctccaacatg agcctctgga 2100
cccagaacat caccgcctgt cctgtgcgga atgtgacacg ggatgggggc ttcggcccat 2160

ggtcaccatg gcaaccatgt gagcacttgg atggggacaa ctcaggctct tgcctgtgtc 2220
gagctcgatc ctgtgattcc cctcgacccc gctgtggggg ccttgactgc ctggggccag 2280
ccatccacat cgccaactgc tccaggaatg gggcgtggac cccgtggtca tcgtgggcgc 2340
tgtgcagcac gtcctgtggc atcggcttcc aggtccgcca gcgaagttgc agcaaccctg 2400
ctccccgcca cgggggccgc atctgcgtgg gcaagagccg ggaggaacgg ttctgtaatg 2460
agaacacgcc ttgcccggtg cccatcttct gggcttcctg gggctcctgg agcaagtgca 2520
gcagcaactg tggagggggc atgcagtcgc ggcgtcgggc ctgcgagaac ggcaactcct 2580
gcctgggctg cggcgtggag ttcaagacgt gcaaccccga gggctgcccc gaagtgcggc 2640
gcaacacccc ctggacgccg tggctgcccg tgaacgtgac gcagggcggg gcacggcagg 2700
agcagcggtt ccgcttcacc tgccgcgcgc cccttgcaga cccgcacggc ctgcagttcg 2760
gcaggagaag gaccgagacg aggacctgtc ccgcggacgg ctccggctcc tgcgacaccg 2820
acgccctggt ggaggtcctc ctgcgcagcg ggagcacctc cccgcacacg gtgagcgggg 2880
gctgggccgc ctggggcccg tggtcgtcct gctcccggga ctgcgagctg ggcttccgcg 2940
tccgcaagag aacgtgcact aacccggagc cccgcaacgg gggcctgccc tgcgtgggcg 3000
atgctgccga gtaccaggac tgcaaccccc aggcttgccc agttcggggt gcttggtcct 3060
gctggacctc atggtctcca tgctcagctt cctgtggtgg gggtcactat caacgcaccc 3120
gttcctgcac cagccccgca ccctccccag gtgaggacat ctgtctcggg ctgcacacgg 3180
aggaggcact atgtgccaca caggcctgcc cagaaggctg gtcgccctgg tctgagtgga 3240
gtaagtgcac tgacgacgga gcccagagcc gaagccggca ctgtgaggag ctcctcccag 3300
ggtccagcgc mtgtgctgga aacagcagcc agagccgccc ctgcccctac agcgagattc 3360
scgtcatcct gccagcctcc agcatggagg aggccaccga ctgtgcaggg ttcaatctca 3420
tccacttggt ggccacgggc atctcctgct tcttgggctc tgggctcctg accctagcag 3480
tgtacctgtc ttgccagcac tgccagcgtc agtcccagga gtccacactg gtccatcctg 3540
ccacccccaa ccatttgcac tacaagggcg gaggcacccc gaagaatgaa aagtacacac 3600
ccatggaatt caagaccctg aacaagaata acttgatccc tgatgacaga gccaacttct 3660
acccattgca gcagaccaat gtgtacacga ctacttacta cccaagcccc ctgaacaaac 3720
acagcttccg gcccgaggcc tcacctggac aacggtgctt ccccaacagc tgataccgcc 3780
gtcctgggga cttgggcttc ttgccttcat aaggcacaga gcagatggag atgggacagt 3840
ggagccagtt tggttttctc cctctgcact aggccaagaa cttgctgcct tgcctgtggg 3900
gggtcccatc cggcttcaga gagctctggc tggcattgac catgggggaa agggctggtt 3960
tcaggctgac atatggccgc aggtccagtt cagcccaggt ctmtcatggt tatcttccaa 4020
cccactgtca cgctgacact atgctgccat gcctgggctg tggacctact gggcatttga 4080
ggaaytggag aatggagatg gcaagagggc aggcttttaa gtttgggttg gagacaactt 4140
cctgtggccc ccacaagctg agtctggcct tctccagctg gccccaaaaa aggcctttgc 4200
tacatcctga ttatctctga aagtaatcaa tcaagtggct ccagtagctc tggattttct 4260
gccagggctg ggccattgtg gtgctgcccc agtatgacat gggaccaagg ccagcgcagg 4320
ttatccacct ctgcctggaa gtctatactc tacccagggc atccctctgg tcagaggcag 4380
tgagtactgg gaactggagg ctgacctgtg cttagaagtc ctttaatctg ggctggtaca 4440
ggcctcagcc ttgccctcaa tgcacgaaag gtggcccagg agagaggatc aatgccatag 4500
gaggcagaag tctggcctct gtgcctctat ggagactatc ttccagttgc tgctcaacag 4560
agttgttggc tgagacctgc ttgggagtct ctgctggccc ttcatctgtt caggaacaca 4620
cacacacaca cactcacaca cgcacacaca atcacaattt gctacagcaa caaaaaagac 4680
attgggctgt ggcattatta attaaagatg atatccagtc aaaaaaaaac t 4731

<210> 31
<211> 4703
<212> DNA
<213> Homo sapiens

<400> 31
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagccagc 120
ttgggtccgg gttgcactgg gccctgccag ggctgtggtc ggcgcatctg ggctgcagcg 180
gcgatgggga cccgggaccc aggcctggag aaggagacgg acgagtgagg ctgagggacg 240
gagggacaga gtgagtggtt ccagctggtg cctggcctgt gtctcttgga tgccctgtgg 300
cttcagtccg tctcctgttg cccaccacct cgtccctggg ccgcctgata ccccagccca 360
acagctaagg tgtggatgga cagtaggggg ctggcttctc tcactggtca ggggtcttct 420
cccctgtctg cctcccggag ctaggactgc agaggggcct atcatggtgc ttgcaggccc 480
cctggctgtc tcgctgttgc tgcccagcct cacactgctg gtgtcccacc tctccagctc 540
ccaggatgtc tccagtgagc ccagcagtga gcagcagctg tgcgccctta gcaagcaccc 600

caccgtggcc tttgaagacc tgcagccgtg ggtctctaac ttcacctacc ctggagcccg 660
ggatttctcc cagctggctt tggacccctc cgggaaccag ctcatcgtgg gagccaggaa 720
ctacctcttc agactcagcc ttgccaatgt ctctcttctt caggccacag agtgggcctc 780
cagtgaggac acgcgccgct cctgccaaag caaagggaag actgaggagg agtgtcagaa 840
ctacgtgcga gtcctgatcg tcgccggccg gaaggtgttc atgtgtggaa ccaatgcctt 900
ttcccccatg tgcaccagca gacaggtggg gaacctcagc cggactactg agaagatcaa 960
tggtgtggcc cgctgccccct atgacccacg ccacaactcc acagctgtca tctcctccca 1020
gggggagctc tatgcagcca cggtcatcga cttctcaggt cgggaccctg ccatctaccg 1080
cagcctgggc agtgggccac cgcttcgcac tgcccaatat aactccaagt ggcttaatga 1140
gccaaacttc gtggcagcct atgatattgg gctgtttgca tacttcttcc tgcgggagaa 1200
cgcagtggag cacgactgtg gacgcaccgt gtactctcgc gtggcccgcg tgtgcaagaa 1260
tgacgtgggg ggccgattcc tgctggagga cacatggacc acattcatga aggcccggct 1320
caactgctcc cgcccgggcg aggtcccctt ctactataac gagctgcaga gtgccttcca 1380
cttgccagag caggacctca tctatggagt tttcacaacc aacgtaaaca gcatcgcggc 1440
ttctgctgtc tgcgccttca acctcagtgc tatctcccag gctttcaatg gcccatttcg 1500
ctaccaggag aaccccaggg ctgcctggct ccccatagcc aaccccatcc ccaatttcca 1560
gtgtggcacc ctgcctgaga ccggtcccaa cgagaacctg acggagcgca gcctgcagga 1620
cgcgcagcgc ctcttcctga tgagcgaggc cgtgcagccg gtgacacccg agccctgtgt 1680
cacccaggac agcgtgcgct tctcacacct cgtggtggac ctggtgcagg ctaaagacac 1740
gctctaccat gtactctaca ttggcaccga gtcgggcacc atcctgaagg cgctgtccac 1800
ggcgagccgc agcctccacg gctgctacct ggaggagctg cacgtgctgc cccccgggcg 1860
ccgcgagccc ctgcgcagcc tgcgcatcct gcacagcgcc cgcgcgctct tcgtggggct 1920
gagagacggc gtcctgcggg tcccactgga gaggtgcgcc gcctaccgca gccaggggc 1980
atgcctgggg gcccgggacc cgtactgtgg ctgggacggg aagcagcaac gttgcagcac 2040
actcgaggac agctccaaca tgagcctctg gacccagaac atcaccgcct gtcctgtgcg 2100
gaatgtgaca cgggatgggg gcttcggccc atggtcacca tggcaaccat gtgagcactt 2160
ggatggggac aactcaggct cttgcctgtg tcgagctcga tcctgtgatt cccctcgacc 2220
ccgctgtggg ggccttgact gcctggggcc agccatccac atcgccaact gctccaggaa 2280
tggggcgtgg accccgtggt catcgtgggc gctgtgcagc acgtcctgtg gcatcggctt 2340
ccaggtccgc cagcgaagtt gcagcaaccc tgctccccgc cacgggggcc gcatctgcgt 2400
gggcaagagc cgggaggaac ggttctgtaa tgagaacacg ccttgcccgg tgcccatctt 2460
ctgggcttcc tggggctcct ggagcaagtg cagcagcaac tgtggagggg gcatgcagtc 2520
gcggcgtcgg gcctgcgaga acggcaactc ctgcctgggc tgcggcgtgg agttcaagac 2580
gtgcaacccc gagggctgcc ccgaagtgcg gcgcaacacc ccctggacgc cgtggctgcc 2640
cgtgaacgtg acgcagggcg gggcacggca ggagcagcgg ttccgcttca cctgccgcgc 2700
gccccttgca gacccgcacg gcctgcagtt cggcaggaga aggaccgaga cgaggacctg 2760
tcccgcggac ggctccggct cctgcgacac cgacgccctg gtggaggtcc tcctgcgcag 2820
cgggagcacc tccccgcaca cggtgagcgg gggctgggcc gcctggggcc cgtggtcgtc 2880
ctgctcccgg gactgcgagc tgggcttccg cgtccgcaag agaacgtgca ctaacccgga 2940
gccccgcaac gggggcctgc cctgcgtggg cgatgctgcc gagtaccagg actgcaaccc 3000
ccaggcttgc ccagttcggg gtgcttggtc ctgctggacc tcatggtctc catgctcagc 3060
ttcctgtggt gggggtcact atcaacgcac ccgttcctgc accagccccg caccctcccc 3120
aggtgaggac atctgtctcg ggctgcacac ggaggaggca ctatgtgcca cacaggcctg 3180
cccagaaggc tggtcgccct ggtctgagtg gagtaagtgc actgacgacg gagcccagag 3240
ccgaagccgg cactgtgagg agctcctccc agggtccagc gcmtgtgctg gaaacagcag 3300
ccagagccgc ccctgccct acagcgagat tcscgtcatc ctgccagcct ccagcatgga 3360
ggaggccacc gactgtgcag ggttcaatct catccacttg gtggccacgg gcatctcctg 3420
cttcttgggc tctgggctcc tgaccctagc agtgtacctg tcttgccagc actgccagcg 3480
tcagtcccag gagtccacac tggtccatcc tgccaccccc aaccatttgc actacaaggg 3540
cggaggcacc ccgaagaatg aaaagtacac acccatggaa ttcaagaccc tgaacaagaa 3600
taacttgatc cctgatgaca gagccaactt ctacccattg cagcagacca atgtgtacac 3660
gactacttac tacccaagcc ccctgaacaa acacagcttc cggcccgagg cctcacctgg 3720
acaacggtgc ttccccaaca gctgataccg ccgtcctggg gacttgggct tcttgccttc 3780
ataaggcaca gagcagatgg agatgggaca gtggagccag tttggttttc tccctctgca 3840
ctaggccaag aacttgctgc cttgcctgtg gggggtccca tccggcttca gagagctctg 3900
gctggcattg accatggggg aaagggctgg tttcaggctg acatatggcc gcaggtccag 3960
ttcagcccag gtctmtcatg gttatcttcc aacccactgt cacgctgaca ctatgctgcc 4020
atgcctgggc tgtggaccta ctgggcattt gaggaaytgg agaatggaga tggcaagagg 4080
gcaggctttt aagtttgggt tggagacaac ttcctgtggc ccccacaagc tgagtctggc 4140
cttctccagc tggccccaaa aaaggccttt gctacatcct gattatctct gaaagtaatc 4200
aatcaagtgg ctccagtagc tctggatttt ctgccagggc tgggccattg tggtgctgcc 4260

ccagtatgac atgggaccaa ggccagcgca ggttatccac ctctgcctgg aagtctatac 4320
tctacccagg gcatccctct ggtcagaggc agtgagtact gggaactgga ggctgacctg 4380
tgcttagaag tcctttaatc tgggctggta caggcctcag ccttgccctc aatgcacgaa 4440
aggtggccca ggagagagga tcaatgccat aggaggcaga agtctggcct ctgtgcctct 4500
atggagacta tcttccagtt gctgctcaac agagttgttg gctgagacct gcttgggagt 4560
ctctgctggc ccttcatctg ttcaggaaca cacacacaca cacactcaca cacgcacaca 4620
caatcacaat ttgctacagc aacaaaaaag acattgggct gtggcattat taattaaaga 4680
tgatatccag tcaaaaaaaa act 4703

<210> 32
<211> 4405
<212> DNA
<213> Homo sapiens

<400> 32
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggaggtctt 120
ctcccctgtc tgcctcccgg agctaggact gcagaggggc ctatcatggt gcttgcaggc 180
cccctggctg tctcgctgtt gctgcccagc ctcacactgc tggtgtccca cctctccagc 240
tcccaggatg tctccagtga gcccagcagt gagcagcagc tgtgcgccct tagcaagcac 300
cccaccgtgg cctttgaaga cctgcagccg tgggtctcta acttcaccta ccctggagcc 360
cgggatttct cccagctggc tttggacccc tccgggaacc agctcatcgt gggagccagg 420
aactacctct tcagactcag ccttgccaat gtctctcttc ttcaggccac agagtgggcc 480
tccagtgagg acacgcgccg ctcctgccaa agcaaaggga agactgagga ggagtgtcag 540
aactacgtgc gagtcctgat cgtcgccggc cggaaggtgt tcatgtgtgg aaccaatgcc 600
ttttccccca tgtgcaccag cagacaggtg gggaacctca gccggactac tgagaagatc 660
aatggtgtgg cccgctgccc ctatgaccca cgccacaact ccacagctgt catctcctcc 720
caggggagc tctatgcagc cacggtcatc gacttctcag gtcgggaccc tgccatctac 780
cgcagcctgg gcagtgggcc accgcttcgc actgcccaat ataactccaa gtggcttaat 840
gagccaaact tcgtggcagc ctatgatatt gggctgtttg catacttctt cctgcgggag 900
aacgcagtgg agcacgactg tggacgcacc gtgtactctc gcgtggcccg cgtgtgcaag 960
aatgacgtgg ggggccgatt cctgctggag gacacatgga ccacattcat gaaggcccgg 1020
ctcaactgct cccgcccggg cgaggtcccc ttctactata acgagctgca gagtgccttc 1080
cacttgccag agcaggacct catctatgga gttttcacaa ccaacgtaaa cagcatcgcg 1140
gcttctgctg tctgcgcctt caacctcagt gctatctccc aggctttcaa tggcccattt 1200
cgctaccagg agaaccccag ggctgcctgg ctccccatag ccaaccccat ccccaatttc 1260
cagtgtggca ccctgcctga gaccggtccc aacgagaacc tgacggagcg cagcctgcag 1320
gacgcgcagc gcctcttcct gatgagcgag gccgtgcagc cggtgacacc cgagccctgt 1380
gtcacccagg acagcgtgcg cttctcacac ctcgtggtgg acctggtgca ggctaaagac 1440
acgctctacc atgtactcta cattggcacc gagtcgggca ccatcctgaa ggcgctgtcc 1500
acggcgagcc gcagcctcca cggctgctac ctggaggagc tgcacgtgct gccccccggg 1560
cgccgcgagc ccctgcgcag cctgcgcatc ctgcacagcg cccgcgcgct cttcgtgggg 1620
ctgagagacg gcgtcctgcg ggtcccactg gagaggtgcg ccgcctaccg cagccagggg 1680
gcatgcctgg gggcccggga cccgtactgt ggctgggacg ggaagcagca acgttgcagc 1740
acactcgagg acagctccaa catgagcctc tggacccaga acatcaccgc ctgtcctgtg 1800
cggaatgtga cacgggatgg gggcttcggc ccatggtcac catggcaacc atgtgagcac 1860
ttggatgggg acaactcagg ctcttgcctg tgtcgagctc gatcctgtga ttcccctcga 1920
ccccgctgtg ggggccttga ctgcctgggg ccagccatcc acatcgccaa ctgctccagg 1980
aatggggcgt ggaccccgtg gtcatcgtgg gcgctgtgca gcacgtcctg tggcatcggc 2040
ttccaggtcc gccagcgaag ttgcagcaac cctgctcccc gccacggggg ccgcatctgc 2100
gtgggcaaga gccgggagga acggttctgt aatgagaaca cgccttgccc ggtgcccatc 2160
ttctgggctt cctggggctc ctggagcaag tgcagcagca actgtggagg gggcatgcag 2220
tcgcggcgtc gggcctgcga gaacggcaac tcctgcctgg gctgcggcgt ggagttcaag 2280
acgtgcaacc ccgagggctg ccccgaagtg cggcgcaaca ccccctggac gccgtggctg 2340
cccgtgaacg tgacgcaggg cggggcacgg caggagcagc ggttccgctt cacctgccgc 2400
gcgccccttg cagacccgca cggcctgcag ttcggcagga gaaggaccga gacgaggacc 2460
tgtcccgcgg acggctccgg ctcctgcgac accgacgccc tggtggaggt cctcctgcgc 2520
agcgggagca cctccccgca cacggtgagc gggggctggg ccgcctgggg cccgtggtcg 2580
tcctgctccc gggactgcga gctgggcttc cgcgtccgca agagaacgtg cactaacccg 2640
gagccccgca acgggggcct gccctgcgtg ggcgatgctg ccgagtacca ggactgcaac 2700

ccccaggctt gcccagttcg gggtgcttgg tcctgctgga cctcatggtc tccatgctca 2760
gcttcctgtg gtgggggtca ctatcaacgc acccgttcct gcaccagccc cgcaccctcc 2820
ccaggtgagg acatctgtct cgggctgcac acggaggagg cactatgtgc cacacaggcc 2880
tgcccagaag gctggtcgcc ctggtctgag tggagtaagt gcactgacga cggagcccag 2940
agccgaagcc ggcactgtga ggagctcctc ccagggtcca gcgcmtgtgc tggaaacagc 3000
agccagagcc gcccctgccc ctacagcgag attcscgtca tcctgccagc ctccagcatg 3060
gaggaggcca ccgactgtgc agggttcaat ctcatccact tggtggccac gggcatctcc 3120
tgcttcttgg gctctgggct cctgacccta gcagtgtacc tgtcttgcca gcactgccag 3180
cgtcagtccc aggagtccac actggtccat cctgccaccc ccaaccattt gcactacaag 3240
ggcggaggca ccccgaagaa tgaaaagtac acacccatgg aattcaagac cctgaacaag 3300
aataacttga tccctgatga cagagccaac ttctacccat tgcagcagac caatgtgtac 3360
acgactactt actacccaag ccccctgaac aaacacagct tccggcccga ggcctcacct 3420
ggacaacggt gcttccccaa cagctgatac cgccgtcctg gggacttggg cttcttgcct 3480
tcataaggca cagagcagat ggagatggga cagtggagcc agtttggttt tctccctctg 3540
cactaggcca agaacttgct gccttgcctg tgggggtcc catccggctt cagagagctc 3600
tggctggcat tgaccatggg ggaaagggct ggtttcaggc tgacatatgg ccgcaggtcc 3660
agttcagccc aggtctmtca tggttatctt ccaacccact gtcacgctga cactatgctg 3720
ccatgcctgg gctgtggacc tactgggcat ttgaggaayt ggagaatgga gatggcaaga 3780
gggcaggctt ttaagtttgg gttggagaca acttcctgtg gcccccacaa gctgagtctg 3840
gccttctcca gctggcccca aaaaaggcct ttgctacatc ctgattatct ctgaaagtaa 3900
tcaatcaagt ggctccagta gctctggatt ttctgccagg gctgggccat tgtggtgctg 3960
ccccagtatg acatgggacc aaggccagcg caggttatcc acctctgcct ggaagtctat 4020
actctaccca gggcatccct ctggtcagag gcagtgagta ctgggaactg gaggctgacc 4080
tgtgcttaga agtcctttaa tctgggctgg tacaggcctc agccttgccc tcaatgcacg 4140
aaaggtggcc caggagagag gatcaatgcc ataggaggca gaagtctggc ctctgtgcct 4200
ctatggagac tatcttccag ttgctgctca acagagttgt tggctgagac ctgcttggga 4260
gtctctgctg gcccttcatc tgttcaggaa cacacacaca cacacactca cacacgcaca 4320
cacaatcaca atttgctaca gcaacaaaaa agacattggg ctgtggcatt attaattaaa 4380
gatgatatcc agtcaaaaaa aaact 4405

<210> 33
<211> 3938
<212> DNA
<213> Homo sapiens

<400> 33
gcggccgccc cattcccaga ccggccgcca gcccatctgg ttagctcccg ccgctccgcg 60
ccgcccggga gtcgggagcc gcggggaacc gggcacctgc acccgcctct gggagtgagt 120
ggttccagct ggtgcctggc ctgtgtctct tggatgccct gtggcttcag tccgtctcct 180
gttgcccacc acctcgtccc tgggccgcct gataccccag cccaacagct aaggtgtgga 240
tggacagtag ggggctggct tctctcactg gtcaggggtc ttctcccctg tctgcctccc 300
ggagctagga ctgcagaggg gcctatcatg gtgcttgcag gccccctggc tgtctcgctg 360
ttgctgccca gcctcacact gctggtgtcc cacctctcca gctcccagga tgtctccagt 420
gagcccagca gtgagcagca gctgtgcgcc cttagcaagc accccaccgt ggcctttgaa 480
gacctgcagc cgtgggtctc taacttcacc taccctggag cccgggattt ctcccagctg 540
gctttggacc cctccgggaa ccagctcatc gtgggagcca ggaactacct cttcagactc 600
agccttgcca atgtctctct tcttcaggcc acagagtggg cctccagtga ggacacgcgc 660
cgctcctgcc aaagcaaagg gaagactgag gaggagtgtc agaactacgt gcgagtcctg 720
atcgtcgccg gccggaaggt gttcatgtgt ggaaccaatg ccttttcccc catgtgcacc 780
agcagacagg tggggaacct cagccggact actgagaaga tcaatggtgt ggcccgctgc 840
ccctatgacc cacgccacaa ctccacagct gtcatctcct cccaggggga gctctatgca 900
gccacggtca tcgacttctc aggtcgggac cctgccatct accgcagcct gggcagtggg 960
ccaccgcttc gcactgccca atataactcc aagtggctta atgagccaaa cttcgtggca 1020
gcctatgata ttgggctgtt tgcatacttc ttcctgcggg agaacgcagt ggagcacgac 1080
tgtggacgca ccgtgtactc tcgcgtggcc cgcgtgtgca agaatgacgt gggggggccga 1140
ttcctgctgg aggacacatg gaccacattc atgaaggccc ggctcaactg ctcccgcccg 1200
ggcgaggtcc ccttctacta taacgagctg cagagtgcct tccacttgcc agagcaggac 1260
ctcatctatg gagttttcac aaccaacgta aacagcatcg cggcttctgc tgtctgcgcc 1320
ttcaacctca gtgctatctc ccaggctttc aatggcccat ttcgctacca ggagaacccc 1380
agggctgcct ggctccccat agccaacccc atccccaatt tccagtgtgg caccctgcct 1440

```
gagaccggtc ccaacgagaa cctgacggag cgcagcctgc aggacgcgca gcgcctcttc   1500
ctgatgagcg aggccgtgca gccggtgaca cccgagccct gtgtcaccca ggacagcgtg   1560
cgcttctcac acctcgtggt ggacctggtg caggctaaag acacgctcta ccatgtactc   1620
tacattggca ccgagtcggg caccatcctg aaggcgctgt ccacggcgag ccgcagcctc   1680
cacggctgct acctggagga gctgcacgtg ctgccccccg ggcgccgcga gcccctgcgc   1740
agcctgcgca tcctgcacag cgcccgcgcg ctcttcgtgg ggctgagaga cggcgtcctg   1800
cgggtcccac tggagaggtg cgccgcctac cgcagccagg gggcatgcct gggggcccgg   1860
gacccgtact gtggctggga cgggaagcag caacgttgca gcacactcga ggacagctcc   1920
aacatgagcc tctggaccca gaacatcacc gcctgtcctg tgcggaatgt gacacgggat   1980
gggggcttcg gcccatggtc accatggcaa ccatgtgagc acttggatgg ggacaactca   2040
ggctcttgcc tgtgtcgagc tcgatcctgt gattcccctc gaccccgctg tgggggcctt   2100
gactgcctgg ggccagccat ccacatcgcc aactgctcca ggaatggggc gtggaccccg   2160
tggtcatcgt gggcgctgtg cagcacgtcc tgtggcatcg gcttccaggt ccgccagcga   2220
agttgcagca accctgctcc ccgccacggg ggccgcatct gcgtgggcaa gagccgggag   2280
gaacggttct gtaatgagaa cacgccttgc ccggtgccca tcttctgggc ttcctggggc   2340
tcctggagca agtgcagcag caactgtgga gggggcatgc agtcgcggcg tcgggcctgc   2400
gagaacggca actcctgcct gggctgcggc gtggagttca agacgtgcaa ccccgagggc   2460
tgccccgaag tgcggcgcaa cacccccctgg acgccgtggc tgcccgtgaa cgtgacgcag   2520
ggcggggcac ggcaggagca gcggttccgc ttcacctgcc gcgcgccccct tgcagacccg   2580
cacggcctgc agttcggcag gagaaggacc gagacgagga cctgtcccgc ggacggctcc   2640
ggctcctgcg acaccgacgc cctggtggag gtcctcctgc gcagcgggag cacctccccg   2700
cacacggtga gcgggggctg ggccgcctgg ggcccgtggt cgtcctgctc ccgggactgc   2760
gagctgggct tccgcgtccg caagagaacg tgcactaacc cggagccccg caacgggggc   2820
ctgccctgcg tgggcgatgc tgccgagtac caggactgca acccccaggc ttgcccagtt   2880
cggggtgctt ggtcctgctg gacctcatgg tctccatgct cagcttcctg tggtgggggt   2940
cactatcaac gcacccgttc ctgcaccagc cccgcaccct ccccaggtga ggacatctgt   3000
ctcgggctgc acacggagga ggcactatgt gccacacagg cctgcccaga aggctggtcg   3060
ccctggtctg agtggagtaa gtgcactgac gacggagccc agagccgaag ccggcactgt   3120
gaggagctcc tcccagggtc cagcgcmtgt gctggaaaca gcagccagag ccgcccctgc   3180
ccctacagcg agattcscgt catcctgcca gcctccagca tggaggaggc caccgactgt   3240
gcagggttca atctcatcca cttggtggcc acgggcatct cctgcttctt gggctctggg   3300
ctcctgaccc tagcagtgta cctgtcttgc cagcactgcc agcgtcagtc ccaggagtcc   3360
acactggtcc atcctgccac ccccaaccat ttgcactaca agggcggagg caccccgaag   3420
aatgaaaagt acacacccat ggaattcaag accctgaaca agaataactt gatccctgat   3480
gacagagcca acttctaccc attgcagcag accaatgcca gcgcaggtta tccacctctg   3540
cctggaagtc tatactctac ccagggcatc cctctggtca gaggcagtga gtactgggaa   3600
ctggaggctg acctgtgctt agaagtcctt taatctgggc tggtacaggc ctcagccttg   3660
ccctcaatgc acgaaaggtg gcccaggaga gaggatcaat gccataggag gcagaagtct   3720
ggcctctgtg cctctatgga gactatcttc cagttgctgc tcaacagagt tgttggctga   3780
gacctgcttg ggagtctctg ctggcccttc atctgttcag gaacacacac acacacacac   3840
tcacacacgc acacacaatc acaatttgct acagcaacaa aaaagacatt gggctgtggc   3900
attattaatt aaagatgata tccagtcaaa aaaaaact                           3938
```

<210> 34
<211> 1095
<212> PRT
<213> Homo sapiens

<400> 34

```
Met Val Leu Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu
1               5                  10                  15

Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu
            20                  25                  30

Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val
        35                  40                  45

Ala Phe Glu Asp Leu Gln Pro Trp Val Ser Asn Phe Thr Tyr Pro Gly
    50                  55                  60
```

```
Ala Arg Asp Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu
65                  70                  75                  80

Ile Val Gly Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val
                85                  90                  95

Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg
            100                 105                 110

Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val
        115                 120                 125

Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn
    130                 135                 140

Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg
145                 150                 155                 160

Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg
                165                 170                 175

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

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

Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu
    210                 215                 220

Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr
225                 230                 235                 240

Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val
                245                 250                 255

Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe
            260                 265                 270

Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys
        275                 280                 285

Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala
    290                 295                 300

Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn
305                 310                 315                 320

Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala
                325                 330                 335

Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg
            340                 345                 350

Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly
        355                 360                 365

Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu
    370                 375                 380
```

```
Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val
385                 390                 395                 400

Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu
                405                 410                 415

Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr
            420                 425                 430

Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser
        435                 440                 445

Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro
    450                 455                 460

Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg
465                 470                 475                 480

Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu
                485                 490                 495

Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp
            500                 505                 510

Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu
        515                 520                 525

Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro
    530                 535                 540

Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp
545                 550                 555                 560

Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys
                565                 570                 575

Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp
            580                 585                 590

Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Gly
        595                 600                 605

Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro Val
    610                 615                 620

Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro
625                 630                 635                 640

Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe
                645                 650                 655

Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp
            660                 665                 670

Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser
        675                 680                 685

Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val
    690                 695                 700

Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn
```

705 710 715 720

Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala
725 730 735

Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp
740 745 750

Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys
755 760 765

Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val
770 775 780

Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp
785 790 795 800

Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly
805 810 815

Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly
820 825 830

Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro
835 840 845

Gln Ala Cys Pro Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser
850 855 860

Pro Cys Ser Ala Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser
865 870 875 880

Cys Thr Ser Pro Ala Pro Ser Pro Gly Glu Asp Ile Cys Leu Gly Leu
885 890 895

His Thr Glu Glu Ala Leu Cys Ala Thr Gln Ala Cys Pro Glu Gly Trp
900 905 910

Ser Pro Trp Ser Glu Trp Ser Lys Cys Thr Asp Asp Gly Ala Gln Ser
915 920 925

Arg Ser Arg His Cys Glu Glu Leu Leu Pro Gly Ser Ser Ala Cys Ala
930 935 940

Gly Asn Ser Ser Gln Ser Arg Pro Cys Pro Tyr Ser Glu Ile Arg Val
945 950 955 960

Ile Leu Pro Ala Ser Ser Met Glu Glu Ala Thr Asp Cys Ala Gly Phe
965 970 975

Asn Leu Ile His Leu Val Ala Thr Gly Ile Ser Cys Phe Leu Gly Ser
980 985 990

Gly Leu Leu Thr Leu Ala Val Tyr Leu Ser Cys Gln His Cys Gln Arg
995 1000 1005

Gln Ser Gln Glu Ser Thr Leu Val His Pro Ala Thr Pro Asn His
1010 1015 1020

Leu His Tyr Lys Gly Gly Gly Thr Pro Lys Asn Glu Lys Tyr Thr
1025 1030 1035

```
Pro Met  Glu Phe Lys Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp
    1040                1045                1050

Asp Arg  Ala Asn Phe Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr
    1055                1060                1065

Thr Thr  Tyr Tyr Pro Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro
    1070                1075                1080

Glu Ala  Ser Pro Gly Gln Arg  Cys Phe Pro Asn Ser
    1085                1090                1095


<210>  35
<211>  1248
<212>  PRT
<213>  Homo sapiens

<400>  35
Arg Pro Pro His Ser Gln Thr Gly Arg Gln Pro Ile Trp Leu Ala Pro
1               5                  10                  15

Ala Ala Pro Arg Arg Pro Gly Val Gly Ser Arg Gly Glu Pro Gly Thr
            20                  25                  30

Cys Thr Arg Leu Trp Glu Pro Ala Trp Val Arg Val Ala Leu Gly Pro
        35                  40                  45

Ala Arg Ala Val Val Gly Ala Ser Gly Leu Gln Arg Arg Trp Gly Pro
    50                  55                  60

Gly Thr Gln Ala Trp Arg Arg Arg Arg Thr Ser Glu Ala Glu Gly Arg
65                  70                  75                  80

Arg Asp Arg Val Ser Gly Ser Ser Trp Cys Leu Ala Cys Val Ser Trp
                85                  90                  95

Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His Leu Val Pro
            100                 105                 110

Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly Trp Thr Val
        115                 120                 125

Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro Cys Leu Pro
    130                 135                 140

Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro
145                 150                 155                 160

Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His
                165                 170                 175

Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln
            180                 185                 190

Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln
        195                 200                 205

Pro Trp Val Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln
    210                 215                 220
```

```
Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn
225                 230                 235                 240

Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr
                245                 250                 255

Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly
            260                 265                 270

Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala
        275                 280                 285

Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys
    290                 295                 300

Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu Lys Ile Asn
305                 310                 315                 320

Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val
                325                 330                 335

Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser
            340                 345                 350

Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu
        355                 360                 365

Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val
    370                 375                 380

Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn
385                 390                 395                 400

Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg
                405                 410                 415

Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp
            420                 425                 430

Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val
        435                 440                 445

Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln
    450                 455                 460

Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala
465                 470                 475                 480

Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn
                485                 490                 495

Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile
            500                 505                 510

Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly
        515                 520                 525

Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu
    530                 535                 540
```

```
Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val
545                 550                 555                 560

Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln
                565                 570                 575

Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly
            580                 585                 590

Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys
        595                 600                 605

Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu
    610                 615                 620

Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu
625                 630                 635                 640

Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg
                645                 650                 655

Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp
            660                 665                 670

Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser
        675                 680                 685

Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg
    690                 695                 700

Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu
705                 710                 715                 720

Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp
                725                 730                 735

Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile
            740                 745                 750

His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val Ile
        755                 760                 765

Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe Gln Val Arg
    770                 775                 780

Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly Arg Ile Cys
785                 790                 795                 800

Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys
                805                 810                 815

Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser
            820                 825                 830

Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn
        835                 840                 845

Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro
    850                 855                 860

Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu
```

865 870 875 880

Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg
885 890 895

Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly
900 905 910

Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser
915 920 925

Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr
930 935 940

Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser
945 950 955 960

Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr
965 970 975

Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp
980 985 990

Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro Val Arg Gly
995 1000 1005

Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala Ser Cys
1010 1015 1020

Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro Ala
1025 1030 1035

Pro Ser Pro Gly Glu Asp Ile Cys Leu Gly Leu His Thr Glu Glu
1040 1045 1050

Ala Leu Cys Ala Thr Gln Ala Cys Pro Glu Gly Trp Ser Pro Trp
1055 1060 1065

Ser Glu Trp Ser Lys Cys Thr Asp Asp Gly Ala Gln Ser Arg Ser
1070 1075 1080

Arg His Cys Glu Glu Leu Leu Pro Gly Ser Ser Ala Cys Ala Gly
1085 1090 1095

Asn Ser Ser Gln Ser Arg Pro Cys Pro Tyr Ser Glu Ile Arg Val
1100 1105 1110

Ile Leu Pro Ala Ser Ser Met Glu Glu Ala Thr Asp Cys Ala Gly
1115 1120 1125

Phe Asn Leu Ile His Leu Val Ala Thr Gly Ile Ser Cys Phe Leu
1130 1135 1140

Gly Ser Gly Leu Leu Thr Leu Ala Val Tyr Leu Ser Cys Gln His
1145 1150 1155

Cys Gln Arg Gln Ser Gln Glu Ser Thr Leu Val His Pro Ala Thr
1160 1165 1170

Pro Asn His Leu His Tyr Lys Gly Gly Gly Thr Pro Lys Asn Glu
1175 1180 1185

```
Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1190                1195                1200

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1205                1210                1215

Val Tyr  Thr Thr Thr Tyr Tyr  Pro Ser Pro Leu Asn  Lys His Ser
    1220                1225                1230

Phe Arg  Pro Glu Ala Ser Pro  Gly Gln Arg Cys Phe  Pro Asn Ser
    1235                1240                1245
```

<210> 36
<211> 1150
<212> PRT
<213> Homo sapiens

<400> 36

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala
        35                  40                  45

Arg Thr Ala Glu Gly Pro Ile Met Val Leu Ala Gly Pro Leu Ala Val
    50                  55                  60

Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu Val Ser His Leu Ser Ser
65                  70                  75                  80

Ser Gln Asp Val Ser Ser Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala
                85                  90                  95

Leu Ser Lys His Pro Thr Val Ala Phe Glu Asp Leu Gln Pro Trp Val
            100                 105                 110

Ser Asn Phe Thr Tyr Pro Gly Ala Arg Asp Phe Ser Gln Leu Ala Leu
        115                 120                 125

Asp Pro Ser Gly Asn Gln Leu Ile Val Gly Ala Arg Asn Tyr Leu Phe
    130                 135                 140

Arg Leu Ser Leu Ala Asn Val Ser Leu Leu Gln Ala Thr Glu Trp Ala
145                 150                 155                 160

Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu
                165                 170                 175

Glu Glu Cys Gln Asn Tyr Val Arg Val Leu Ile Val Ala Gly Arg Lys
            180                 185                 190

Val Phe Met Cys Gly Thr Asn Ala Phe Ser Pro Met Cys Thr Ser Arg
        195                 200                 205

Gln Val Gly Asn Leu Ser Arg Thr Thr Glu Lys Ile Asn Gly Val Ala
```

```
    210                 215                 220

Arg Cys Pro Tyr Asp Pro Arg His Asn Ser Thr Ala Val Ile Ser Ser
225                 230                 235                 240

Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile Asp Phe Ser Gly Arg Asp
                245                 250                 255

Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala
            260                 265                 270

Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr
        275                 280                 285

Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu
    290                 295                 300

His Asp Cys Gly Arg Thr Val Tyr Ser Arg Val Ala Arg Val Cys Lys
305                 310                 315                 320

Asn Asp Val Gly Gly Arg Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe
                325                 330                 335

Met Lys Ala Arg Leu Asn Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr
            340                 345                 350

Tyr Asn Glu Leu Gln Ser Ala Phe His Leu Pro Glu Gln Asp Leu Ile
        355                 360                 365

Tyr Gly Val Phe Thr Thr Asn Val Asn Ser Ile Ala Ala Ser Ala Val
    370                 375                 380

Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe
385                 390                 395                 400

Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro
                405                 410                 415

Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu
            420                 425                 430

Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala Gln Arg Leu Phe Leu Met
        435                 440                 445

Ser Glu Ala Val Gln Pro Val Thr Pro Glu Pro Cys Val Thr Gln Asp
    450                 455                 460

Ser Val Arg Phe Ser His Leu Val Val Asp Leu Val Gln Ala Lys Asp
465                 470                 475                 480

Thr Leu Tyr His Val Leu Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu
                485                 490                 495

Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu His Gly Cys Tyr Leu Glu
            500                 505                 510

Glu Leu His Val Leu Pro Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu
        515                 520                 525

Arg Ile Leu His Ser Ala Arg Ala Leu Phe Val Gly Leu Arg Asp Gly
    530                 535                 540
```

```
Val Leu Arg Val Pro Leu Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly
545                 550                 555                 560

Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln
                565                 570                 575

Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser Asn Met Ser Leu Trp Thr
            580                 585                 590

Gln Asn Ile Thr Ala Cys Pro Val Arg Asn Val Thr Arg Asp Gly Gly
        595                 600                 605

Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys Glu His Leu Asp Gly Asp
    610                 615                 620

Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg
625                 630                 635                 640

Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly Pro Ala Ile His Ile Ala
                645                 650                 655

Asn Cys Ser Arg Asn Gly Ala Val Asp Pro Val Val His Arg Gly Pro
            660                 665                 670

Leu Cys Ser His Val Leu Trp His Ala Ala Ser Arg Ser Ala Ser Glu
        675                 680                 685

Val Ala Ala Thr Leu Leu Pro Ala Thr Gly Ala Ala Ser Ala Trp Ala
    690                 695                 700

Arg Ala Trp Glu Glu Arg Phe Cys Asn Glu Asn Thr Pro Cys Pro Val
705                 710                 715                 720

Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn
                725                 730                 735

Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn
            740                 745                 750

Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr Cys Asn Pro Glu Gly
        755                 760                 765

Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr Pro Trp Leu Pro Val
    770                 775                 780

Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr
785                 790                 795                 800

Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu Gln Phe Gly Arg Arg
                805                 810                 815

Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp
            820                 825                 830

Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser Gly Ser Thr Ser Pro
        835                 840                 845

His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys
    850                 855                 860
```

```
Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg Lys Arg Thr Cys Thr
865                 870                 875                 880

Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys Val Gly Asp Ala Ala
                885                 890                 895

Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro Val Arg Gly Ala Trp
            900                 905                 910

Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala Ser Cys Gly Gly Gly
        915                 920                 925

His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro Ala Pro Ser Pro Gly
    930                 935                 940

Glu Asp Ile Cys Leu Gly Leu His Thr Glu Glu Ala Leu Cys Ala Thr
945                 950                 955                 960

Gln Ala Cys Pro Glu Gly Trp Ser Pro Trp Ser Glu Trp Ser Lys Cys
                965                 970                 975

Thr Asp Asp Gly Ala Gln Ser Arg Ser Arg His Cys Glu Glu Leu Leu
            980                 985                 990

Pro Gly Ser Ser Ala Cys Ala Gly  Asn Ser Ser Gln Ser  Arg Pro Cys
        995                 1000                1005

Pro Tyr  Ser Glu Ile Arg Val  Ile Leu Pro Ala Ser  Ser Met Glu
    1010                1015                1020

Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1025                1030                1035

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1040                1045                1050

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1055                1060                1065

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1070                1075                1080

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
    1085                1090                1095

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1100                1105                1110

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1115                1120                1125

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1130                1135                1140

Gln Arg  Cys Phe Pro Asn Ser
    1145                1150


<210>  37
<211>  1211
<212>  PRT
```

<213> Homo sapiens

<400> 37

```
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140

Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

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

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
    290                 295                 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
```

305 310 315 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
325 330 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
340 345 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
355 360 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
370 375 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385 390 395 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
405 410 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
420 425 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
435 440 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
450 455 460

Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465 470 475 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
485 490 495

Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
500 505 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
515 520 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
530 535 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545 550 555 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
565 570 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
580 585 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
595 600 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
610 615 620

Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625 630 635 640

```
Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
                645                 650                 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
            660                 665                 670

Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
        675                 680                 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
    690                 695                 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705                 710                 715                 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
                725                 730                 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
            740                 745                 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
        755                 760                 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
    770                 775                 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
785                 790                 795                 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
                805                 810                 815

Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
            820                 825                 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
        835                 840                 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
    850                 855                 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865                 870                 875                 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
                885                 890                 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
            900                 905                 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
        915                 920                 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
    930                 935                 940

Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945                 950                 955                 960
```

```
Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
                965                 970                 975

Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
            980                 985                 990

Ala Pro Ser Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
        995                 1000                1005

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1010                1015                1020

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1025                1030                1035

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1040                1045                1050

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1055                1060                1065

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1070                1075                1080

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1085                1090                1095

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1100                1105                1110

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                1120                1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1130                1135                1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                1150                1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                1165                1170

Ala Ser  Ala Gly Tyr Pro Pro  Leu Pro Gly Ser Leu  Tyr Ser Thr
    1175                1180                1185

Gln Gly  Ile Pro Leu Val Arg  Gly Ser Glu Tyr Trp  Glu Leu Glu
    1190                1195                1200

Ala Asp  Leu Cys Leu Glu Val  Leu
    1205                1210


<210>  38
<211>  1203
<212>  PRT
<213>  Homo sapiens

<400>  38
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                   10                  15
```

```
Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60

Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Leu Leu Pro
                85                  90                  95

Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro Ile Met Val Leu
            100                 105                 110

Ala Gly Pro Leu Ala Val Ser Leu Leu Leu Pro Ser Leu Thr Leu Leu
        115                 120                 125

Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser Glu Pro Ser Ser
    130                 135                 140

Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr Val Ala Phe Glu
145                 150                 155                 160

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

Phe Ser Gln Leu Ala Leu Asp Pro Ser Gly Asn Gln Leu Ile Val Gly
            180                 185                 190

Ala Arg Asn Tyr Leu Phe Arg Leu Ser Leu Ala Asn Val Ser Leu Leu
        195                 200                 205

Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg Arg Ser Cys Gln
    210                 215                 220

Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr Val Arg Val Leu
225                 230                 235                 240

Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr Asn Ala Phe Ser
                245                 250                 255

Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser Arg Thr Thr Glu
            260                 265                 270

Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro Arg His Asn Ser
        275                 280                 285

Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala Ala Thr Val Ile
    290                 295                 300

Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser Leu Gly Ser Gly
305                 310                 315                 320

Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp Leu Asn Glu Pro
                325                 330                 335

Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala Tyr Phe Phe Leu
```

```
            340                 345                 350

Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr Val Tyr Ser Arg
        355                 360                 365

Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg Phe Leu Leu Glu
    370                 375                 380

Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn Cys Ser Arg Pro
385                 390                 395                 400

Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser Ala Phe His Leu
                405                 410                 415

Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr Asn Val Asn Ser
            420                 425                 430

Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser Ala Ile Ser Gln
        435                 440                 445

Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro Arg Ala Ala Trp
    450                 455                 460

Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys Gly Thr Leu Pro
465                 470                 475                 480

Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser Leu Gln Asp Ala
                485                 490                 495

Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro Val Thr Pro Glu
            500                 505                 510

Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His Leu Val Val Asp
        515                 520                 525

Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu Tyr Ile Gly Thr
    530                 535                 540

Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala Ser Arg Ser Leu
545                 550                 555                 560

His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro Pro Gly Arg Arg
                565                 570                 575

Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala Arg Ala Leu Phe
            580                 585                 590

Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu Glu Arg Cys Ala
        595                 600                 605

Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg Asp Pro Tyr Cys
    610                 615                 620

Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu Glu Asp Ser Ser
625                 630                 635                 640

Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys Pro Val Arg Asn
                645                 650                 655

Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro Trp Gln Pro Cys
            660                 665                 670
```

```
Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu Cys Arg Ala Arg
        675                 680                 685

Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu Asp Cys Leu Gly
    690                 695                 700

Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly Ala Val Asp Pro
705                 710                 715                 720

Val Val Ile Val Gly Arg Cys Ala Ala Thr Ser Cys Gly Ile Gly Phe
                725                 730                 735

Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala Pro Arg His Gly Gly
            740                 745                 750

Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg Phe Cys Asn Glu Asn
        755                 760                 765

Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser Trp Gly Ser Trp Ser
    770                 775                 780

Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln Ser Arg Arg Arg Ala
785                 790                 795                 800

Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly Val Glu Phe Lys Thr
                805                 810                 815

Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg Asn Thr Pro Trp Thr
            820                 825                 830

Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly Ala Arg Gln Glu Gln
        835                 840                 845

Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala Asp Pro His Gly Leu
    850                 855                 860

Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr Cys Pro Ala Asp Gly
865                 870                 875                 880

Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu Val Leu Leu Arg Ser
                885                 890                 895

Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly Trp Ala Ala Trp Gly
            900                 905                 910

Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu Gly Phe Arg Val Arg
        915                 920                 925

Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn Gly Gly Leu Pro Cys
    930                 935                 940

Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn Pro Gln Ala Cys Pro
945                 950                 955                 960

Val Arg Gly Ala Trp Ser Cys Trp Thr Ser Trp Ser Pro Cys Ser Ala
                965                 970                 975

Ser Cys Gly Gly Gly His Tyr Gln Arg Thr Arg Ser Cys Thr Ser Pro
            980                 985                 990
```

```
Ala Pro Ser Pro Gly Glu Asp Ile  Cys Leu Gly Leu His  Thr Glu Glu
        995                 1000                1005

Ala Leu  Cys Ala Thr Gln Ala  Cys Pro Glu Gly Trp  Ser Pro Trp
    1010                1015                1020

Ser Glu  Trp Ser Lys Cys Thr  Asp Asp Gly Ala Gln  Ser Arg Ser
    1025                1030                1035

Arg His  Cys Glu Glu Leu Leu  Pro Gly Ser Ser Ala  Cys Ala Gly
    1040                1045                1050

Asn Ser  Ser Gln Ser Arg Pro  Cys Pro Tyr Ser Glu  Ile Arg Val
    1055                1060                1065

Ile Leu  Pro Ala Ser Ser Met  Glu Glu Ala Thr Asp  Cys Ala Gly
    1070                1075                1080

Phe Asn  Leu Ile His Leu Val  Ala Thr Gly Ile Ser  Cys Phe Leu
    1085                1090                1095

Gly Ser  Gly Leu Leu Thr Leu  Ala Val Tyr Leu Ser  Cys Gln His
    1100                1105                1110

Cys Gln  Arg Gln Ser Gln Glu  Ser Thr Leu Val His  Pro Ala Thr
    1115                1120                1125

Pro Asn  His Leu His Tyr Lys  Gly Gly Gly Thr Pro  Lys Asn Glu
    1130                1135                1140

Lys Tyr  Thr Pro Met Glu Phe  Lys Thr Leu Asn Lys  Asn Asn Leu
    1145                1150                1155

Ile Pro  Asp Asp Arg Ala Asn  Phe Tyr Pro Leu Gln  Gln Thr Asn
    1160                1165                1170

Val Tyr  Thr Thr Thr Tyr Tyr  Pro Ser Pro Leu Asn  Lys His Ser
    1175                1180                1185

Phe Arg  Pro Glu Ala Ser Pro  Gly Gln Arg Cys Phe  Pro Asn Ser
    1190                1195                1200


<210>  39
<211>  1240
<212>  PRT
<213>  Homo sapiens

<400>  39
Ala Ala Ala Pro Phe Pro Asp Arg Pro Pro Ala His Leu Val Ser Ser
1               5                  10                  15

Arg Arg Ser Ala Pro Pro Gly Ser Arg Glu Pro Arg Gly Thr Gly His
            20                  25                  30

Leu His Pro Pro Leu Gly Val Ser Gly Ser Ser Trp Cys Leu Ala Cys
        35                  40                  45

Val Ser Trp Met Pro Cys Gly Phe Ser Pro Ser Pro Val Ala His His
    50                  55                  60
```

```
Leu Val Pro Gly Pro Pro Asp Thr Pro Ala Gln Gln Leu Arg Cys Gly
65                  70                  75                  80

Trp Thr Val Gly Gly Trp Leu Leu Ser Leu Val Arg Gly Arg Lys Pro
                85                  90                  95

Ser Gly Asp Phe Glu Trp Arg Gln Gly Trp Arg Gly Pro Gly Glu Glu
            100                 105                 110

Asp Trp Pro Glu Ser Pro Ser Pro Lys Val Leu Met Asp Ser Ala Gly
        115                 120                 125

Gly Leu Leu Pro Cys Leu Pro Pro Gly Ala Arg Thr Ala Glu Gly Pro
    130                 135                 140

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

Leu Thr Leu Leu Val Ser His Leu Ser Ser Ser Gln Asp Val Ser Ser
                165                 170                 175

Glu Pro Ser Ser Glu Gln Gln Leu Cys Ala Leu Ser Lys His Pro Thr
            180                 185                 190

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

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

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

Val Ser Leu Leu Gln Ala Thr Glu Trp Ala Ser Ser Glu Asp Thr Arg
                245                 250                 255

Arg Ser Cys Gln Ser Lys Gly Lys Thr Glu Glu Glu Cys Gln Asn Tyr
            260                 265                 270

Val Arg Val Leu Ile Val Ala Gly Arg Lys Val Phe Met Cys Gly Thr
        275                 280                 285

Asn Ala Phe Ser Pro Met Cys Thr Ser Arg Gln Val Gly Asn Leu Ser
    290                 295                 300

Arg Thr Thr Glu Lys Ile Asn Gly Val Ala Arg Cys Pro Tyr Asp Pro
305                 310                 315                 320

Arg His Asn Ser Thr Ala Val Ile Ser Ser Gln Gly Glu Leu Tyr Ala
                325                 330                 335

Ala Thr Val Ile Asp Phe Ser Gly Arg Asp Pro Ala Ile Tyr Arg Ser
            340                 345                 350

Leu Gly Ser Gly Pro Pro Leu Arg Thr Ala Gln Tyr Asn Ser Lys Trp
        355                 360                 365

Leu Asn Glu Pro Asn Phe Val Ala Ala Tyr Asp Ile Gly Leu Phe Ala
    370                 375                 380

Tyr Phe Phe Leu Arg Glu Asn Ala Val Glu His Asp Cys Gly Arg Thr
```

385 390 395 400

Val Tyr Ser Arg Val Ala Arg Val Cys Lys Asn Asp Val Gly Gly Arg
405 410 415

Phe Leu Leu Glu Asp Thr Trp Thr Thr Phe Met Lys Ala Arg Leu Asn
420 425 430

Cys Ser Arg Pro Gly Glu Val Pro Phe Tyr Tyr Asn Glu Leu Gln Ser
435 440 445

Ala Phe His Leu Pro Glu Gln Asp Leu Ile Tyr Gly Val Phe Thr Thr
450 455 460

Asn Val Asn Ser Ile Ala Ala Ser Ala Val Cys Ala Phe Asn Leu Ser
465 470 475 480

Ala Ile Ser Gln Ala Phe Asn Gly Pro Phe Arg Tyr Gln Glu Asn Pro
485 490 495

Arg Ala Ala Trp Leu Pro Ile Ala Asn Pro Ile Pro Asn Phe Gln Cys
500 505 510

Gly Thr Leu Pro Glu Thr Gly Pro Asn Glu Asn Leu Thr Glu Arg Ser
515 520 525

Leu Gln Asp Ala Gln Arg Leu Phe Leu Met Ser Glu Ala Val Gln Pro
530 535 540

Val Thr Pro Glu Pro Cys Val Thr Gln Asp Ser Val Arg Phe Ser His
545 550 555 560

Leu Val Val Asp Leu Val Gln Ala Lys Asp Thr Leu Tyr His Val Leu
565 570 575

Tyr Ile Gly Thr Glu Ser Gly Thr Ile Leu Lys Ala Leu Ser Thr Ala
580 585 590

Ser Arg Ser Leu His Gly Cys Tyr Leu Glu Glu Leu His Val Leu Pro
595 600 605

Pro Gly Arg Arg Glu Pro Leu Arg Ser Leu Arg Ile Leu His Ser Ala
610 615 620

Arg Ala Leu Phe Val Gly Leu Arg Asp Gly Val Leu Arg Val Pro Leu
625 630 635 640

Glu Arg Cys Ala Ala Tyr Arg Ser Gln Gly Ala Cys Leu Gly Ala Arg
645 650 655

Asp Pro Tyr Cys Gly Trp Asp Gly Lys Gln Gln Arg Cys Ser Thr Leu
660 665 670

Glu Asp Ser Ser Asn Met Ser Leu Trp Thr Gln Asn Ile Thr Ala Cys
675 680 685

Pro Val Arg Asn Val Thr Arg Asp Gly Gly Phe Gly Pro Trp Ser Pro
690 695 700

Trp Gln Pro Cys Glu His Leu Asp Gly Asp Asn Ser Gly Ser Cys Leu
705 710 715 720

```
Cys Arg Ala Arg Ser Cys Asp Ser Pro Arg Pro Arg Cys Gly Gly Leu
                725                 730                 735

Asp Cys Leu Gly Pro Ala Ile His Ile Ala Asn Cys Ser Arg Asn Gly
            740                 745                 750

Gly Arg Gly Pro Arg Gly Ala Ser Trp Ala Ala Val Gln Ala Arg Pro
        755                 760                 765

Val Ala Ser Gly Phe Gln Val Arg Gln Arg Ser Cys Ser Asn Pro Ala
    770                 775                 780

Pro Arg His Gly Gly Arg Ile Cys Val Gly Lys Ser Arg Glu Glu Arg
785                 790                 795                 800

Phe Cys Asn Glu Asn Thr Pro Cys Pro Val Pro Ile Phe Trp Ala Ser
                805                 810                 815

Trp Gly Ser Trp Ser Lys Cys Ser Ser Asn Cys Gly Gly Gly Met Gln
            820                 825                 830

Ser Arg Arg Arg Ala Cys Glu Asn Gly Asn Ser Cys Leu Gly Cys Gly
        835                 840                 845

Val Glu Phe Lys Thr Cys Asn Pro Glu Gly Cys Pro Glu Val Arg Arg
    850                 855                 860

Asn Thr Pro Trp Thr Pro Trp Leu Pro Val Asn Val Thr Gln Gly Gly
865                 870                 875                 880

Ala Arg Gln Glu Gln Arg Phe Arg Phe Thr Cys Arg Ala Pro Leu Ala
                885                 890                 895

Asp Pro His Gly Leu Gln Phe Gly Arg Arg Arg Thr Glu Thr Arg Thr
            900                 905                 910

Cys Pro Ala Asp Gly Ser Gly Ser Cys Asp Thr Asp Ala Leu Val Glu
        915                 920                 925

Val Leu Leu Arg Ser Gly Ser Thr Ser Pro His Thr Val Ser Gly Gly
    930                 935                 940

Trp Ala Ala Trp Gly Pro Trp Ser Ser Cys Ser Arg Asp Cys Glu Leu
945                 950                 955                 960

Gly Phe Arg Val Arg Lys Arg Thr Cys Thr Asn Pro Glu Pro Arg Asn
                965                 970                 975

Gly Gly Leu Pro Cys Val Gly Asp Ala Ala Glu Tyr Gln Asp Cys Asn
            980                 985                 990

Pro Gln Ala Cys Pro Val Arg Gly  Ala Trp Ser Cys Trp  Thr Ser Trp
        995                 1000                1005

Ser Pro  Cys Ser Ala Ser Cys  Gly Gly Gly His Tyr  Gln Arg Thr
    1010                1015                1020

Arg Ser  Cys Thr Ser Pro Ala  Pro Ser Pro Gly Glu  Asp Ile Cys
    1025                1030                1035
```

```
Leu Gly  Leu His Thr Glu Glu  Ala Leu Cys Ala Thr  Gln Ala Cys
    1040                1045                1050

Pro Glu  Gly Trp Ser Pro Trp  Ser Glu Trp Ser Lys  Cys Thr Asp
    1055                1060                1065

Asp Gly  Ala Gln Ser Arg Ser  Arg His Cys Glu Glu  Leu Leu Pro
    1070                1075                1080

Gly Ser  Ser Ala Cys Ala Gly  Asn Ser Ser Gln Ser  Arg Pro Cys
    1085                1090                1095

Pro Tyr  Ser Glu Ile Arg Val  Ile Leu Pro Ala Ser  Ser Met Glu
    1100                1105                1110

Glu Ala  Thr Asp Cys Ala Gly  Phe Asn Leu Ile His  Leu Val Ala
    1115                1120                1125

Thr Gly  Ile Ser Cys Phe Leu  Gly Ser Gly Leu Leu  Thr Leu Ala
    1130                1135                1140

Val Tyr  Leu Ser Cys Gln His  Cys Gln Arg Gln Ser  Gln Glu Ser
    1145                1150                1155

Thr Leu  Val His Pro Ala Thr  Pro Asn His Leu His  Tyr Lys Gly
    1160                1165                1170

Gly Gly  Thr Pro Lys Asn Glu  Lys Tyr Thr Pro Met  Glu Phe Lys
    1175                1180                1185

Thr Leu  Asn Lys Asn Asn Leu  Ile Pro Asp Asp Arg  Ala Asn Phe
    1190                1195                1200

Tyr Pro  Leu Gln Gln Thr Asn  Val Tyr Thr Thr Thr  Tyr Tyr Pro
    1205                1210                1215

Ser Pro  Leu Asn Lys His Ser  Phe Arg Pro Glu Ala  Ser Pro Gly
    1220                1225                1230

Gln Arg  Cys Phe Pro Asn Ser
    1235                1240
```