

(19) World Intellectual Property Organization  
International Bureau(43) International Publication Date  
7 January 2010 (07.01.2010)

PCT

(10) International Publication Number  
**WO 2010/003118 A1**

## (51) International Patent Classification:

C12N 15/62 (2006.01) C07K 16/24 (2006.01)  
C07K 14/715 (2006.01) A61K 38/17 (2006.01)  
C07K 16/28 (2006.01)

## (21) International Application Number:

PCT/US2009/049616

## (22) International Filing Date:

2 July 2009 (02.07.2009)

## (25) Filing Language:

English

## (26) Publication Language:

English

## (30) Priority Data:

61/134,091	2 July 2008 (02.07.2008)	US
61/134,090	2 July 2008 (02.07.2008)	US
61/134,089	2 July 2008 (02.07.2008)	US
61/134,088	2 July 2008 (02.07.2008)	US
61/134,085	2 July 2008 (02.07.2008)	US
61/134,087	2 July 2008 (02.07.2008)	US
61/134,086	2 July 2008 (02.07.2008)	US
61/181,232	26 May 2009 (26.05.2009)	US

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PE, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(81) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

## Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))

## Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))
- with sequence listing part of description (Rule 5.2(a))

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(54) Title: TGF-B ANTAGONIST MULTI-TARGET BINDING PROTEINS

(57) Abstract: This disclosure provides a multi-target fusion protein composed of a TGF $\beta$  antagonist domain and another binding domain antagonistic for a heterologous target (such as IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF) or agonistic for a heterologous target (such as GPCR). The multi-specific fusion protein may also include an intervening domain that separates the binding domains and allows for dimerization. This disclosure also provides polynucleotides encoding the multi-specific fusion proteins, compositions of the fusion proteins, and methods of using the multi-specific fusion proteins and compositions.

## **TGF- $\beta$ ANTAGONIST MULTI-TARGET BINDING PROTEINS**

### **TECHNICAL FIELD**

[001] This disclosure relates generally to the field of multi-target binding molecules and therapeutic applications thereof and more specifically to a fusion protein composed of either a transforming growth factor-beta (TGF $\beta$ ) antagonist domain and another binding domain antagonistic for a heterologous target, such as IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF1 or IGF2, or a TGF $\beta$  antagonist domain and another binding domain agonistic for a heterologous target, such as GPCR, as well as compositions and therapeutic uses thereof.

### **BACKGROUND**

[002] Transforming growth factor-beta (TGF $\beta$ ) is a potent cytokine that has significant effects on the immune system. The main function of TGF $\beta$  in the immune system is to maintain tolerance and initial immune responses against foreign pathogens. Three isoforms of TGF $\beta$  have been identified in mammals, TGF $\beta$ 1, TGF $\beta$ 2 and TGF $\beta$ 3, with TGF $\beta$ 1 being the predominant isoform. TGF $\beta$  is secreted in a latent form and only a small percentage of total secreted TGF $\beta$  is activated under physiological conditions. The biological effects of TGF $\beta$  occur mostly through binding of TGF $\beta$  to the receptors ALK5 and TGF $\beta$  receptor II (TGF $\beta$ R2). Specifically, active TGF $\beta$  dimer binds to a tetrameric ALK5 and TGF $\beta$ R2 complex to initiate cell signaling. ALK5 is not required for the initial binding of TGF $\beta$ , but is required for signaling.

[003] TGF $\beta$  has been shown to influence many cellular functions such as cell proliferation, differentiation, cell-cell and cell-matrix adhesion, cell motility and activation of lymphocytes. (For a review of the role of TGF $\beta$  in regulating immune responses, see Li *et al.* (2006) *Annu. Rev. Immunol.* 24:99-146.) Furthermore, TGF $\beta$  is believed to induce or mediate the progression of many diseases such as osteoporosis, hypertension, atherosclerosis, hepatic cirrhosis and fibrotic diseases of the kidney, liver and lungs, and tumor progression. TGF $\beta$  can augment end-organ damage caused by chronic inflammation and TGF $\beta$  antagonists have been shown to be effective in attenuating this damage in animal models of diseases such as diabetic kidney disease, glomerulonephritis, cyclosporine-mediated renal injury and systemic lupus erythematosus (SLE) (Border *et al.* (1990) *Nature* 346:371-374; Border *et al.* (1992) *Nature* 360:361-364; Isaka *et al.* (1999) *Kidney Int.* 55:465-475; Sharma *et al.* (1996)

Diabetes 45:522; Xin *et al.* (2004) Transplantation 15:1433; Benigni *et al.* (2003) J. Am. Soc. Nephrol. 14:1816). With respect to cancer, TGF $\beta$  can have a direct inhibitory activity on malignant cells and can augment the production or activity of a range of tumor growth factors and angiogenic factors.

[004] While TGF $\beta$  knock-out mice have severe pathology related to unrestrained inflammation and autoimmunity, administration of TGF $\beta$  antagonists is well tolerated in mice and humans (Rusek *et al.* (2003) Immunopharmacol. Immunotoxicol. 25:235-57; Denton *et al.* (2007) Arthritis Rheum. 56:323-33). Methods of treatment using TGF antagonists known in the art include use of antibodies against TGF $\beta$ , use of TGF $\beta$ R2 ectodomain Ig fusion proteins, and use of small molecule inhibitors of TGF $\beta$ RI kinase activity. All of these methods have modest beneficial impact in rodent models of disease or in clinical trials in humans (Denton *et al.* (2007) Arthritis Rheum. 56:323). Indeed, chronic use of a TGF $\beta$  antagonist in mice shows no evidence of activation of the immune system as might be expected from the phenotype of TGF $\beta$ -/- knock-out mice. This is likely to reflect, in part, the complex nature of the biology of cytokines, interleukins, chemokines and growth factors in human diseases and the requirement to inhibit more than one pathway simultaneously to maximize the benefit to patients.

#### **BRIEF DESCRIPTION OF THE FIGURES**

[005] Figures 1A-1C show that multi-specific (Xceptor) fusion proteins containing one of various different Hyper-IL6 binding domains fused to a TNFR ectodomain bind to Hyper-IL6 specifically as measured by ELISA, and that these multi-specific fusion proteins preferentially bind Hyper-IL6 over IL6 and IL6R alone. Only two fusion proteins tested bound IL6 and none bound sIL6R.

[006] Figure 2 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to one of various different Hyper-IL6 binding domains bind to TNF- $\alpha$  as measured by ELISA.

[007] Figure 3 shows that multi-specific fusion proteins containing one of various different Hyper-IL6 binding domains fused to a TNFR ectodomain can simultaneously bind to Hyper-IL6 and TNF- $\alpha$  as measured by ELISA.

[008] Figure 4 shows that multi-specific fusion proteins containing one of various different Hyper-IL6 binding domains fused to a TNFR ectodomain block gp130 from binding to Hyper-IL6 as measured by ELISA.

[0009] Figures 5A and 5B show that multi-specific fusion proteins containing one of various different Hyper-IL6 binding domains fused to a TNFR ectodomain block (A) IL6 or (B) Hyper-IL6 induced proliferation of TF-1 cells.

[0010] Figure 6 shows that multi-specific fusion proteins containing one of various different Hyper-IL6 binding domains fused to a TNFR ectodomain block TNF- $\alpha$  from binding to TNFR as measured by ELISA.

[0011] Figure 7 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to one of various different Hyper-IL6 binding domains block TNF- $\alpha$  induced killing of L929 cells.

[0012] Figure 8 shows that multi-specific fusion proteins containing a TGF $\beta$ R2 ectodomain fused to one of various different Hyper-IL6 binding domains bind to TGF $\beta$ 1 as measured by ELISA.

[0013] Figure 9 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to a TGF $\beta$ RII ectodomain block TGF $\beta$ -1 induced inhibition of IL-4 proliferation of HT2 cells.

[0014] Figure 10 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to an IL6 binding domain did not bind to HepG2 (liver) cells.

[0015] Figure 11 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to an IL6 binding domain blocked the HIL6-induced SAA response in mice.

[0016] Figure 12 shows that multi-specific fusion proteins containing a TNFR ectodomain fused to an IL6 binding domain blocked the HIL6-induced sgp130 response in mice.

[0017] Figures 13A and B show the results of studies on the ability of multi-specific fusion proteins containing a TNFR ectodomain fused to an IL6 binding domain to block the TNF $\alpha$ -induced SAA response in mice, at 2 hours and 24 hours post-administration, respectively.

#### **DETAILED DESCRIPTION**

[0018] The present disclosure provides multi-specific fusion proteins, referred to herein as Xceptor molecules. Exemplary structures of such multi-specific fusion proteins, include N-BD-ID-ED-C, N-ED-ID-BD-C, and N-ED1-ID-ED2-C, wherein N- and -C represent the amino- and carboxy-terminus, respectively, BD is an immunoglobulin-like or

immunoglobulin variable region binding domain, ID is an intervening domain, and ED is an ectodomain (*e.g.* an extracellular domain), such as a receptor ligand binding domain, cysteine rich domain (A domain; *see* WO 02/088171 and WO 04/044011), semaphorin or semaphorin-like domain, or the like. In some constructs, the ID can comprise an immunoglobulin constant region or sub-region disposed between the first and second binding domains. In still further constructs, the BD and ED are each linked to the ID via the same or different linker (*e.g.*, a linker comprising one to 50 amino acids), such as an immunoglobulin hinge region (made up of, for example, the upper and core regions) or functional variant thereof, or a lectin interdomain region or functional variant thereof, or a cluster of differentiation (CD) molecule stalk region or functional variant thereof.

[0019] Prior to setting forth this disclosure in more detail, it may be helpful to an understanding thereof to provide definitions of certain terms to be used herein. Additional definitions are set forth throughout this disclosure.

[0020] In the present description, any concentration range, percentage range, ratio range, or integer range is to be understood to include the value of any integer within the recited range and, when appropriate, fractions thereof (such as one tenth and one hundredth of an integer), unless otherwise indicated. Also, any number range recited herein relating to any physical feature, such as polymer subunits, size or thickness, are to be understood to include any integer within the recited range, unless otherwise indicated. As used herein, "about" or "consisting essentially of" mean  $\pm 20\%$  of the indicated range, value, or structure, unless otherwise indicated. It should be understood that the terms "a" and "an" as used herein refer to "one or more" of the enumerated components. The use of the alternative (*e.g.*, "or") should be understood to mean either one, both, or any combination thereof of the alternatives. As used herein, the terms "include" and "comprise" are used synonymously. In addition, it should be understood that the individual compounds, or groups of compounds, derived from the various combinations of the structures and substituents described herein, are disclosed by the present application to the same extent as if each compound or group of compounds was set forth individually. Thus, selection of particular structures or particular substituents is within the scope of the present disclosure.

[0021] A "binding domain" or "binding region" according to the present disclosure may be, for example, any protein, polypeptide, oligopeptide, or peptide that possesses the ability to specifically recognize and bind to a biological molecule (*e.g.*, TGF $\beta$  or IL6) or complex of more than one of the same or different molecule or assembly or aggregate,

whether stable or transient (*e.g.*, IL6/IL6R complex). Such biological molecules include proteins, polypeptides, oligopeptides, peptides, amino acids, or derivatives thereof; lipids, fatty acids, or derivatives thereof; carbohydrates, saccharides, or derivatives thereof; nucleotides, nucleosides, peptide nucleic acids, nucleic acid molecules, or derivatives thereof; glycoproteins, glycopeptides, glycolipids, lipoproteins, proteolipids, or derivatives thereof; other biological molecules that may be present in, for example, a biological sample; or any combination thereof. A binding region includes any naturally occurring, synthetic, semi-synthetic, or recombinantly produced binding partner for a biological molecule or other target of interest. A variety of assays are known for identifying binding domains of the present disclosure that specifically bind with a particular target, including Western blot, ELISA, or Biacore analysis.

**[0022]** Binding domains and fusion proteins thereof of this disclosure can be capable of binding to a desired degree, including “specifically or selectively binding” a target while not significantly binding other components present in a test sample, if they bind a target molecule with an affinity or  $K_a$  (*i.e.*, an equilibrium association constant of a particular binding interaction with units of  $1/M$ ) of, for example, greater than or equal to about  $10^5 M^{-1}$ ,  $10^6 M^{-1}$ ,  $10^7 M^{-1}$ ,  $10^8 M^{-1}$ ,  $10^9 M^{-1}$ ,  $10^{10} M^{-1}$ ,  $10^{11} M^{-1}$ ,  $10^{12} M^{-1}$ , or  $10^{13} M^{-1}$ . “High affinity” binding domains refers to those binding domains with a  $K_a$  of at least  $10^7 M^{-1}$ , at least  $10^8 M^{-1}$ , at least  $10^9 M^{-1}$ , at least  $10^{10} M^{-1}$ , at least  $10^{11} M^{-1}$ , at least  $10^{12} M^{-1}$ , at least  $10^{13} M^{-1}$ , or greater. Alternatively, affinity may be defined as an equilibrium dissociation constant ( $K_d$ ) of a particular binding interaction with units of  $M$  (*e.g.*,  $10^{-5} M$  to  $10^{-13} M$ ). Affinities of binding domain polypeptides and fusion proteins according to the present disclosure can be readily determined using conventional techniques (*see, e.g.*, Scatchard *et al.* (1949) Ann. N.Y. Acad. Sci. 51:660; and U.S. Patent Nos. 5,283,173; 5,468,614; Biacore® analysis; or the equivalent).

**[0023]** Binding domains of this disclosure can be generated as described herein or by a variety of methods known in the art (*see, e.g.*, US Patent Nos. 6,291,161; 6,291,158). Sources include antibody gene sequences from various species (which can be formatted as antibodies, sFvs, scFvs or Fabs, such as in a phage library), including human, camelid (from camels, dromedaries, or llamas; Hamers-Casterman *et al.* (1993) Nature, 363:446 and Nguyen *et al.* (1998) J. Mol. Biol., 275:413), shark (Roux *et al.* (1998) Proc. Nat’l. Acad. Sci. (USA) 95:11804), fish (Nguyen *et al.* (2002) Immunogenetics, 54:39), rodent, avian, ovine, sequences that encode random peptide libraries or sequences that encode an engineered diversity of amino acids in loop regions of alternative non-antibody scaffolds, such as

fibrinogen domains (*see, e.g.*, Weisel *et al.* (1985) Science 230:1388), Kunitz domains (*see, e.g.*, US Patent No. 6,423,498), lipocalin domains (*see, e.g.*, WO 2006/095164), V-like domains (*see, e.g.*, US Patent Application Publication No. 2007/0065431), C-type lectin domains (Zelensky and Gready (2005) FEBS J. 272:6179), mAb<sup>2</sup> or Fcab<sup>TM</sup> (*see, e.g.*, PCT Patent Application Publication Nos. WO 2007/098934; WO 2006/072620), or the like. Additionally, traditional strategies for hybridoma development using a synthetic single chain IL6/IL6R complex, such as a human IL6/IL6R complex or Hyper-IL6 (IL6 joined by a peptide linker to IL6R), as an immunogen in convenient systems (*e.g.*, mice, HuMAb mouse®, TC mouse<sup>TM</sup>, KM-mouse®, llamas, chicken, rats, hamsters, rabbits, *etc.*) can be used to develop binding domains of this disclosure.

**[0024]** Terms understood by those in the art as referring to antibody technology are each given the meaning acquired in the art, unless expressly defined herein. For example, the terms “V<sub>L</sub>” and “V<sub>H</sub>” refer to the variable binding region derived from an antibody light and heavy chain, respectively. The variable binding regions are made up of discrete, well-defined sub-regions known as “complementarity determining regions” (CDRs) and “framework regions” (FRs). The terms “C<sub>L</sub>” and “C<sub>H</sub>” refer to an “immunoglobulin constant region,” *i.e.*, a constant region derived from an antibody light or heavy chain, respectively, with the latter region understood to be further divisible into C<sub>H1</sub>, C<sub>H2</sub>, C<sub>H3</sub> and C<sub>H4</sub> constant region domains, depending on the antibody isotype (IgA, IgD, IgE, IgG, IgM) from which the region was derived. A portion of the constant region domains makes up the Fc region (the “fragment crystallizable” region), which contains domains responsible for the effector functions of an immunoglobulin, such as ADCC (antibody-dependent cell-mediated cytotoxicity), ADCP (antibody-dependent cell-mediated phagocytosis), CDC (complement-dependent cytotoxicity) and complement fixation, binding to Fc receptors, greater half-life *in vivo* relative to a polypeptide lacking an Fc region, protein A binding, and perhaps even placental transfer (*see* Capon *et al.* (1989) Nature, 337:525). Further, a polypeptide containing an Fc region allows for dimerization or multimerization of the polypeptide. A “hinge region,” also referred to herein as a “linker,” is an amino acid sequence interposed between and connecting the variable binding and constant regions of a single chain of an antibody, which is known in the art as providing flexibility in the form of a hinge to antibodies or antibody-like molecules.

**[0025]** The domain structure of immunoglobulins is amenable to engineering, in that the antigen binding domains and the domains conferring effector functions may be exchanged between immunoglobulin classes and subclasses. Immunoglobulin structure and function are reviewed, for example, in Harlow *et al.*, Eds., *Antibodies: A Laboratory Manual*, Chapter 14

(Cold Spring Harbor Laboratory, Cold Spring Harbor, 1988). An extensive introduction as well as detailed information about all aspects of recombinant antibody technology can be found in the textbook *Recombinant Antibodies* (John Wiley & Sons, NY, 1999). A comprehensive collection of detailed antibody engineering lab Protocols can be found in R. Kontermann and S. Dübel, Eds., *The Antibody Engineering Lab Manual* (Springer Verlag, Heidelberg/New York, 2000).

**[0026]** "Derivative" as used herein refers to a chemically or biologically modified version of a compound that is structurally similar to a parent compound and (actually or theoretically) derivable from that parent compound. Generally, a "derivative" differs from an "analogue" in that a parent compound may be the starting material to generate a "derivative," whereas the parent compound may not necessarily be used as the starting material to generate an "analogue." An analogue may have different chemical or physical properties of the parent compound. For example, a derivative may be more hydrophilic or it may have altered reactivity (*e.g.*, a CDR having an amino acid change that alters its affinity for a target) as compared to the parent compound.

**[0027]** The term "biological sample" includes a blood sample, biopsy specimen, tissue explant, organ culture, biological fluid or any other tissue or cell or other preparation from a subject or a biological source. A subject or biological source may, for example, be a human or non-human animal, a primary cell culture or culture adapted cell line including genetically engineered cell lines that may contain chromosomally integrated or episomal recombinant nucleic acid sequences, somatic cell hybrid cell lines, immortalized or immortalizable cell lines, differentiated or differentiable cell lines, transformed cell lines, or the like. In further embodiments of this disclosure, a subject or biological source may be suspected of having or being at risk for having a disease, disorder or condition, including a malignant disease, disorder or condition or a B cell disorder. In certain embodiments, a subject or biological source may be suspected of having or being at risk for having a hyperproliferative, inflammatory, or autoimmune disease, and in certain other embodiments of this disclosure the subject or biological source may be known to be free of a risk or presence of such disease, disorder, or condition.

**[0028]** In certain embodiments, the present disclosure makes possible the depletion or modulation of cells associated with aberrant TGF $\beta$  activity by providing multi-specific fusion proteins that bind both a TGF $\beta$  and a second target other than TGF $\beta$ , such as IL6, IL6R, an IL6/IL6R complex, IL10, GITR, VEGF, TNF, HGF, Tumor necrosis factor-like



weak inducer of apoptosis (TWEAK; also known as tumor necrosis factor (ligand) superfamily, member 12, TNFSF12), IGF1 or IGF2. In certain embodiments, a multi-specific fusion protein comprises a first and second binding domain, a first and second linker, and an intervening domain, wherein one end of the intervening domain is fused via a linker to a first binding domain that is a TGF $\beta$ R2 ectodomain (*e.g.* an extracellular domain) and at the other end fused via a linker to a second binding domain. In some embodiments, less than an entire TGF $\beta$ R2 ectodomain is employed. Specifically, domains within the ectodomain that function as a TGF $\beta$  antagonist or confer ligand binding are employed.

**[0029]** In certain embodiments, the second binding domain is an IL6 antagonist (such as an immunoglobulin variable region that is specific for an IL6, IL6R, or IL6/IL6Ra complex), an IL10 antagonist (such as an immunoglobulin variable region that is specific for IL10, an IL10R1 ectodomain (*e.g.* SEQ ID NO:745) or a sub-domain of an IL10R1 ectodomain), a GITR agonist (such as an immunoglobulin variable region that is specific for GITR, a GITRL ectodomain (for example, amino acids 74-181 of Genbank Accession NP\_005083.2, SEQ ID NO:746) or a sub-domain of a GITRL ectodomain), a VEGF antagonist (such as an immunoglobulin variable region that is specific for VEGF, a VEGFR2 ectodomain (see, Genbank Accession NP\_002244.1, SEQ ID NO:747) or a sub-domain of a VEGFR2 ectodomain), a TNF antagonist (such as an immunoglobulin variable region that is specific for TNF, a TNFR1 ectodomain (see, Genbank Accession NP\_001056.1; SEQ ID NO:749), a sub-domain of a TNFR1 ectodomain, a TNFR2 ectodomain (see, Genbank Accession NP\_001057.1; SEQ ID NO:748), or a sub-domain of a TNFR2 ectodomain), a HGF antagonist (such as an immunoglobulin variable region that is specific for HGF, a c-Met ectodomain or a sub-domain of a c-Met ectodomain (*e.g.* SEQ ID NO:750-752)), a TWEAK antagonist (such as an immunoglobulin binding domain specific for TWEAK or TWEAKR, or a TWEAKR ectodomain (*e.g.* SEQ ID NO:761) or TWEAK binding fragment thereof), or an IGF1 or IGF2 antagonist (such as an immunoglobulin variable region that is specific for IGF1 or IGF2, an IGF1R ectodomain (for example, an IGF1R ectodomain of Genbank Accession no. NP\_000866.1 (SEQ ID NO:753) or a sub-domain thereof), or an IGFBP (for example, an IGFBP ectodomain of Genbank Accession no. NP\_000587.1 (IGFBP1; SEQ ID NO:754), NP\_000588.2 (IGFBP2; SEQ ID NO:755), NP\_001013416.1 (IGFBP3 isoform a; SEQ ID NO:756), NP\_000589.2 (IGFBP3 isoform b; SEQ ID NO:757), NP\_001543.2 (IGFBP4; SEQ ID NO:758), NP\_000590.1 (IGFBP5; SEQ ID NO:759) or NP\_002169.1 (IGFBP6; SEQ ID NO:760)), or a sub-domain thereof).



<sup>3</sup>/sec, about 10<sup>-4</sup>/sec, about 10<sup>-5</sup>/sec, about 10<sup>-6</sup>/sec, about 10<sup>-7</sup>/sec, about 10<sup>-8</sup>/sec, about 10<sup>-9</sup>/sec, about 10<sup>-10</sup>/sec, or less (*see* Graff *et al.* (2004) Protein Eng. Des. Sel. 17:293). In some embodiments, a TGFβ antagonist or fusion protein thereof of this disclosure will bind TGFβ with higher affinity and have a lower k<sub>OFF</sub> rate as compared to the cognate TGFβ receptor binding to TGFβ. In further embodiments, a TGFβ antagonist or fusion protein thereof of this disclosure that blocks or alters TGFβ dimerization or other cell surface activity may have a more moderate affinity (*i.e.*, a K<sub>d</sub> of about 10<sup>-8</sup> M to about 10<sup>-9</sup> M) and a more moderate off rate (*i.e.*, a k<sub>OFF</sub> closer to about 10<sup>-4</sup>/sec) as compared to the affinity and dimerization rate of cognate TGFβ receptor.

**[0034]** Exemplary binding domains that function as TGFβ antagonists of this disclosure can be generated as described herein or by a variety of methods known in the art (*see, e.g.*, US Patent Nos. 6,291,161; 6,291,158). Sources include antibody gene sequences from various species (which can be formatted as scFvs or Fabs, such as in a phage library), including human, camelid (from camels, dromedaries, or llamas; Hamers-Casterman *et al.* (1993) Nature, 363:446 and Nguyen *et al.* (1998) J. Mol. Biol., 275:413), shark (Roux *et al.* (1998) Proc. Nat'l. Acad. Sci. (USA) 95:11804), fish (Nguyen *et al.* (2002) Immunogenetics, 54:39), rodent, avian, ovine, sequences that encode random peptide libraries or sequences that encode an engineered diversity of amino acids in loop regions of alternative non-antibody scaffolds, such as fibrinogen domains (*see, e.g.*, Weisel *et al.* (1985) Science 230:1388), Kunitz domains (*see, e.g.*, US Patent No. 6,423,498), lipocalin domains (*see, e.g.*, WO 2006/095164), V-like domains (*see, e.g.*, US Patent Application Publication No. 2007/0065431), C-type lectin domains (Zelensky and Gready (2005) FEBS J. 272:6179), or the like. Additionally, traditional strategies for hybridoma development using a synthetic TGFβ or single chain TGFβR2 ectodomain as an immunogen in convenient systems (*e.g.*, mice, HuMAb mouse®, TC mouse™, KM-mouse®, llamas, chicken, rats, hamsters, rabbits, *etc.*) can be used to develop binding domains of this disclosure.

**[0035]** In an illustrative example, TGFβ antagonists of this disclosure specific for a TGFβ can be identified using a Fab phage library of fragments (*see, e.g.*, Hoet *et al.* (2005) Nature Biotechnol. 23:344) by screening for binding to a synthetic or recombinant TGFβ (using an amino acid sequence or fragment thereof as set forth in GenBank Accession No. NP\_000651.3). A TGFβ, as described herein or known in the art, can be used for such a screening. In certain embodiments, a TGFβ used to generate a TGFβ antagonist can further comprise an intervening domain or a dimerization domain, as described herein, such as an immunoglobulin Fc domain or fragment thereof.

**[0036]** In some embodiments, TGF $\beta$  antagonist domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains as described herein. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent (*e.g.*, mouse, rat), humanized, or human. In further embodiments, there are provided TGF $\beta$  antagonist domains of this disclosure that have a sequence that is at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to the amino acid sequence of one or more light chain variable regions (V<sub>L</sub>) or to one or more heavy chain variable regions (V<sub>H</sub>), or both, wherein each CDr has up to three amino acid changes (*i.e.*, many of the changes are in the framework region(s)), as set forth herein.

**[0037]** In further embodiments, TGF $\beta$  antagonist domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains as set forth herein, which are at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of such V<sub>H</sub> domain, V<sub>L</sub> domain, or both wherein each CDr has at most up to three amino acid changes (*i.e.*, many of the changes are in the framework region(s)).

**[0038]** The terms "identical" or "percent identity," in the context of two or more polypeptide or nucleic acid molecule sequences, means two or more sequences or subsequences that are the same or have a specified percentage of amino acid residues or nucleotides that are the same over a specified region (*e.g.*, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identity), when compared and aligned for maximum correspondence over a comparison window, or designated region, as measured using methods known in the art, such as a sequence comparison algorithm, by manual alignment, or by visual inspection. For example, preferred algorithms suitable for determining percent sequence identity and sequence similarity are the BLAST and BLAST 2.0 algorithms, which are described in Altschul *et al.* (1977) *Nucleic Acids Res.* 25:3389 and Altschul *et al.* (1990) *J. Mol. Biol.* 215:403, respectively.

**[0039]** In any of these or other embodiments described herein, the V<sub>L</sub> and V<sub>H</sub> domains may be arranged in either orientation and may be separated by about a five to about a thirty amino acid linker as disclosed herein or any other amino acid sequence capable of providing a spacer function compatible with interaction of the two sub-binding domains. In certain embodiments, a linker joining the V<sub>H</sub> and V<sub>L</sub> domains comprises an amino acid sequence as set forth in SEQ ID NO:497-604 and 1223-1228, such as Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576). Multi-specific binding domains will have at least

two specific sub-binding domains, by analogy to camelid antibody organization, or at least four specific sub-binding domains, by analogy to the more conventional mammalian antibody organization of paired V<sub>H</sub> and V<sub>L</sub> chains.

**[0040]** In further embodiments, TGF $\beta$  antagonist domains and fusion proteins thereof of this disclosure may comprise a binding domain including one or more complementarity determining region ("CDR"), or multiple copies of one or more such CDRs, which have been obtained, derived, or designed from variable regions of an anti-TGF $\beta$  or anti-TGF $\beta$ R2 scFv or Fab fragment or from heavy or light chain variable regions thereof.

**[0041]** CDRs are defined in various ways in the art, including the Kabat, Chothia, AbM, and contact definitions. The Kabat definition is based on sequence variability and is the most commonly used definition to predict CDR regions (Johnson *et al.* (2000) *Nucleic Acids Res.* 28:214). The Chothia definition is based on the location of the structural loop regions (Chothia *et al.* (1986) *J. Mol. Biol.* 196:901; Chothia *et al.* (1989) *Nature* 342:877). The AbM definition, a compromise between the Kabat and Chothia definitions, is an integral suite of programs for antibody structure modeling produced by the Oxford Molecular Group (Martin *et al.* (1989) *Proc. Nat'l. Acad. Sci. (USA)* 86:9268; Rees *et al.*, ABMTM, a computer program for modeling variable regions of antibodies, Oxford, UK; Oxford Molecular, Ltd.). An additional definition, known as the contact definition, has been recently introduced (*see* MacCallum *et al.* (1996) *J. Mol. Biol.* 5:732), which is based on an analysis of available complex crystal structures.

**[0042]** By convention, the CDR domains in the heavy chain are referred to as H1, H2, and H3, which are numbered sequentially in order moving from the amino terminus to the carboxy terminus. The CDR-H1 is about ten to 12 residues in length and starts four residues after a Cys according to the Chothia and AbM definitions, or five residues later according to the Kabat definition. The H1 can be followed by a Trp, Trp-Val, Trp-Ile, or Trp-Ala. The length of H1 is approximately ten to 12 residues according to the AbM definition, while the Chothia definition excludes the last four residues. The CDR-H2 starts 15 residues after the end of H1 according to the Kabat and AbM definitions, which is generally preceded by sequence Leu-Glu-Trp-Ile-Gly (but a number of variations are known) and is generally followed by sequence Lys/Arg-Leu/Ile/Val/Phe/Thr/Ala-Thr/Ser/Ile/Ala. According to the Kabat definition, the length of H2 is about 16 to 19 residues, while the AbM definition predicts the length to be nine to 12 residues. The CDR-H3 usually starts 33 residues after the end of H2, is generally preceded by the amino acid sequence Cys-Ala-Arg

and followed by the amino acid Gly, and has a length that ranges from three to about 25 residues.

[0043] By convention, the CDR regions in the light chain are referred to as L1, L2, and L3, which are numbered sequentially in order moving from the amino terminus to the carboxy terminus. The CDR-L1 generally starts at about residue 24 and generally follows a Cys. The residue after the CDR-L1 is always Trp, which begins one of the following sequences: Trp-Tyr-Gln, Trp-Leu-Gln, Trp-Phe-Gln, or Trp-Tyr-Leu. The length of CDR-L1 is approximately ten to 17 residues. The CDR-L2 starts about 16 residues after the end of L1 and will generally follow residues Ile-Tyr, Val-Tyr, Ile-Lys, or Ile-Phe. The CDR-L2 is about seven residues in length. The CDR-L3 usually starts 33 residues after the end of L2 and generally follows a Cys, which is generally followed by the sequence Phe-Gly-XXX-Gly and has a length of about seven to 11 residues. A binding domain of this disclosure can comprise a single CDR from a variable region of an anti-TGF $\beta$  or anti-TGF $\beta$ R2, or it can comprise multiple CDRs that can be the same or different.

[0044] Thus, a binding domain of this disclosure can comprise a single CDR from a variable region of an anti-TGF $\beta$  or anti-TGF $\beta$ R2, or it can comprise multiple CDRs that can be the same or different. In certain embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a TGF $\beta$  or TGF $\beta$ R2 comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises an amino acid sequence of a heavy chain CDR3; or (b) the V<sub>L</sub> domain comprises an amino acid sequence of a light chain CDR3; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b) and wherein the V<sub>H</sub> and V<sub>L</sub> are found in the same reference sequence. In further embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a TGF $\beta$  or TGF $\beta$ R2 comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises an amino acid sequence of a heavy chain CDR1, CDR2, and CDR3; or (b) the V<sub>L</sub> domain comprises an amino acid sequence of a light chain CDR1, CDR2, and CDR3; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b), wherein the V<sub>H</sub> and V<sub>L</sub> amino acid sequences are from the same reference sequence.

[0045] In any of the embodiments described herein comprising specific CDRs, a binding domain can comprise (i) a V<sub>H</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the amino

acid sequence of a  $V_H$  domain, wherein each CDR has at most three amino acid changes (*i.e.*, many of the changes will be in the framework regions); or (ii) a  $V_L$  domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the amino acid sequence of a  $V_L$  domain, wherein each CDR has at most three amino acid changes (*i.e.*, many of the changes will be in the framework regions); or (iii) both a  $V_H$  domain of (i) and a  $V_L$  domain of (ii); or both a  $V_H$  domain of (i) and a  $V_L$  domain of (ii) wherein the  $V_H$  and  $V_L$  are from the same reference sequence.

[0046] A TGF $\beta$  antagonist domain of fusion proteins of this disclosure may be an immunoglobulin-like domain such as an immunoglobulin scaffold. Immunoglobulin scaffolds contemplated by this disclosure include a scFv, a domain antibody or a heavy chain-only antibody. In a scFv, this disclosure contemplates the heavy and light chain variable regions are joined by any linker peptide known in the art to be compatible with domain or region joiner in a binding molecule. Exemplary linkers are linkers based on the Gly<sub>4</sub>Ser linker motif, such as (Gly<sub>4</sub>Ser)<sub>n</sub>, wherein n=1-5. If a binding domain of a fusion protein of this disclosure is based on a non-human immunoglobulin or includes non-human CDRs, the binding domain may be “humanized” according to methods known in the art.

[0047] Alternatively, a TGF $\beta$  antagonist domain of fusion proteins of this disclosure may be a scaffold other than an immunoglobulin scaffold. Other scaffolds contemplated by this disclosure present the TGF $\beta$ -specific CDR(s) in a functional conformation. Other scaffolds contemplated include, but are not limited to, an A domain molecule, a fibronectin III domain, an anticalin, an ankyrin-repeat engineered binding molecule, an adnectin, a Kunitz domain or a protein AZ domain affibody.

### **IL6 Antagonists**

[0048] As noted above, in certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is an IL6 antagonist (*e.g.*, preferentially inhibits IL6 trans-signaling or inhibits both IL6 cis- and trans-signaling). In certain embodiments, the present disclosure provides multi-specific fusion proteins containing a binding region or domain specific for an IL6/IL6R complex that has one or more of the following properties: (1) greater or equal affinity for an IL6/IL6R complex than for IL6 or IL6R $\alpha$  alone or has greater affinity for IL6R $\alpha$  alone or an IL6/IL6R complex than for IL6 alone, (2) competes with membrane gp130 for binding with a sIL6/IL6R complex or augments soluble gp130 binding to sIL6/IL6R complex, (3) preferentially inhibits IL6 trans-signaling over IL6 cis-signaling, or (4) does not inhibit signaling of gp130 family cytokines

other than IL6. In certain preferred embodiments, a binding domain specific for an IL6/IL6R complex according to this disclosure has the following properties: (1) greater affinity for IL6R $\alpha$  alone or an IL6xR complex than for IL6 alone, (2) augments soluble gp130 binding to sIL6/IL6R complex, (3) preferentially inhibits IL6 trans-signaling over IL6 cis-signaling, and (4) does not inhibit signaling of gp130 family cytokines other than IL6. For example, a binding region or domain specific for an IL6/IL6R complex may be an immunoglobulin variable binding domain or derivative thereof, such as an antibody, Fab, scFv, or the like. In the context of this disclosure, it should be understood that a binding region or domain specific for an IL6/IL6R complex is not gp130 as described herein.

[0049] As used herein, “IL6xR complex” or “IL6xR” refers to a complex of an IL6 with an IL6 receptor, wherein the IL6 receptor (also known as, for example, IL6R $\alpha$ , IL6RA, IL6R1, and CD126) is either a membrane protein (referred to herein as mIL6R or mIL6R $\alpha$ ) or a soluble form (referred to herein as sIL6R or sIL6R $\alpha$ ). The term “IL6R” encompasses both mIL6R $\alpha$  and sIL6R $\alpha$ . In one embodiment, IL6xR comprises a complex of IL6 and mIL6R $\alpha$ . In certain embodiments, the IL6xR complex is held together via one or more covalent bonds. For example, the carboxy terminus of an IL6R can be fused to the amino-terminus of an IL6 via a peptide linker, which is known in the art as a Hyper-IL6 (*see, e.g., Fischer et al. (1997) Nat. Biotechnol. 15:142*). A Hyper-IL6 linker can be comprised of a cross-linking compound, a one to 50 amino acid sequence, or a combination thereof. A Hyper-IL6 may further include a dimerization domain, such as an immunoglobulin Fc domain or an immunoglobulin constant domain sub-region. In certain embodiments, the IL6xR complex is held together via non-covalent interactions, such as by hydrogen bonding, electrostatic interactions, Van der Waal’s forces, salt bridges, hydrophobic interactions, or the like, or any combination thereof. For example, an IL6 and IL6R can naturally associate non-covalently (*e.g., as found in nature, or as synthetic or recombinant proteins*) or each can be fused to a domain that promotes multimerization, such as an immunoglobulin Fc domain, to further enhance complex stability.

[0050] As used herein, “gp130” refers to a signal transduction protein that binds to an IL6xR complex. The gp130 protein can be in a membrane (m gp130), soluble (s gp130), or any other functional form thereof. Exemplary gp130 proteins have a sequence as set forth in GenBank Accession No. NP\_002175.2 or any soluble or derivative form thereof (*see, e.g., Narazaki et al. (1993) Blood 82:1120 or Diamant et al. (1997) FEBS Lett. 412:379*). By way of illustration and not wishing to be bound by theory, an m gp130 protein can bind to either an IL6/mILR or an IL6/sILR complex, whereas a s gp130 primarily binds with an IL6/sILR



complex (*see Scheller et al.* (2006) *Scand. J. Immunol.* 63:321). Thus, certain embodiments of binding domains, or fusion proteins thereof, of the instant disclosure can inhibit IL6xR complex trans-signaling by binding with higher affinity to IL6xR than to either IL6 or IL6R $\alpha$  alone and preferably by competing with sIL6xR complex binding to gp130. A binding domain of the instant disclosure “competes” with gp130 binding to a sIL6xR when (1) a binding domain or fusion protein thereof prevents gp130 from binding a sIL6xR and the binding domain binds sIL6xR with equal or higher affinity as compared to the binding of gp130 with sIL6xR, or (2) a binding domain or fusion protein thereof enhances or promotes gp130 binding to sIL6xR.

**[0051]** In one aspect, an IL6 antagonist of this disclosure has an affinity for IL6 or IL6xR complex that is at least 2-fold to 1000-fold greater than for IL6R $\alpha$  alone or has an affinity for IL6R $\alpha$  or IL6xR complex that is at least 2-fold to 1000-fold greater than for IL6 alone. By binding to IL6, IL6R, or IL6xR complex, an IL6 antagonist of this disclosure preferentially inhibits IL6 cis- and trans-signaling. In certain embodiments, the affinity of a binding domain for IL6 or sIL6xR complex is about the same as the affinity of gp130 for IL6xR complex – with “about the same” meaning equal or up to about 2-fold higher affinity. In certain embodiments, the affinity of the binding domain for IL6, IL6R, or IL6xR complex is higher than the affinity of gp130 for IL6xR complex by at least 2-fold, at least 3-fold, at least 4-fold, at least 5-fold, at least 6-fold, at least 7-fold, at least 8-fold, at least 9-fold, at least 10-fold, at least 15-fold, at least 20-fold, at least 25-fold, at least 50-fold, at least 100-fold, 1000-fold, or greater. For example, if the affinity of gp130 for a IL6xR complex is about 2 nM (*see, e.g., Gaillard et al.* (1999) *Eur. Cytokine Netw.* 10:337), then a binding domain having at least a 10-fold higher affinity for the IL6xR complex would have a dissociation constant ( $K_d$ ) of about 0.2 nM or less.

**[0052]** In further embodiments, an IL6 antagonist binding domain of this disclosure comprises a polypeptide sequence that (a) binds to a sIL6xR complex with an affinity at least 2-fold, 10-fold, 25-fold, 50-fold, 75-fold to 100-fold, 100-fold to 1000-fold higher than for either IL6 or IL6R $\alpha$  alone and (b) competes with membrane gp130 for binding to sIL6xR complex or augments soluble gp130 binding to sIL6xR complex. In further embodiments, a polypeptide binding domain of this disclosure that binds to a sIL6xR complex with an affinity at least 2-fold, 10-fold, 25-fold, 50-fold, 75-fold to 100-fold, 100-fold to 1000-fold higher than for either IL6 or IL6R $\alpha$  alone may also (i) more significantly or preferentially inhibit IL6 trans-signaling over IL6 cis-signaling, (ii) not inhibit signaling of gp130 cytokine family members other than IL6, (iii) preferentially inhibit IL6 trans-signaling over IL6 cis-signaling

and not detectably inhibit signaling of gp130 family cytokines other than IL6, (iv) may have two or more of these properties, or (v) may have all of these properties.

[0053] In certain embodiments, a polypeptide IL6 antagonist binding domain of this disclosure binds to a sIL6xR complex with an affinity at least 2-fold to 1000-fold higher than for either IL6 or IL6R $\alpha$  alone and more significantly or preferentially inhibits IL6 trans-signaling over IL6 cis-signaling. To “preferentially inhibit IL6 trans-signaling over IL6 cis-signaling” refers to altering trans-signaling to an extent that sIL6xR activity is measurably decreased while the decrease in IL6 cis-signaling is not substantially altered (*i.e.*, meaning inhibition is minimal, non-existent, or not measurable). For example, a biomarker for sIL6xR activity (*e.g.*, acute phase expression of antichymotrypsin (ACT) in HepG2 cells) can be measured to detect trans-signaling inhibition. A representative assay is described by Jostock *et al.* (Eur. J. Biochem., 2001) – briefly, HepG2 cells can be stimulated to overexpress ACT in the presence of sIL6xR (trans-signaling) or IL6 (cis-signaling), but adding spg130 will inhibit the overexpression of ACT induced by sIL6xR while not substantially affecting IL6 induced expression. Similarly, a polypeptide binding domain of this disclosure that preferentially inhibits IL6 trans-signaling over IL6 cis-signaling will inhibit the overexpression of ACT induced by sIL6xR (*i.e.*, inhibit trans-signaling) while not substantially affecting IL6 induced expression (*i.e.*, not measurably decrease cis-signaling). This and other assays known in the art can be used to measure preferential inhibition of IL6 trans-signaling over IL6 cis-signaling (*see, e.g.*, other biomarkers described in Sporri *et al.* (1999) Int. Immunol. 11:1053; Mihara *et al.* (1995) Br. J. Rheum. 34:321; Chen *et al.* (2004) Immun. 20:59).

[0054] In further embodiments, signaling by gp130 family cytokines other than IL6 is not substantially inhibited by binding domain polypeptides or multi-specific fusion proteins thereof of this disclosure. For example, cis- and trans-signaling by an IL6xR complex via gp130 will be inhibited, but signaling by one or more other gp130 family cytokines will be minimally affected or unaffected, such as signaling via leukemia inhibitory factor (LIF), ciliary neurotropic factor (CNTF), neuropoietin (NPN), cardiotropin like cytokine (CLC), oncostatin M (OSM), IL-11, IL-27, IL-31, cardiotrophin-1 (CT-1), or any combination thereof.

[0055] It will be appreciated by those skilled in the art that the preferred *in vivo* half-life of a binding domain of this disclosure is on the order of days or weeks, but while the binding domain concentration may be low, the target may be plentiful as both IL6 and sIL6 production can be quite elevated in disease states (*see, e.g.*, Lu *et al.* (1993) Cytokine 5:578).

Thus, in certain embodiments, a binding domain of this disclosure has a  $k_{\text{OFF}}$  of about  $10^{-5}$ /sec (e.g., about a day) or less. In certain embodiments, the  $k_{\text{OFF}}$  can range from about  $10^{-1}$ /sec, about  $10^{-2}$ /sec, about  $10^{-3}$ /sec, about  $10^{-4}$ /sec, about  $10^{-5}$ /sec, about  $10^{-6}$ /sec, about  $10^{-7}$ /sec, about  $10^{-8}$ /sec, about  $10^{-9}$ /sec, about  $10^{-10}$ /sec, or less.

**[0056]** In an illustrative example, binding domains of this disclosure specific for an IL6 or IL6xR complex were identified in a Fab phage library of fragments (*see* Hoet *et al.* (2005) Nature Biotechnol. 23:344) by screening for binding to a synthetic IL6xR complex. The synthetic IL6xR complex used for this screening comprises a structure of N-IL6R $\alpha$ (frag)-L1-IL6(frag)-L2-ID-C, wherein N is the amino-terminus and C is the carboxy-terminus, IL6R $\alpha$ (frag) is a fragment of full length IL6R $\alpha$ , IL6(frag) is a fragment of IL6, L1 and L2 are linkers, and ID is an intervening or dimerization domain, such as an immunoglobulin Fc domain.

**[0057]** More specifically, an IL6xR (which is a form of Hyper IL6) used to identify the binding domains specific for IL6xR complex has a structure, from amino-terminus to carboxy-terminus, as follows: (a) a central fragment of 212 amino acids from IL6R $\alpha$  that is missing the first 110 amino acids of the full length protein and a carboxy-terminal portion that will depend on the isoform used (*see* GenBank Accession No. NP\_000556.1, isoform 1 or NP\_852004.1, isoform 2) fused to (2) a linker of G<sub>3</sub>S that is in turn fused to (3) a 175 amino acid carboxy-terminal fragment of IL6 (*i.e.*, missing the first 27 amino acids of the full length protein; GenBank Accession No. NP\_000591.1) that is in turn fused to (4) a linker that is an IgG2A hinge as set forth in SEQ ID NO:589, which is finally fused to a dimerization domain comprised of an immunoglobulin G1 (IgG1) Fc domain. In certain embodiments, the dimerization domain comprised of an IgG1 Fc domain has one or more of the following amino acids mutated (*i.e.*, have a different amino acid at that position): leucine at position 234 (L234), leucine at position 235 (L235), glycine at position 237 (G237), glutamate at position 318 (E318), lysine at position 320 (K320), lysine at position 322 (K322), or any combination thereof (EU numbering). For example, any one of these amino acids can be changed to alanine. In a further embodiment, an IgG1 Fc domain has each of L234, L235, G237, E318, K320, and K322 (according to Kabat numbering) mutated to an alanine (*i.e.*, L234A, L235A, G237A, E318A, K320A, and K322A, respectively).

**[0058]** In one embodiment, an IL6xR complex used to identify the IL6 antagonist binding domains of this disclosure has an amino acid sequence as set forth in SEQ ID NO:606. In certain embodiments, there are provided polypeptides containing a binding domain specific for an IL6xR complex, wherein the IL6xR is a sIL6xR and has the amino

acid sequence as set forth in SEQ ID NO:606. In further embodiments, polypeptides containing a binding domain specific for an IL6xR complex (1) have greater or equal affinity for an IL6xR complex than for IL6 or IL6R $\alpha$  alone, or have greater affinity for IL6R $\alpha$  alone or an IL6xR complex than for IL6 alone, (2) compete with membrane gp130 for binding with a sIL6xR complex or augment soluble gp130 binding to sIL6xR complex, (3) preferentially inhibit IL6 trans-signaling over IL6 cis-signaling, or (4) do not inhibit signaling of gp130 family cytokines other than IL6, (5) have any combination thereof of properties (1) – (4), or (6) have all of the properties of (1) – (4). Other exemplary IL6xR complexes that may be used to identify binding domains of the instant disclosure or used as a reference complex to measure any of the aforementioned binding properties are described, for example, in US Patent Publication Nos. 2007/0172458; 2007/0031376; and US Patent Nos. 7,198,781; 5,919,763.

**[0059]** In some embodiments, IL6 antagonist binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an IL6, IL6R, or IL6xR complex as described herein, and preferably human IL6, human IL6R, or human IL6xR complex. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent, (*e.g.*, mouse, rat), humanized, or human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for IL6, IL6R, or IL6xR are set forth in SEQ ID NOS:435-496 and 373-434, respectively. In further embodiments, there are provided polypeptide binding domains specific for an IL6xR wherein the binding domain comprises a sequence that is at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to the amino acid sequence of one or more light chain variable regions (V<sub>L</sub>) or to one or more heavy chain variable regions (V<sub>H</sub>), or both, as set forth in SEQ ID NOS:373-434 and 435-496, respectively, wherein each CDR has up to three amino acid changes (*i.e.*, many of the changes are found in one or more of the framework regions).

**[0060]** In further embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an IL6xR as set forth in SEQ ID NOS:435-496 and 373-434, respectively, which are at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of such V<sub>H</sub> domain, V<sub>L</sub> domain, or both, wherein each CDR has zero, one, two, or three amino acid changes. For example, the amino acid sequence of a V<sub>H</sub> domain, V<sub>L</sub> domain, or both of this disclosure can

be at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of V<sub>H</sub> domain (*e.g.*, amino acids 512 to 636), V<sub>L</sub> domain (*e.g.*, amino acids 652 to 759), or both, respectively, from an exemplary xceptor molecule containing binding domain TRU6-1002 (*see* SEQ ID NO:608), wherein each CDR has zero, one, two, or three amino acid changes.

**[0061]** In any of these or other embodiments described herein, the V<sub>L</sub> and V<sub>H</sub> domains may be arranged in either orientation and may be separated by up to about a ten amino acid linker as disclosed herein or any other amino acid sequence capable of providing a spacer function compatible with interaction of the two sub-binding domains. In certain embodiments, a linker joining the V<sub>H</sub> and V<sub>L</sub> domains comprises an amino acid sequence as set forth in SEQ ID NO:497-604 and SEQ ID NO:1223-1228, such as Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576).

**[0062]** In further embodiments, IL6 antagonist binding domains of this disclosure may comprise one or more complementarity determining region ("CDR"), or multiple copies of one or more such CDRs, which have been obtained, derived, or designed from variable regions of an anti-IL6, anti-IL6R, or anti-IL6xR complex scFv or Fab fragment or from heavy or light chain variable regions thereof. Thus, a binding domain of this disclosure can comprise a single CDR from a variable region of an IL6 or anti-IL6xR, or it can comprise multiple CDRs that can be the same or different. In certain embodiments, IL6 antagonist binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises the amino acid sequence of a heavy chain CDR3 found in any one of SEQ ID NOS:435-496; or (b) the V<sub>L</sub> domain comprises the amino acid sequence of a light chain CDR3 found in any one of SEQ ID NOS:373-434; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b) and wherein the V<sub>H</sub> and V<sub>L</sub> are found in the same reference sequence. In further embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an IL6xR complex comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises the amino acid sequence of a heavy chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:435-496; or (b) the V<sub>L</sub> domain comprises the amino acid sequence of a light chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:373-434; or (c) the binding domain comprises a

V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b), wherein the V<sub>H</sub> and V<sub>L</sub> amino acid sequences are from the same reference sequence. Exemplary light and heavy chain variable domain CDRs directed against IL6, IL6R, or IL6xR complex are provided in SEQ ID NO:1-186 and 1187-1192, and 187-372 and 1193-1198, respectively.

[0063] Amino acid sequences of IL6 antagonist light chain variable regions are provided in SEQ ID NO:373-434 and 1199-1204, with the corresponding heavy chain variable regions being provided in SEQ ID NO:435-496 and 1205-1210, respectively.

[0064] In any of the embodiments described herein comprising specific CDRs against IL6, IL6R, or IL6xR, a binding domain can comprise (i) a V<sub>H</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the amino acid sequence of a V<sub>H</sub> domain found in any one of SEQ ID NOS:435-496; or (ii) a V<sub>L</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the amino acid sequence of a V<sub>L</sub> domain found in any one of SEQ ID NOS:373-434; or (iii) both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii); or both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii) wherein the V<sub>H</sub> and V<sub>L</sub> are from the same reference sequence.

[0065] In certain embodiments, a binding domain of this disclosure may be an immunoglobulin-like domain, such as an immunoglobulin scaffold. Immunoglobulin scaffolds contemplated in this disclosure include a scFv, Fab, a domain antibody, or a heavy chain-only antibody. In further embodiments, there are provided anti-IL6 or anti-IL6xR antibodies (*e.g.*, non-human such as mouse or rat, chimeric, humanized, human) or Fab fragments or scFv fragments that have an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% identical to the amino acid sequence of a V<sub>H</sub> and V<sub>L</sub> domain set in any one of SEQ ID NOS:435-496 and 373-434, respectively, which may also have one or more of the following properties: (1) have greater or equal affinity for an IL6xR complex than for IL6 or IL6R $\alpha$  alone, or have greater affinity for IL6R $\alpha$  alone or an IL6xR complex than for IL6 alone, (2) compete with membrane gp130 for binding with a sIL6xR complex or augment soluble gp130 binding to sIL6xR complex, (3) preferentially inhibit IL6 trans-signaling over IL6 cis-signaling, or (4) do not inhibit signaling of gp130 family cytokines other than IL6. Such antibodies, Fabs, or scFvs can be used in any of the methods described herein. In certain embodiments, the present disclosure provides polypeptides containing a binding domain that is an IL6 antagonist (*i.e.*, can inhibit IL6 cis- and trans-signaling). In further embodiments, an IL6 antagonist according to this

disclosure does not inhibit signaling of gp130 family cytokines other than IL6. Exemplary IL6 antagonists include binding domains specific for an IL6 or IL6xR, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like).

[0066] Alternatively, binding domains of this disclosure may be part of a scaffold other than an immunoglobulin. Other scaffolds contemplated include an A domain molecule, a fibronectin III domain, an anticalin, an ankyrin-repeat engineered binding molecule, an adnectin, a Kunitz domain, or a protein AZ domain affibody.

### **IL10 Antagonists**

[0067] In certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is an IL10 antagonist (*i.e.*, can inhibit IL10 signaling). Exemplary IL10 antagonists include binding domains specific for an IL10 or IL10R1, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or an IL10R1 ectodomain.

[0068] IL10 is a member of a cytokine superfamily that share an alpha-helical structure. Although no empirical evidence exists, it has been suggested that all possess six alpha-helices (Fickenscher, H. et al., 2002, Trends Immunol. 23: 89). IL10 has four cysteines, only one of which is conserved among family members. Since IL10 demonstrates a V-shaped fold that contributes to its dimerization, it appears that disulfide bonds are not critical to this structure. Amino acid identity of family members to IL10 ranges from 20% (IL-19) to 28% (IL-20) (Dumouter et al., 2002, Eur. Cytokine Netw. 13: 5).

[0069] IL10 was first described as a Th2 cytokine in mice that inhibited IFN- $\alpha$  and GM-CSF cytokine production by Th1 cells (Moore et al., 2001, Annu. Rev. Immunol. 19: 683; Fiorentino et al., 1989, J. Exp. Med. 170:2081).

[0070] Human IL10 is 178 amino acids in length with an 18 amino acid signal sequence and a 160 amino acid mature segment and a molecular weight of approximately 18 kDa (monomer). Human IL10 contains no potential N-linked glycosylation site and is not glycosylated (Dumouter et al., 2002, Eur. Cytokine Netw. 13: 5; Vieira et al., 1991, Proc. Natl. Acad. Sci. USA 88:1172). It contains four cysteine residues that form two intrachain disulfide bonds. The length of  $\alpha$ -helices A to F in human IL10 are 21, 8, 19, 20, 12 and 23 amino acids, respectively. Helices A to D of one monomer noncovalently interact with helices E and F of a second monomer, forming a noncovalent V-shaped homodimer. Functional areas have been mapped on the IL10 molecule. In the N-terminus, pre-helix A

residues no. 1-9 are involved in mast cell proliferation, while in the C-terminus, helix F residues no. 152-160 mediate leukocyte secretion and chemotaxis.

[0071] Cells known to express IL10 include CD8<sup>+</sup> T cells, microglia, CD14<sup>+</sup> (but not CD16<sup>+</sup>) monocytes, Th2 CD4<sup>+</sup> cells (mice), keratinocytes, hepatic stellate cells, Th1 and Th2 CD4<sup>+</sup> T cells (human), melanoma cells, activated macrophages, NK cells, dendritic cells, B cells (CD5<sup>+</sup> and CD19<sup>+</sup>) and eosinophils.

[0072] On T cells, the initial observations of IL10 inhibition of IFN- $\gamma$  production are now believed to be an indirect effect mediated by accessory cells. Additional effects on T cells, however, include: IL10 induced CD8<sup>+</sup> T cell chemotaxis, an inhibition of CD4<sup>+</sup> T cell chemotaxis towards IL-8, suppression of IL-2 production following activation, an inhibition of T cell apoptosis via Bcl-2 up-regulation, and an interruption of T cell proliferation following low antigen exposure accompanied by B7/CD28 costimulation (Akdis et al., 2001, Immunology 103: 131).

[0073] On B cells, IL10 has a number of related, yet distinct functions. In conjunction with TNF- $\beta$  and CD40L, IL10 induces IgA production in naïve (IgD<sup>+</sup>) B cells. It is believed that TGF- $\beta$ /CD40L promotes class switching while IL10 initiates differentiation and growth. When TGF- $\beta$  is not present, IL10 cooperates with CD40L in inducing IgG1 and IgG3 (human), and thus may be a direct switch factor for IgG subtypes. IL10 has divergent effects on IL-4 induced IgE secretion. If IL10 is present at the time of IL-4 induced class switching, it reverses the effect; if it is present after IgE commitment, it augments IgE secretion. CD27/CD70 interaction in the presence of IL10 promotes plasma cell formation from memory B cells (Agematsu et al., 1998, Blood 91: 173).

[0074] Mast cells and NK cells are also impacted by IL10. On mast cells, IL10 induces histamine release while blocking GM-CSF and TNF- $\alpha$  release. This effect may be autocrine as IL10 is known to be released by mast cells in rat. As evidence of its pleiotrophic nature, IL10 has the opposite effects on NK cells. Rather than blocking TNF- $\alpha$  and GM-CSF production, IL10 actually promotes this function on NK cells. In addition, it potentiates IL-2 induced NK cell proliferation and facilitates IFN- $\gamma$  secretion in NK cells primed by IL-18. In concert with both IL-12 and/or IL-18, IL10 potentiates NK cell cytotoxicity (Cai et al., 1999, Eur. J. Immunol. 29: 2658).

[0075] IL10 has a pronounced anti-inflammatory impact on neutrophils. It inhibits the secretion of the chemokines MIP-1 $\alpha$ , MIP-1 $\beta$  and IL-8, and blocks production of the proinflammatory mediators IL-1  $\beta$  and TNF- $\alpha$ . In addition, it decreases the ability of



neutrophils to produce superoxide, and as a result interferes with PMN-mediated antibody-dependent cellular cytotoxicity. IL10 also blocks IL-8 and fMLP-induced chemotaxis, possibly via CXCR1 (Vicioso et al., 1998 Eur. Cytokine Netw. 9: 247).

**[0076]** On dendritic cells (DCs), IL10 generally exhibits immunosuppressive effects. It would appear to promote CD14<sup>+</sup> macrophage differentiation at the expense of DCs. Macrophages, while phagocytic, are poor antigen-presenting cells. IL10 seems to decrease the ability of DCs to stimulate T cells, particularly for Th1 type cells. How IL10 accomplishes this is unclear, as the data within the literature is conflicting. Relative to MHC-II expression, it can be down-regulated, unchanged, or up-regulated (Sharma et al., 1999, J. Immunol. 163:5020). With respect to B7-1/CD80, IL10 will either up-regulate or down-regulate its expression. B7-2/CD86 plays a key role in T cell activation. For this molecule, IL10 is involved in both up-regulation and down-regulation. Perhaps the most significant modulation, however, occurs with CD40 (IL10 seems to reduce its expression). At the regional level, IL10 may block immunostimulation by inhibiting Langerhans cell migration in response to proinflammatory cytokines. Alternatively, IL10 blocks an inflammation-induced DC maturation step that normally involves CCR1, CCR2 and CCR5 down-regulation and CCR7 up-regulation. This blockage, with retention of CCR1, CCR2 and CCR5, results in a failure of DCs to migrate to regional nodes. The result is an immobile DC that will not stimulate T cells but will bind (and clear) proinflammatory chemokines without responding to them (D-Amico et al., 2000 Nat. Immunol. 1:387).

**[0077]** On monocytes, IL10 has a number of documented effects. For example, IL10 seems to clearly reduce cell surface MHC-II expression. It also inhibits IL-12 production following stimulation. While it promotes a monocyte to macrophage transition in conjunction with M-CSF, the phenotype of the macrophage is not clear (i.e. CD16<sup>+</sup>/cytotoxic vs. CD16<sup>-</sup>). IL10 also reduces monocyte GM-CSF secretion and IL-8 production, while promoting IL-1ra release (Gesser et al., 1997, Proc. Natl. Acad. Sci. USA 94:14620). Hyaluronectin, a connective tissue component, is now known to be secreted by monocytes in response to IL10. This may have some importance in cell migration, particularly tumor cell metastases, where hyaluronectin is known to interrupt cell migration through extracellular space (Gesser et al., 1997).

**[0078]** Human IL10R1 is a 90-110 kDa, single-pass type I transmembrane glycoprotein that is expressed on a limited number of cell types (Liu et al., 1994, J. Immunol. 152:1821). Weak expression is seen in pancreas, skeletal muscle, brain, heart and kidney. Placenta, lung, and liver showed intermediate levels of expression, while monocytes, B-cells,

large granular lymphocytes and T-cells express high levels (Liu et al., 1994). The expressed protein is a 578 amino acid protein that contains a 21 amino acid signal peptide, a 215 amino acid extracellular region, a 25 amino acid transmembrane segment, and a 317 amino acid cytoplasmic domain. There are two FNIII motifs within the extracellular region and a STAT3 docking site plus a JAK1 association region within the cytoplasmic domain (Kotenko et al., 2000 *Oncogene* 19:2557; Kotenko et al., 1997, *EMBO J.* 16:5894). IL10R1 binds human IL10 with a K<sub>d</sub> of about 200 pM.

[0079] In some embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an IL10 or an IL10R1. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent (*e.g.*, mouse, rat), humanized, or human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for IL10 include, but are not limited to, those disclosed in US Patent Application Publication no. US 2007/0178097A1. Binding domains of this disclosure may also, or alternatively, comprise an IL10R1 ectodomain as shown, for example, in SEQ ID NO:745, or a fragment thereof. In further embodiments, there are provided polypeptide binding domains specific for IL10, wherein the binding domain comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to an amino acid sequence of SEQ ID NO:745 or to amino acids 22-401 of SEQ ID NO:745, wherein the polypeptide binding domain binds to IL10 and inhibits the activity thereof.

### **GITR Agonists**

[0080] In certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is a GITR agonist (*i.e.*, can increase GITR signaling). Exemplary GITR agonists include binding domains specific for a GITR or GITRL, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or a GITRL ectodomain.

[0081] Glucocorticoid-induced tumor necrosis factor receptor (GITR; also known as AITR) is a type I transmembrane protein and a member of the TNF receptor superfamily (Nocentini et al., (2007) *Eur. J. Immunol.* 37:1165-9). The cytoplasmic domain has homology to the cytoplasmic domain of 4-1BB and CD27. GITR is expressed in peripheral blood T cells, bone marrow, thymus, spleen, and lymph nodes, and is constitutively expressed in CD4<sup>+</sup>CD25<sup>+</sup> regulatory T cells (Kwon et al., (2003) *Exp. Mol. Med.* 35:13). In addition, it is constitutively expressed at low levels in natural killer (NK) cells and is induced upon

stimulation by either Toll-like receptor ligand or IL-15 (Liu et al., (2008) J. Biol. Chem. 283:8202).

**[0082]** Expression of GITR is increased following T cell activation. Activation of GITR coactivates effector T lymphocytes and modulates regulatory T cell activity. Binding of GITR to its ligand GITRL has been shown to render CD4<sup>+</sup>CD25<sup>-</sup> effector T cells resistant to the inhibitory effects of CD4<sup>+</sup>CD25<sup>+</sup> regulatory T cells.

**[0083]** GITR ligand (GITRL) is a type II membrane protein. It is 173 amino acids long with a predicted molecular weight of 20 kDa. The experimental molecular weight of 25-28 kDa is suggestive of glycosylation. GITRL is expressed in antigen presenting cells (APC) and is constitutively expressed in human umbilical vein endothelial cells (Nocentini et al. *Ibid*). It is not, however, expressed in resting or stimulated T cells, B cell lines, or peripheral blood mononuclear cells.

**[0084]** The GITR/GITRL system has been shown to increase resistance to tumors and viral infections (Nocentini et al., *ibid*). Specifically, the anti-GITR monoclonal antibody DTA-1 was shown to inhibit regulatory T cell-dependent suppression and enhance T cell responses. Administration of DTA-1 in mice induced B16 melanoma tumor rejection. GITR is also involved in autoimmune/inflammatory processes and regulates leukocyte extravasation. GITR<sup>-/-</sup> mice exhibit decreased sensitivity to inflammatory disease conditions, indicating a positive role for GITR in inflammation.

**[0085]** In some embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a GITR or a GITRL. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent (e.g., mouse, rat), humanized, or human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for GITR include, but are not limited to, those disclosed in US Patent Application Publication no. US 2007/0098719A1. Binding domains of this disclosure may also, or alternatively, comprise a GITRL ectodomain (e.g. amino acids 74-181 of Genbank Accession NP\_005083.2 (SEQ ID NO:746) or a fragment thereof. In further embodiments, there are provided polypeptide binding domains specific for GITR, wherein the binding domain comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to amino acids 74-181 of SEQ ID NO:746, wherein the polypeptide binding domain binds to GITR and increases the activity thereof.

**VEGF Antagonists**

[0086] In certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is a VEGF antagonist (*i.e.*, can inhibit VEGF signaling). Exemplary VEGF antagonists include binding domains specific for a VEGF or VEGFR2, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or a VEGFR2 ectodomain.

[0087] Vascular endothelial growth factor (VEGF or VEGF-A) is an evolutionarily conserved homodimeric glycoprotein and a potent endothelial cell-specific mitogen that plays a critical role in angiogenesis and vasculogenesis (Lee et al. (2007) PLOS Medicine 6:1101-1116). VEGF induces various intracellular signaling and physiologic responses that are essential for angiogenesis, such as intracellular  $\text{Ca}^{2+}$  influx, chemotaxis (migration), expression of plasminogen activators, urokinase receptor and collagenases, and vascular permeability. Its biological effects are elicited through two high-affinity receptor tyrosine kinases, namely VEGF receptors 1 (VEGFR1) and 2 (VEGFR2), which are mainly expressed in endothelial cells.

[0088] VEGFA is a secreted protein that is a homodimer linked by disulfide bonds. It is also found as heterodimer with PlGF. Alternative splicing of VEGF mRNA results in various isoforms, which include VEGF121, VEGF145, VEGF165, VEGF189 and VEGF206, in humans and VEGF120, VEGF164 and VEGF188 in mice. Studies of genetically engineered mice expressing only one VEGF isoform indicate that VEGF isoforms have distinct yet some overlapping roles in vascular development and function as evidenced by tissue-specific vascular defects in these mice. The VEGF isoforms display differences in their biochemical properties, including receptor binding with VEGF165 and VEGF188 but not VEGF120 binding to neuropilins and heparan sulfate. The differential affinity to heparan sulfate is important in their binding to VEGFR1 and VEGFR2, as heparan sulfate can mediate the binding and transactivation of these receptors. Furthermore, differential binding to heparan sulfate is reported to lead to different VEGF actions, including endothelial cell survival, adhesion and vascular branch formation. Both VEGF164 and VEGF188 bind heparan sulfate, making them partially or fully cell-bound, respectively, whereas VEGF120 does not bind heparan sulfate, and is freely diffusible.

[0089] The VEGF isoforms display tissue-specific patterns of expression. The VEGF189, VEGF-165 and VEGF-121 isoforms are widely expressed, whereas the VEGF206 and VEGF-145 are uncommon. Its expression is regulated by growth factors, cytokines, gonadotropins, nitric oxide, hypoxia, hypoglycemia and oncogenic mutations.

[0090] The classical role of VEGF in tumor progression is as a positive regulator of angiogenesis, the process of forming new capillaries from preexisting blood vessels. Tumor growth is highly dependent on the ability of tumors to induce their own vascularization. VEGF expression has been reported in a number of cancer cell lines and in several clinical specimens derived from breast, brain, and ovarian cancers. Thus, antagonism of VEGF can effectively prevent tumor growth through incomplete blood vessel formation. VEGF exerts its effects on endothelial cells in a paracrine mode after its release by other cells such as tumor cells, or in an autocrine manner in VEGF-producing endothelial cells. VEGF binds to its cognate receptors VEGFR1 (also known as FLT1), VEGFR2 (also known as KDR or FLK1), and neuropilin 1 (NRP1).

[0091] VEGF expression in the adult is cell-type specific and is controlled at many levels from transcription to translation, and is upregulated in tumors and in various pathologic states. One of the best-characterized stimuli of VEGF transcription is hypoxia, which acts by stabilization of the hypoxia-inducible factor-1 alpha (HIF1 $\alpha$ ) transcription factor. Hypoxic regulation of VEGF also takes place post-transcriptionally via mRNA stabilization. VEGF expression is induced by other growth factors and cytokines including IGF-1, IL-6, IL-1, PDGF, TNF- $\alpha$ , TGF- $\beta$  and FGF-4. In addition, VEGF expression is also stimulated by physical forces, including stretch, with one putative transcription factor being the Kruppel like factor-2. Analysis of the VEGF promoter reveals many other potential transcription factor responsive elements, of which several pathways have been elucidated, for example EGF and HGF signaling via the SP1 responsive element.

[0092] Members of the VEGF family promote two very important processes *in vivo*, angiogenesis and lymphangiogenesis, which involve growth of new blood and lymphatic vessels from pre-existing vasculature, respectively. These processes control the normal processes of wound healing, ovarian-follicular development, endometrium growth and pathological processes such as retinopathies, rheumatoid arthritis and solid tumor growth. A newly identified splice variant of VEGF, VEGF165b, is postulated to have an inhibitory effect on angiogenesis. Lymphangiogenesis is correlated with lymph node metastasis and cancer spread via the lymphatic system.

[0093] VEGF activities are mediated by high-affinity receptor tyrosine kinases expressed primarily in endothelial cells. These are: VEGFR-1 (Flt-1) and VEGFR-2 (Flk-1/KDR), which are mainly expressed by blood vessel endothelial cells and VEGFR-3 (Flt-4) expressed in lymphatic endothelial cells. These receptors are characterized by seven extracellular immunoglobulin-like domains, which bind the growth factor, followed by a

single membrane-spanning region and a conserved intracellular tyrosine kinase domain interrupted by a kinase insert sequence. These receptors are themselves enzymes and once activated by ligand binding, they dimerize and undergo autophosphorylation. This step enhances the capacity of the receptor to directly activate other target proteins by phosphorylating them on specific tyrosine residues.

**[0094]** The VEGF-kinase ligand/receptor signaling system plays a key role in vascular development and regulation of vascular permeability. In case of HIV-1 infection, the interaction with extracellular viral Tat protein seems to enhance angiogenesis in Kaposi's sarcoma lesions.

**[0095]** Although VEGF binds to VEGFR1, VEGFR2, Nrp-1 and Nrp-2, its main signaling receptor in the endothelium is VEGFR2. VEGFR2 belongs to the family of receptor tyrosine kinases, and upon VEGF binding, there is dimerization and activation of the tyrosine kinase, resulting in phosphorylation of specific tyrosine residues on the cytoplasmic tail, which in turn promotes docking of signal transducing molecules. VEGFR2 is responsible for initiating signal transduction pathways within endothelial cells. Following the binding of VEGF to VEGFR2, VEGF mediates its effects on proliferation, survival, adhesion, migration, capillary morphogenesis, and gene expression in endothelial cells. VEGFR1 has a relatively minor role in VEGF-mediated signal transduction as compared to VEGFR2, since its kinase activity is 10-fold less than that of VEGFR2. Breast cancer cell lines express both VEGF and the VEGF receptors VEGFR1, VEGFR2, and NRP1. Recent studies have shown that VEGF acts as an autocrine growth and survival factor for VEGF receptor-expressing tumor cells. However, the mechanism by which VEGF mediates the survival of tumor cells needs to be investigated in depth (Lee *et al.*, 2007, PLOS Medicine 6: 1101-1116).

**[0096]** Although VEGFR1 is also expressed by endothelial cells (EC), it is believed to act primarily to modulate VEGFR2 signaling. Mitogenesis, chemotaxis, cell survival and changes in the morphology of endothelial cells are mainly mediated by VEGFR-2. The mitogenic signal is induced by activation of the Raf-Mek-Erk pathway, while the antiapoptotic effects and chemotaxis are mediated by PI3K/Akt activation. VEGF binding to VEGFR-2 also results in activation of several integrins, which are adhesion molecules involved in angiogenesis, in a PI3K/Akt dependent manner. Apart from being expressed in endothelial cells, VEGFR-2 is also found in haematopoietic stem cells, where it increases their survival, and in retinal progenitor cells, where it plays a critical role in neurogenesis and vasculogenesis.

[0097] In some embodiments, binding domains of this disclosure comprise  $V_H$  and  $V_L$  domains specific for a VEGF or a VEGFR2. In certain embodiments, the  $V_H$  and  $V_L$  domains are rodent (*e.g.*, mouse, rat), humanized, or human. Examples of binding domains containing such  $V_H$  and  $V_L$  domains specific for VEGF include, but are not limited to, those disclosed in US Patent Application Publication no. US 2007/0141065A1. Binding domains of this disclosure may also, or alternatively, comprise a VEGFR2 ectodomain (see, Genbank Accession NP\_002244.1, SEQ ID NO:747) or a fragment thereof. In further embodiments, there are provided polypeptide binding domains specific for VEGF, wherein the binding domain comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to an amino acid sequence of SEQ ID NO:747, wherein the polypeptide binding domain binds to VEGF and inhibits the activity thereof.

#### **TNF $\alpha$ Antagonists**

[0098] In certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is a TNF $\alpha$  antagonist (*i.e.*, can inhibit TNF $\alpha$  signaling). Exemplary TNF $\alpha$  antagonists include binding domains specific for a TNF $\alpha$ , such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or a TNFR1 or TNFR2 ectodomain.

[0099] Tumor Necrosis Factor Receptor (TNFR) is a member of the tumor necrosis factor receptor superfamily and is the receptor for Tumor Necrosis Factor- $\alpha$  (TNF $\alpha$ ), also known as CD120 or cachectin. There are two variants of this cytokine receptor, TNFR1 and TNFR2, (CD120a receptor and CD120b receptor). TNFR1 (Genbank accession no. NP\_001056.1) has a molecular weight of about 55 KD and is therefore sometimes referred to as p55. A TNFR domain that may be used as a TNF $\alpha$  binding domain in the disclosed fusion proteins is located at amino acids 44-149 of the TNFR1 sequence. TNFR2 (Genbank accession no. NP\_001057.1) has a molecular weight of about 75 KD and is therefore sometimes referred to as p75. A TNFR domain that may be used as a TNF $\alpha$  binding domain in the disclosed fusion proteins is located at amino acids 40-141 of the TNFR2 sequence.

[00100] A majority of cell types and tissues appear to express both TNF receptors. Both exist in cell surface as well as soluble forms and both are active in signal transduction, although they are able to mediate distinct cellular responses. TNFR1 appears to be responsible for signaling most TNF responses. Among other activities, TNFR2 stimulates

thymocyte proliferation, activates NF- $\kappa$ B, and is an accessory to TNFR1 in the signaling of responses primarily mediated by TNF-R1, like cytotoxicity.

[00101] TNF antagonists, such as anti-TNF antibodies, can positively affect various inflammatory conditions. For example, infliximab is indicated in the United States for the treatment of rheumatoid arthritis, Crohn's disease, ankylosing spondylitis, psoriatic arthritis, plaque psoriasis, and ulcerative colitis. Recently, perispinal delivery of the TNF $\alpha$  inhibitor etanercept has been shown to reduce symptoms in patients with Alzheimer's disease (Tobinick and Gross (2008) BMC Neurol. 8:27-36; Griffin (2008) J. Neuroinflammation, 5:3-6).

[00102] According to REMICADE® (infliximab) prescribing information, biological activities attributed to TNF include: induction of pro-inflammatory cytokines such as interleukins (IL) 1 and 6, enhancement of leukocyte migration by increasing endothelial layer permeability and expression of adhesion molecules by endothelial cells and leukocytes, activation of neutrophil and eosinophil functional activity, induction of acute phase reactants and other liver proteins, as well as tissue degrading enzymes produced by synoviocytes and/or chondrocytes.

[00103] In some embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a TNF $\alpha$ . In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for TNF $\alpha$  include, but are not limited to, those disclosed in US Patent Application Publication no. US 2007/0249813. Binding domains of this disclosure may also, or alternatively, comprise a TNFR1 ectodomain (see, Genbank Accession NP\_001056.1, SEQ ID NO:749) or a fragment thereof, or a TNFR2 ectodomain (see, Genbank Accession NP\_001057.1, SEQ ID NO:748) or a fragment thereof. TNFR1 and TNFR2 ectodomains are described in US Patent Application Publication no. US 2007/ 0128177. In further embodiments, there are provided polypeptide binding domains specific for TNF $\alpha$ , wherein the binding domain comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to an amino acid sequence of SEQ ID NO:748 or 749, wherein the polypeptide binding domain binds to TNF $\alpha$  and inhibits the activity thereof.

### **HGF Antagonists**

[00104] As noted above, in certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is a HGF antagonist (*i.e.*, can inhibit



HGF signaling). Exemplary HGF antagonists include binding domains specific for a HGF, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or a c-Met ectodomain or sub-domain thereof (*e.g.*, a Sema domain, a PSI domain, or both domains of c-Met).

**[00105]** The tyrosine kinase receptor c-Met (also known as the hepatocyte growth factor receptor, HGFR, because hepatocyte growth factor (HGF) is one of its ligands) is active during the normal processes of embryogenesis and tissue repair. In both of these processes, cells dissociate from neighboring cells and enter the bloodstream. In the bloodstream, c-Met-induced protection from apoptosis and ability to grow in an anchorage-independent manner allow the cells to survive until they extravasate, proliferate and eventually differentiate. In tissue repair, c-Met is involved in the process of epithelial-mesenchymal transition when epithelial cells adjacent to the injury detach, change shape and migrate toward the injured area where they proliferate and reconstitute the epithelial layer.

**[00106]** However, when c-Met is constitutively activated, the cells expressing it become tumorigenic and metastatic. Constitutive c-Met activation has been demonstrated to occur by multiple mechanisms. The most common is over-expression of the receptor, which occurs as a result of *c-Met* gene amplification (*e.g.*, in colorectal tumors), enhanced *c-Met* transcription induced by other oncogenes, or hypoxia-activated transcription. Another mechanism includes *c-Met* gene structural alterations including point mutations (*e.g.*, in hereditary papillary renal carcinomas, childhood hepatocellular carcinomas, sporadic papillary renal carcinomas, gastric carcinomas and head and neck squamous-cell carcinomas) and chromosomal translocations. Yet another mechanism includes c-Met structural alterations such as abnormal posttranslational processing, lack of cleavage of the precursor protein, mutations that prevent receptor downregulation and truncation of the receptor (*e.g.*, in musculoskeletal tumors). Still another mechanism is HGF-dependent autocrine/paracrine activation. Paracrine activation can become pathological in the presence of abnormal HGF production by mesenchymal cells. Autocrine activation occurs when tumor cells aberrantly express both c-Met and HGF (*e.g.*, in osteosarcomas, rhabdomyosarcomas, gliomas and carcinomas of the thyroid, breast and lung). Finally, constitutive c-Met activation can also be caused by transactivation by other membrane receptors (*e.g.*, RON, EGF-receptor family members, FAS and B plexins). *See Corso et al., TRENDS Mol. Med. 11:284 (2005).*

**[00107]** Anti-cancer strategies targeting the c-Met signaling pathway are also discussed in Corso *et al., supra*. These have included antagonism or neutralization of HGF, inhibition of c-Met kinase activity, prevention of c-Met dimerization, inhibition of c-Met

intracellular activities, and silencing of *c-Met* or *Hgf* expression. Michielli *et al.*, *Cancer Cell*, 6: 61-73 (2004) describe a soluble c-Met receptor, termed “decoy Met,” that interferes with both HGF binding to c-Met and c-Met homodimerization. Delivery of the decoy Met by a lentiviral vector in mice was reported to inhibit tumor cell proliferation and survival in human xenografts. Decoy Met was observed to impair tumor angiogenesis, suppress formation of spontaneous metastases, and synergize with radiotherapy in inducing tumor regression.

**[00108]** In some embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a HGF. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent (*e.g.*, mouse, rat), humanized, or human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for HGF include, but are not limited to, those disclosed in US Patent Application Publication no. US 2005/0118643. Binding domains of this disclosure may also, or alternatively, comprise a cMet ectodomain of SEQ ID NO:750, 751 or 752, or a fragment thereof. In further embodiments, there are provided polypeptide binding domains specific for HGF, wherein the binding domain comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to an amino acid sequence of SEQ ID NO:750, 751 or 752, wherein the polypeptide binding domain binds to HGF and inhibits the activity thereof.

**[00109]** In some embodiments, binding domains of this disclosure are c-Met antagonist domains that comprise V<sub>H</sub> and V<sub>L</sub> domains as described herein. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains are set forth in SEQ ID NOS:1132-1184 and 1079-1131, respectively. In further embodiments, there are provided c-Met antagonist domains of this disclosure that have a sequence that is at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to the amino acid sequence of one or more light chain variable regions (V<sub>L</sub>) or to one or more heavy chain variable regions (V<sub>H</sub>), or both, as set forth in SEQ ID NOS:1079-1131 and 1132-1184, respectively, wherein each CDR has at most up to three amino acid changes.

**[00110]** In further embodiments, c-Met antagonist domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains as set forth in SEQ ID NOS:1132-1184 and 1079-1131, respectively, which are at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at

least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of such V<sub>H</sub> domain, V<sub>L</sub> domain, or both, wherein each CDR has no more than zero, one, two, or three mutations. For example, the amino acid sequence of a V<sub>H</sub> domain, V<sub>L</sub> domain, or both of this disclosure can be at least 80%, at least 81%, at least 82%, at least 83%, at least 84%, at least 85%, at least 86%, at least 87%, at least 88%, at least 89%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, or at least 99.5% identical to the amino acid sequence of V<sub>H</sub> domain (SEQ ID NO:1174), V<sub>L</sub> domain (SEQ ID NO:1121), or both, respectively, from exemplary binding domain TRU(H)-343.

**[00111]** A binding domain of this disclosure can comprise a single CDR from a variable region of an anti-HGF or anti-c-Met, or it can comprise multiple CDRs that can be the same or different. In certain embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an HGF or c-Met comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises an amino acid sequence of a heavy chain CDR3 found in any one of SEQ ID NOS:1132-1184; or (b) the V<sub>L</sub> domain comprises an amino acid sequence of a light chain CDR3 found in any one of SEQ ID NOS:1079-1131; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b) and wherein the V<sub>H</sub> and V<sub>L</sub> are found in the same reference sequence. In further embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for an HGF or c-Met comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises an amino acid sequence of a heavy chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:1132-1184; or (b) the V<sub>L</sub> domain comprises an amino acid sequence of a light chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:1079-1131; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b), wherein the V<sub>H</sub> and V<sub>L</sub> amino acid sequences are from the same reference sequence.

**[00112]** In any of the embodiments described herein comprising specific CDRs, a binding domain can comprise (i) a V<sub>H</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% identical to the amino acid sequence of a V<sub>H</sub> domain found in any one of SEQ ID NOS:1132-1184; or (ii) a V<sub>L</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%,

94%, 95%, 96%, 97%, 98%, or 99% identical to the amino acid sequence of a V<sub>L</sub> domain found in any one of SEQ ID NOS:1079-1131; or (iii) both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii); or both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii) wherein the V<sub>H</sub> and V<sub>L</sub> are from the same reference sequence. Exemplary light and heavy chain variable domain CDRs directed against c-Met are provided in SEQ ID NO:762-920 and 921-1078, respectively.

[00113] Amino acid sequences of c-Met antagonist light chain and heavy chain variable regions are provided in SEQ ID NO:1079-1131 and 1132-1184, respectively.

### **TWEAK Antagonists**

[00114] In certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is a TWEAK antagonist (*i.e.*, can inhibit TWEAKR signaling). Exemplary TWEAK antagonists include binding domains specific for a TWEAK, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or a TWEAKR ectodomain or fragment thereof.

[00115] TWEAK is a cytokine that belongs to the tumor necrosis factor (TNF) ligand family and regulates multiple cellular responses including pro-inflammatory activity, angiogenesis and cell proliferation. TWEAK is a type II-transmembrane protein that is cleaved to generate a soluble cytokine with biological activity. The position of various domains within the TWEAK protein is shown, for example, in US Published Patent Application No. 2007/0280940. TWEAK has overlapping signaling functions with TNF, but displays a much wider tissue distribution. TWEAK can induce apoptosis via multiple pathways of cell death in a cell type-specific manner and has also been found to promote proliferation and migration of endothelial cells, and thus acts as a regulator of angiogenesis.

[00116] The cognate TWEAK receptor, TWEAKR or fibroblast growth factor-inducible 14 (Fn14), is a TNF receptor superfamily member expressed by non-lymphoid cell types (Wiley *et al.* (2001) *Immunity* 15:837). Expression of TWEAK and TWEAKR is relatively low in normal tissues but undergoes dramatic upregulation in settings of tissue injury and diseases. The TWEAK/R pathway facilitates acute tissue repair functions and thus functions physiologically after acute injury but functions pathologically in chronic inflammatory disease settings. In contrast to TNF, TWEAK plays no apparent role in development or homeostasis. A review of the TWEAK/R pathway is provided in Burkly *et al.* (2007) *Cytokine* 40:1. Persistently activated TWEAK promotes chronic inflammation, pathological hyperplasia and angiogenesis, and potentially impedes tissue repair by inhibiting differentiation of progenitor cells. TWEAK protein has been identified on the surface of

activated monocytes and T cells and on tumor cell lines, and intracellularly in resting and activated monocytes, dendritic cells and NK cells. TWEAK expression is significantly increased locally in target tissues in contexts of acute injury, inflammatory disease and cancer, all of which are associated with infiltration of inflammatory cells and/or activation of resident innate immune cell types. Circulating TWEAK levels have been shown to be significantly increased in patients with chronic inflammatory diseases such as multiple sclerosis and systemic lupus erythematosus.

[00117] TWEAK blocking monoclonal antibodies have been shown to be effective in a mouse collagen-induced arthritis (CIA) model (Kamata *et al.* (2006) J. Immunol. 177:6433; Perper *et al.* (2006) J. Immunol. 177:2610). The arthritogenic activities of TWEAK and TNF on human synoviocytes are often additive or synergistic and appear independent of one another, indicating that TWEAK and TNF may act in parallel in pathology of rheumatoid arthritis. It has been speculated that the heterogeneity of RA patients with respect to their clinical response to TNF inhibitors may reflect a pathological contribution by TWEAK.

[00118] US Patent No. 7,169,387 describes the preparation of a monoclonal antibody specific for TWEAK and its use to block aspects of the development of graft-versus-host disease (GVHD) using a mouse model of chronic GVHD. US Patent Application Publication No. 2007/0280940 describes TWEAKR decoy receptors and antibodies against TWEAKR and TWEAK and their use in the treatment of central nervous system diseases associated with cerebral edema and cell death.

[00119] In some embodiments, binding domains of this disclosure comprise V<sub>H</sub> and V<sub>L</sub> domains specific for a TWEAK. In certain embodiments, the V<sub>H</sub> and V<sub>L</sub> domains are rodent (*e.g.*, mouse, rat), humanized, or human. Examples of binding domains containing such V<sub>H</sub> and V<sub>L</sub> domains specific for TWEAK, include those disclosed, for example, in US Patent No. 7,169,387. Monoclonal antibodies that block TWEAK have been shown to be effective in a mouse collagen-induced arthritis (CIA) model (Kamata *et al.* (2006) J. Immunol. 177:6433; Perper *et al.* (2006) J. Immunol. 177:2610).

[00120] In certain embodiments, a TWEAK antagonist may be an extracellular domain ("ectodomain") of a TWEAKR (also known as FN14). As used herein, a TWEAKR ectodomain refers to an extracellular portion of TWEAKR, a soluble TWEAKR, or any combination thereof. In certain embodiments, a TWEAK antagonist comprises an amino-terminal portion of TWEAKR, such as the first 70 amino acids of TWEAKR as set forth in GenBank Accession No. NP\_057723.1 (SEQ ID NO:761), or any fragment thereof that continues to function as a TWEAK antagonist. In other embodiments, a TWEAK antagonist

comprises amino acids 28-70 of SEQ ID NO:761 (*i.e.*, without the native leader sequence). In yet further embodiments, a TWEAK antagonist comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% , or at least 100% identical to an amino acid sequence of SEQ ID NO:761, or amino acids 28-70 of SEQ ID NO:761, wherein the antagonist binds to TWEAK and inhibits the activity thereof.

[00121] The ability of binding proteins or fusion proteins described herein to reduce binding of TWEAK to TWEAKR may be determined using assays known to those of skill in the art including those described in US Patent Application Publication No. 2007/0280940.

### **IGF Antagonists**

[00122] As noted above, in certain embodiments the present disclosure provides polypeptides containing a binding region or domain that is an IGF1 or IGF2 antagonist (*i.e.*, can inhibit IGF1 or IGF2 signaling). Exemplary IGF1 or IGF2 antagonists include binding domains specific for IGF1 or IGF2, such as an immunoglobulin variable binding domain or derivative thereof (*e.g.*, an antibody, Fab, scFv, or the like), or an IGF1R or IGFBP ectodomain or sub-domain thereof.

[00123] The insulin-like growth factors (IGFs), comprise a family of peptides that play important roles in mammalian growth and development. Insulin-like growth factor 1 (IGF1) is a secreted protein that has the following features: disulfide bonds (amino acids 54-96,66-109,95-100); D peptide domain (amino acids 111-118); carboxyl-terminal propeptide domain (E peptide) (amino acids 119-153); insulin chain A-like domain (amino acids 90-110); insulin chain B-like domain (amino acids 49-77); insulin connecting C peptide-like domain (amino acids 78-89); propeptide domain (amino acids 22-48); and signal sequence domain (amino acids 1-21).

[00124] IGF1 is synthesized in multiple tissues including liver, skeletal muscle, bone and cartilage. The changes in blood concentrations of IGF1 reflect changes in its synthesis and secretion from the liver, which accounts for 80% of the total serum IGF1 in experimental animals. The remainder of the IGF1 is synthesized in the periphery, usually by connective tissue cell types, such as stromal cells that are present in most tissues. IGF1 that is synthesized in the periphery can function to regulate cell growth by autocrine and paracrine mechanisms. Within these tissues, the newly synthesized and secreted IGF1 can bind to receptors that are present either on the connective tissue cells themselves and stimulate growth (autocrine), or it can bind to receptors on adjacent cell types (often epithelial cell

types) that do not actually synthesize IGF1 but are stimulated to grow by locally secreted IGF1 (paracrine) (Clemmons, 2007, Nat Rev Drug Discov. 6(10): 821-33). IGF1 synthesis is controlled by several factors, including the human pituitary growth hormone (GH, also known as somatotropin). IGF2 concentrations are high during fetal growth but are less GH-dependent in adult life compared with IGF1.

[00125] IGF1 enhances growth and/or survival of cells in a variety of tissues including musculoskeletal systems, liver, kidney, intestines, nervous system tissues, heart, and lung. IGF1 also has an important role in promoting cell growth and consequently IGF1 inhibition is being pursued as a potential adjunctive measure for treating atherosclerosis. Inhibiting IGF1 action has been proposed as a specific treatment either for potentiating the effects of other forms of anticancer therapies or for directly inhibiting tumor cell growth.

[00126] Like IGF1, IGF2 acts through IGF1R. IGF2 is an important autocrine growth factor in tumors due to its mitogenic and antiapoptotic functions (Kaneda et al., 2005, Cancer Res 65(24): 11236-11240). Increased expression of IGF2 is found frequently in a wide variety of malignancies, including colorectal, liver, esophageal and adrenocortical cancer, as well as sarcomas. Paracrine signaling by IGF2 also plays a role in tumors including breast cancers, as abundant expression of IGF2 is found in stromal fibroblasts surrounding malignant breast epithelial cells.

[00127] Insulin-like growth factor 1 receptor (IGF1R) is a tetramer of two alpha and two beta chains linked by disulfide bonds. Cleavage of a precursor generates the alpha and beta subunits. IGF1R is related to the protein kinase superfamily, the tyrosine protein kinase family, and the insulin receptor subfamily. It contains three fibronectin type-III domains, and one protein kinase domain (Lawrence et al., 2007, Current Opinion in Structural Biology 17: 699-705). The alpha chains contribute to the formation of the ligand-binding domain, while the beta chain carries a kinase domain. It is a single-pass type I membrane protein and is expressed in a variety of tissues.

[00128] The kinase domain has tyrosine-protein kinase activity, which is necessary for the activation of the IGF1- or IGF2-stimulated downstream signaling cascade. Autophosphorylation activates the kinase activity. IGF1R interacts with PIK3R1 and with the PTB/PID domains of IRS1 and SHC1 *in vitro* when autophosphorylated on tyrosine residues in the cytoplasmic domain of the beta subunit. IGF1R plays a critical role in transformation events. It is highly over-expressed in most malignant tissues where it functions as an anti-apoptotic agent by enhancing cell survival. Cells lacking this receptor cannot be transformed by most oncogenes, with the exception of v-Src.

[00129] The insulin-like growth factor-binding protein (IGFBP) family comprises six soluble proteins (IGFBP1–6) of approximately 250 residues that bind to IGFs with nanomolar affinities. Because of their sequence homology, IGFBPs are assumed to share a common overall fold and are expected to have closely related IGF-binding determinants. Each IGFBP can be divided into three distinct domains of approximately equal lengths: highly conserved cysteine-rich N and C domains and a central linker domain unique to each IGFBP species. Both the N and C domains participate in the binding to IGFs, although the specific roles of each of these domains in IGF binding have not been decisively determined. The C-terminal domain may be responsible for preferences of IGFBPs for one species of IGF over the other; the C-terminal domain is also involved in regulation of the IGF-binding affinity through interaction with extracellular matrix components and is most probably engaged in mediating IGF1-independent actions. The central linker domain is the least conserved region and has never been cited as part of the IGF-binding site for any IGFBP. This domain is the site of posttranslational modifications, specific proteolysis, and the acid-labile subunit and extracellular matrix associations known for IGFBPs. Proteolytic cleavage in this domain is believed to produce lower-affinity N- and C-terminal fragments that cannot compete with IGF receptors for IGFs, and, thus, the proteolysis is assumed to be the predominant mechanism for IGF release from IGFBPs. However, recent studies indicate that the resulting N- and C-terminal fragments still can inhibit IGF activity and have functional properties that differ from those of the intact proteins (Sitar et al. (2006) *Proc. Natl. Acad. Sci. USA.* 103(35):13028-33).

[00130] IGF-binding proteins are secreted proteins that prolong the half-life of the IGFs and have been shown to either inhibit or stimulate the growth promoting effects of the IGFs on cell culture. They alter the interaction of IGFs with their cell surface receptors and also promote cell migration. They bind equally well to IGF1 and IGF2. The C-terminal domains of all IGFBPs show sequence homology with thyroglobulin type-1 domains and share common elements of secondary structure: an  $\alpha$ -helix and a 3- to 4- $\beta$ -stranded  $\beta$ -sheet. The core of the molecule is connected by the consensus three disulfide pairings, has conserved Tyr/Phe amino acids and has the QC, CWCV motifs. These essential features are preserved in CBP1, CBP4, and CBP-6, the structures of C domains solved so far, although there are significant variations in detail. For example, CBP4 has helix  $\alpha$ 2, whereas the corresponding residues in CBP1 form a short beta-strand seen in other structures of the thyroglobulin type-1 domain superfamily. This particular region of CBPs has high sequence



diversity and is involved in the IGF complex formation and thus may perform the role of an affinity regulator.

[00131] Inhibition of IGF/IGF-receptor binding interferes with cell growth and represents a strategy for the development of IGFBPs and variants as natural IGF antagonists in many common diseases that arise from dysregulation of the IGF system, including diabetes, atherosclerosis, and cancer.

[00132] In some embodiments, binding domains of this disclosure comprise  $V_H$  and  $V_L$  domains specific for IGF1 or IGF2. In certain embodiments, the  $V_H$  and  $V_L$  domains are rodent (*e.g.*, mouse, rat), humanized, or human. Binding domains of this disclosure may also, or alternatively, comprise an IGF1R ectodomain of Genbank Accession no. NP\_000866.1 (SEQ ID NO:753) or a sub-domain thereof, or an IGFBP ectodomain of Genbank Accession no. NP\_000587.1 (IGFBP1; SEQ ID NO:754), NP\_000588.2 (IGFBP2; SEQ ID NO:755), NP\_001013416.1 (IGFBP3 isoform a; SEQ ID NO:756), NP\_000589.2 (IGFBP3 isoform b; SEQ ID NO:757), NP\_001543.2 (IGFBP4; SEQ ID NO:758), NP\_000590.1 (IGFBP5; SEQ ID NO:759) or NP\_002169.1 (IGFBP6; SEQ ID NO:760) or a sub-domain thereof. In yet further embodiments, an IGF1 or IGF2 antagonist comprises a sequence that is at least 80%, at least 85%, at least 90%, at least 91%, at least 92%, at least 93%, at least 94%, at least 95%, at least 96%, at least 97%, at least 98%, at least 99%, at least 99.5% or at least 100% identical to an amino acid sequence of SEQ ID NO:754-760, wherein the antagonist inhibits the activity of at least one or IGF1 and IGF2.

### **Multi-Specific Fusion Proteins**

[00133] The present disclosure provides multi-specific fusion proteins comprising a domain that is an antagonist of TGF $\beta$  ("TGF $\beta$  antagonist domain") and a domain that is an antagonist or agonist of a ligand other than a TGF $\beta$  ligand ("heterologous binding domain"), such as an IL6 antagonist, IL10 antagonist, GPCR agonist, VEGF antagonist, TNF antagonist, HGF antagonist, TWEAK antagonist, or IGF antagonist. It is contemplated that the TGF $\beta$  antagonist domain may be at the amino-terminus and the heterologous binding domain at the carboxy-terminus of a fusion protein, or the heterologous binding domain may be at the amino-terminus and the TGF $\beta$  antagonist may be at the carboxy-terminus. As set forth herein, the binding domains of this disclosure may be fused to each end of an intervening domain (*e.g.*, an immunoglobulin constant region or sub-region thereof). Furthermore, the two or more binding domains may be each joined to an intervening domain via a linker known in the art or as described herein.

[00134] As used herein, an “intervening domain” refers to an amino acid sequence that simply functions as a scaffold for one or more binding domains so that the fusion protein will exist primarily (*e.g.*, 50% or more of a population of fusion proteins) or substantially (*e.g.*, 90% or more of a population of fusion proteins) as a single chain polypeptide in a composition. For example, certain intervening domains can have a structural function (*e.g.*, spacing, flexibility, rigidity) or biological function (*e.g.*, an increased half-life in plasma, such as in human blood). Exemplary intervening domains that can increase half-life of the fusion proteins of this disclosure in plasma include albumin, transferrin, a scaffold domain that binds a serum protein, or the like, or fragments thereof.

[00135] In certain preferred embodiments, the intervening domain contained in a multi-specific fusion protein of this disclosure is a “dimerization domain,” which refers to an amino acid sequence that is capable of promoting the association of at least two single chain polypeptides or proteins via non-covalent or covalent interactions, such as by hydrogen bonding, electrostatic interactions, Van der Waal’s forces, disulfide bonds, hydrophobic interactions, or the like, or any combination thereof. Exemplary dimerization domains include immunoglobulin heavy chain constant regions or sub-regions. It should be understood that a dimerization domain can promote the formation of dimers or higher order multimer complexes (such as trimers, tetramers, pentamers, hexamers, septamers, octamers, *etc.*).

[00136] A “constant sub-region” is a term defined herein to refer to a peptide, polypeptide, or protein sequence that corresponds to or is derived from part or all of one or more immunoglobulin constant region domains, but does not contain all constant region domains found in a source antibody. In preferred embodiments, the constant region domains of a fusion protein of this disclosure contains a CH2 domain and a CH3 domain of IgG, IgA, or IgD, more preferably IgG1 CH2 and CH3, and even more preferably human IgG1 CH2 and CH3. In some embodiments, the constant region domains of a fusion protein of this disclosure lack or have minimal effector functions of antibody-dependent cell-mediated cytotoxicity (ADCC), antibody-dependent cell-mediated phagocytosis (ADCP), and complement activation and complement-dependent cytotoxicity (CDC), while retaining the ability to bind some F<sub>C</sub> receptors (such as F<sub>C</sub>Rn binding) and retaining a relatively long half life *in vivo*. In certain embodiments, a binding domain of this disclosure is fused to a human IgG1 constant region or sub-region, wherein the IgG1 constant region or sub-region has one or more of the following amino acids mutated: leucine at position 234 (L234), leucine at position 235 (L235), glycine at position 237 (G237), glutamate at position 318 (E318), lysine

at position 320 (K320), lysine at position 322 (K322), or any combination thereof (EU numbering).

**[00137]** Methods are known in the art for making mutations inside or outside an Fc domain that can alter Fc interactions with Fc receptors (CD16, CD32, CD64, CD89, FcεR1, FcRn) or with the complement component C1q (*see, e.g.*, US Patent No. 5,624,821; Presta (2002) *Curr. Pharma. Biotechnol.* 3:237). Particular embodiments of this disclosure include compositions comprising immunoglobulin or fusion proteins that have a constant region or sub-region from human IgG wherein binding to FcRn and protein A are preserved and wherein the Fc domain no longer interacts or minimally interacts with other Fc receptors or C1q. For example, a binding domain of this disclosure can be fused to a human IgG1 constant region or sub-region wherein the asparagine at position 297 (N297 under EU numbering) has been mutated to another amino acid to reduce or eliminate glycosylation at this site and, therefore, abrogate efficient Fc binding to FcγR and C1q. Another exemplary mutation is a P331S, which knocks out C1q binding but does not affect Fc binding.

**[00138]** In further embodiments, an immunoglobulin Fc region may have an altered glycosylation pattern relative to an immunoglobulin referent sequence. For example, any of a variety of genetic techniques may be employed to alter one or more particular amino acid residues that form a glycosylation site (*see Co et al.* (1993) *Mol. Immunol.* 30:1361; Jacquemon *et al.* (2006) *J. Thromb. Haemost.* 4:1047; Schuster *et al.* (2005) *Cancer Res.* 65:7934; Warnock *et al.* (2005) *Biotechnol. Bioeng.* 92:831). Alternatively, the host cells in which fusion proteins of this disclosure are produced may be engineered to produce an altered glycosylation pattern. One method known in the art, for example, provides altered glycosylation in the form of bisected, non-fucosylated variants that increase ADCC. The variants result from expression in a host cell containing an oligosaccharide-modifying enzyme. Alternatively, the Potelligent technology of BioWa/Kyowa Hakko is contemplated to reduce the fucose content of glycosylated molecules according to this disclosure. In one known method, a CHO host cell for recombinant immunoglobulin production is provided that modifies the glycosylation pattern of the immunoglobulin Fc region, through production of GDP-fucose.

**[00139]** Alternatively, chemical techniques are used to alter the glycosylation pattern of fusion proteins of this disclosure. For example, a variety of glycosidase and/or mannosidase inhibitors provide one or more of desired effects of increasing ADCC activity, increasing Fc receptor binding, and altering glycosylation pattern. In certain embodiment, cells expressing a multispecific fusion protein of the instant disclosure (containing a TGFβ

antagonist domain linked to a IL6, IL6R, IL6xR, IL10, VEGF, TNF, HGF, TWEAK, IGF antagonist or to a GITR agonist) are grown in a culture medium comprising a carbohydrate modifier at a concentration that increases the ADCC of immunoglycoprotein molecules produced by said host cell, wherein said carbohydrate modifier is at a concentration of less than 800  $\mu$ M. In a preferred embodiment, the cells expressing these multispecific fusion proteins are grown in a culture medium comprising castanospermine or kifunensine, more preferably castanospermine at a concentration of 100-800  $\mu$ M, such as 100  $\mu$ M, 200  $\mu$ M, 300  $\mu$ M, 400  $\mu$ M, 500  $\mu$ M, 600  $\mu$ M, 700  $\mu$ M, or 800  $\mu$ M. Methods for altering glycosylation with a carbohydrate modifier such as castanospermine are provided in US Patent Application Publication No. 2009/0041756 or PCT Publication No. WO 2008/052030.

**[00140]** In another embodiment, the immunoglobulin Fc region may have amino acid modifications that affect binding to effector cell Fc receptors. These modifications can be made using any technique known in the art, such as the approach disclosed in Presta *et al.* (2001) *Biochem. Soc. Trans.* 30:487. In another approach, the Xencor XmAb technology is available to engineer constant sub-regions corresponding to Fc domains to enhance cell killing effector function (*see* Lazar *et al.* (2006) *Proc. Nat'l. Acad. Sci. (USA)* 103:4005). Using this approach, for example, one can generate constant sub-regions with improved specificity and binding for FC $\gamma$ R, thereby enhancing cell killing effector function.

**[00141]** In still further embodiments, a constant region or sub-region can optionally increase plasma half-life or placental transfer in comparison to a corresponding fusion protein lacking such an intervening domain. In certain embodiments, the extended plasma half-life of a fusion protein of this disclosure is at least two, at least three, at least four, at least five, at least ten, at least 12, at least 18, at least 20, at least 24, at least 30, at least 36, at least 40, at least 48 hours, at least several days, at least a week, at least two weeks, at least several weeks, at least a month, at least two months, at least several months, or more in a human.

**[00142]** A constant sub-region may include part or all of any of the following domains: a C<sub>H2</sub> domain, a C<sub>H3</sub> domain (IgA, IgD, IgG, IgE, or IgM), and a C<sub>H4</sub> domain (IgE or IgM). A constant sub-region as defined herein, therefore, can refer to a polypeptide that corresponds to a portion of an immunoglobulin constant region. The constant sub-region may comprise a C<sub>H2</sub> domain and a C<sub>H3</sub> domain derived from the same, or different, immunoglobulins, antibody isotypes, or allelic variants. In some embodiments, the C<sub>H3</sub> domain is truncated and comprises a carboxy-terminal sequence listed in PCT Publication No. WO 2007/146968 as SEQ ID NO:366-371, which sequences are hereby incorporated by

reference. In certain embodiments, a constant sub-region of a polypeptide of this disclosure has a C<sub>H2</sub> domain and C<sub>H3</sub> domain, which may optionally have an amino-terminal linker, a carboxy-terminal linker, or a linker at both ends.

**[00143]** A “linker” is a peptide that joins or links other peptides or polypeptides, such as a linker of about 2 to about 150 amino acids. In fusion proteins of this disclosure, a linker can join an intervening domain (*e.g.*, an immunoglobulin-derived constant sub-region) to a binding domain or a linker can join two variable regions of a binding domain. For example, a linker can be an amino acid sequence obtained, derived, or designed from an antibody hinge region sequence, a sequence linking a binding domain to a receptor, or a sequence linking a binding domain to a cell surface transmembrane region or membrane anchor. In some embodiments, a linker can have at least one cysteine capable of participating in at least one disulfide bond under physiological conditions or other standard peptide conditions (*e.g.*, peptide purification conditions, conditions for peptide storage). In certain embodiments, a linker corresponding or similar to an immunoglobulin hinge peptide retains a cysteine that corresponds to the hinge cysteine disposed toward the amino-terminus of that hinge. In further embodiments, a linker is from an IgG1 or IgG2A hinge and has one cysteine or two cysteines corresponding to hinge cysteines. In certain embodiments, one or more disulfide bonds are formed as inter-chain disulfide bonds between intervening domains. In other embodiments, fusion proteins of this disclosure can have an intervening domain fused directly to a binding domain (*i.e.*, absent a linker or hinge). In some embodiments, the intervening domain is a dimerization domain.

**[00144]** The intervening or dimerization domain of multi-specific fusion proteins of this disclosure may be connected to one or more terminal binding domains by a peptide linker. In addition to providing a spacing function, a linker can provide flexibility or rigidity suitable for properly orienting the one or more binding domains of a fusion protein, both within the fusion protein and between or among the fusion proteins and their target(s). Further, a linker can support expression of a full-length fusion protein and stability of the purified protein both *in vitro* and *in vivo* following administration to a subject in need thereof, such as a human, and is preferably non-immunogenic or poorly immunogenic in those same subjects. In certain embodiments, a linker of an intervening or a dimerization domain of multi-specific fusion proteins of this disclosure may comprise part or all of a human immunoglobulin hinge.

**[00145]** Additionally, a binding domain may comprise a V<sub>H</sub> and a V<sub>L</sub> domain, and these variable region domains may be combined by a linker. Exemplary variable region

binding domain linkers include those belonging to the (Gly<sub>n</sub>Ser) family, such as (Gly<sub>3</sub>Ser)<sub>n</sub>(Gly<sub>4</sub>Ser)<sub>1</sub>, (Gly<sub>3</sub>Ser)<sub>1</sub>(Gly<sub>4</sub>Ser)<sub>n</sub>, (Gly<sub>3</sub>Ser)<sub>n</sub>(Gly<sub>4</sub>Ser)<sub>n</sub>, or (Gly<sub>4</sub>Ser)<sub>n</sub>, wherein n is an integer of 1 to 5 (*see, e.g.*, Linkers 22, 29, 46, 89, 90, and 116 corresponding to SEQ ID NOS:518, 525, 542, 585, 586 and 603, respectively). In preferred embodiments, these (Gly<sub>4</sub>Ser)-based linkers are used to link variable domains and are not used to link a binding domain (*e.g.*, scFv) to an intervening domain (*e.g.*, an IgG CH2CH3).

[00146] Exemplary linkers that can be used join an intervening domain (*e.g.*, an immunoglobulin-derived constant sub-region) to a binding domain or to join two variable regions of a binding domain are provided in SEQ ID NO:497-604 and 1223-1228.

[00147] Linkers contemplated in this disclosure include, for example, peptides derived from any inter-domain region of an immunoglobulin superfamily member (*e.g.*, an antibody hinge region) or a stalk region of C-type lectins, a family of type II membrane proteins. These linkers range in length from about two to about 150 amino acids, or about two to about 40 amino acids, or about eight to about 20 amino acids, preferably about ten to about 60 amino acids, more preferably about 10 to about 30 amino acids, and most preferably about 15 to about 25 amino acids. For example, Linker 1 (SEQ ID NO:497) is two amino acids in length and Linker 116 (SEQ ID NO:603) is 36 amino acids in length.

[00148] Beyond general length considerations, a linker suitable for use in the fusion proteins of this disclosure includes an antibody hinge region selected from an IgG hinge, IgA hinge, IgD hinge, IgE hinge, or variants thereof. In certain embodiments, a linker may be an antibody hinge region (upper and core region) selected from human IgG1, human IgG2, human IgG3, human IgG4, or fragments or variants thereof. As used herein, a linker that is an "immunoglobulin hinge region" refers to the amino acids found between the carboxyl end of CH1 and the amino terminal end of CH2 (for IgG, IgA, and IgD) or the amino terminal end of CH3 (for IgE and IgM). A "wild type immunoglobulin hinge region," as used herein, refers to a naturally occurring amino acid sequence interposed between and connecting the CH1 and CH2 regions (for IgG, IgA, and IgD) or interposed between and connecting the CH2 and CH3 regions (for IgE and IgM) found in the heavy chain of an antibody. In preferred embodiments, the wild type immunoglobulin hinge region sequences are human.

[00149] According to crystallographic studies, an IgG hinge domain can be functionally and structurally subdivided into three regions: the upper hinge region, the core or middle hinge region, and the lower hinge region (Shin *et al.* (1992) *Immunological Reviews* 130:87). Exemplary upper hinge regions include EPKSCDKTHT (SEQ ID NO:1240) as found in IgG1, ERKCCVE (SEQ ID NO:1241) as found in IgG2, ELKTPLGDTT HT (SEQ

ID NO:1242) or EPKSCDTPPP (SEQ ID NO:1243) as found in IgG3, and ESKYGPP (SEQ ID NO:1244) as found in IgG4. Exemplary middle hinge regions include CPPCP (SEQ ID NO:1245) as found in IgG1 and IgG2, CPRCP (SEQ ID NO:1246) as found in IgG3, and CPSCP (SEQ ID NO:1247) as found in IgG4. While IgG1, IgG2, and IgG4 antibodies each appear to have a single upper and middle hinge, IgG3 has four in tandem – one of ELKTPLGDDT HTCPRCP (SEQ ID NO:1248) and three of EPKSCDTPPP CPRCP (SEQ ID NO:1249).

[00150] IgA and IgD antibodies appear to lack an IgG-like core region, and IgD appears to have two upper hinge regions in tandem (*see* SEQ ID NOS:1250 and 1251). Exemplary wild type upper hinge regions found in IgA1 and IgA2 antibodies are set forth in SEQ ID NOS:1252 and 1253.

[00151] IgE and IgM antibodies, in contrast, instead of a typical hinge region have a CH2 region with hinge-like properties. Exemplary wild-type CH2 upper hinge-like sequences of IgE and IgM are set forth in SEQ ID NO:1254 (VCSRDFTPPT VKILQSSSDG GGHPPTIQL LCLVSGYTPG TINITWLEDG QVMDVDLSTA STTQEGELAS TQSELTLSQK HWLSDRTYTC QVTYQGHTFE DSTKKCA) and SEQ ID NO:1255 (VIAELPPKVS VFVPPRDGFF GNP RKSKLIC QATGFSPRQI QVSWLREGKQ VGSGVTTDQV QAEAKESGPT TYKVTSTLTI KESDWLGQSM FTCRVDHRGL TFQQNASSMC VP), respectively.

[00152] An “altered wild type immunoglobulin hinge region” or “altered immunoglobulin hinge region” refers to (a) a wild type immunoglobulin hinge region with up to 30% amino acid changes (*e.g.*, up to 25%, 20%, 15%, 10%, or 5% amino acid substitutions or deletions), (b) a portion of a wild type immunoglobulin hinge region that is at least 10 amino acids (*e.g.*, at least 12, 13, 14 or 15 amino acids) in length with up to 30% amino acid changes (*e.g.*, up to 25%, 20%, 15%, 10%, or 5% amino acid substitutions or deletions), or (c) a portion of a wild type immunoglobulin hinge region that comprises the core hinge region (which portion may be 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, or 15, or at least 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, or 15 amino acids in length). In certain embodiments, one or more cysteine residues in a wild type immunoglobulin hinge region may be substituted by one or more other amino acid residues (*e.g.*, one or more serine residues). An altered immunoglobulin hinge region may alternatively or additionally have a proline residue of a wild type immunoglobulin hinge region substituted by another amino acid residue (*e.g.*, a serine residue).

[00153] Alternative hinge and linker sequences that can be used as connecting regions may be crafted from portions of cell surface receptors that connect IgV-like or IgC-like domains. Regions between IgV-like domains where the cell surface receptor contains multiple IgV-like domains in tandem and between IgC-like domains where the cell surface receptor contains multiple tandem IgC-like regions could also be used as connecting regions or linker peptides. In certain embodiments, hinge and linker sequences are from five to 60 amino acids long, and may be primarily flexible, but may also provide more rigid characteristics, and may contain primarily an  $\alpha$ -helical structure with minimal  $\beta$ -sheet structure. Preferably, sequences are stable in plasma and serum and are resistant to proteolytic cleavage. In some embodiments, sequences may contain a naturally occurring or added motif such as CPPC that confers the capacity to form a disulfide bond or multiple disulfide bonds to stabilize the C-terminus of the molecule. In other embodiments, sequences may contain one or more glycosylation sites. Examples of hinge and linker sequences include interdomain regions between the IgV-like and IgC-like or between the IgC-like or IgV-like domains of CD2, CD4, CD22, CD33, CD48, CD58, CD66, CD80, CD86, CD96, CD150, CD166, and CD244. Alternative hinges may also be crafted from disulfide-containing regions of Type II receptors from non-immunoglobulin superfamily members such as CD69, CD72, and CD161.

[00154] In some embodiments, a hinge linker has a single cysteine residue for formation of an interchain disulfide bond. In other embodiments, a linker has two cysteine residues for formation of interchain disulfide bonds. In further embodiments, a hinge linker is derived from an immunoglobulin interdomain region (*e.g.*, an antibody hinge region comprising an upper and core sequence of, for example, an IgG1 hinge) or a Type II C-type lectin stalk region (derived from a Type II membrane protein; *see, e.g.*, exemplary lectin stalk region sequences set forth in of PCT Application Publication No. WO 2007/146968, such as SEQ ID NOS:111, 113, 115, 117, 119, 121, 123, 125, 127, 129, 131, 133, 135, 149, 151, 153, 155, 157, 159, 161, 163, 165, 167, 169, 231, 233, 235, 237, 239, 241, 243, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 287, 289, 297, 305, 307, 309-311, 313-331, 346, 373-377, 380, or 381 from that publication), which sequences are herein incorporated by reference.

[00155] In one aspect, exemplary multi-specific fusion proteins containing a TGF $\beta$  antagonist as described herein will also contain at least one additional binding region or domain that is specific for a target other than TGF $\beta$ , such as an IL6, IL10, VEGF, TNF,



HGF, TWEAK, IGF antagonist or a GITR agonist. For example, a multi-specific fusion protein of this disclosure has a TGF $\beta$  antagonist domain linked to an IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF antagonist or a GITR agonist domain by an intervening domain (such as a human IgG1 CH2CH3 Fc region). In certain embodiments, a multi-specific fusion protein comprises a first and second binding domain, a first and second linker, and an intervening domain, wherein one end of the intervening domain is fused via the first linker to a first binding domain that is a TGF $\beta$  antagonist (*e.g.*, a TGF $\beta$ R2 ectodomain, an anti-TGF $\beta$ R2 ectodomain, an anti-TGF $\beta$ ) and at the other end is fused via the second linker to a different binding domain that is an IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF antagonist or a GITR agonist.

**[00156]** In certain embodiments, the first linker and second linker of a multi-specific fusion protein of this disclosure are each independently selected from, for example, SEQ ID NO:497-604 and 1223-1228. For example, the first or second linker can be Linker 102 (SEQ ID NO:589), 47 (SEQ ID NO:543), 80 (SEQ ID NO:576), or any combination thereof. In further examples, one linker is Linker 102 (SEQ ID NO:589) and the other linker is Linker 47 (SEQ ID NO:543), or one linker is Linker 102 (SEQ ID NO:589) and the other linker is Linker 80 (SEQ ID NO:576). In further examples, binding domains of this disclosure that comprise V<sub>H</sub> and V<sub>L</sub> domains, such as those specific for IL6, IL6R, IL6xR, IL10, VEGF, TNF, HGF, TWEAK, IGF, GITR, TGF $\beta$ R2 ectodomain, or TGF $\beta$ , can have a further (third) linker between the V<sub>H</sub> and V<sub>L</sub> domains, such as Linker 46 (SEQ ID NO:542). In any of these embodiments, the linkers may be flanked by one to five additional junction amino acids, which may simply be a result of creating such a recombinant molecule (*e.g.*, use of a particular restriction enzyme site to join nucleic acid molecules may result in the insertion of one to several amino acids), or for purposes of this disclosure may be considered a part of any particular linker core sequence.

**[00157]** In further embodiments, the intervening domain of a multi-specific fusion protein of this disclosure is comprised of an immunoglobulin constant region or sub-region (preferably CH2CH3 of IgG, IgA, or IgD; or CH3CH4 of IgE or IgM), wherein the intervening domain is disposed between a TGF $\beta$  antagonist domain and an IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF antagonist binding domain or a GITR agonist binding domain. In certain embodiments, the intervening domain of a multi-specific fusion protein of this disclosure has a TGF $\beta$  antagonist at the amino-terminus and a binding domain specific for an IL6, IL6xR, IL10, VEGF, TNF, HGF, TWEAK, IGF, or GITR at the carboxy-terminus. In other embodiments, the intervening domain of a multi-specific fusion protein of this

disclosure has a binding domain specific for an IL6, IL10, VEGF, TNF, HGF, TWEAK, IGF antagonist binding domain or a GITR agonist binding domain at the amino-terminus and a TGF $\beta$  antagonist at the carboxy-terminus. In further embodiments, the immunoglobulin constant region sub-region includes CH2 and CH3 domains of immunoglobulin G1 (IgG1). In related embodiments, the IgG1 CH2 and CH3 domains have one or more of the following amino acids mutated (*i.e.*, have a different amino acid at that position): leucine at position 234 (L234), leucine at position 235 (L235), glycine at position 237 (G237), glutamate at position 318 (E318), lysine at position 320 (K320), lysine at position 322 (K322), or any combination thereof (EU numbering). For example, any one of these amino acids can be changed to alanine. In a further embodiment, according to Kabat numbering, the CH2 domain has each of L234, L235, G237, E318, K320 and K322 mutated to an alanine (*i.e.*, L234A, L235A, G237A, E318A, K320A and K322A, respectively).

**[00158]** In some embodiments, a multi-specific fusion protein of this disclosure has a TGF $\beta$  antagonist that comprises a TGF $\beta$ R2 ectodomain or a sub-domain of a TGF $\beta$ R2 ectodomain, or any combination thereof. For example, a TGF $\beta$  antagonist can comprise amino acids 73-176 as set forth in GenBank Accession No. NP\_001020018.1, amino acids 48-151 as set forth in GenBank Accession No. NP\_003233.4, or any combination thereof. In further embodiments, the TGF $\beta$  antagonist comprises an amino acid sequence as set forth in SEQ ID NO:743 or 744.

**[00159]** In further embodiments, a multi-specific fusion protein of this disclosure having a TGF $\beta$  antagonist of this disclosure also has an IL6 antagonist binding domain that binds with higher affinity to IL6xR than to either IL6 or IL6R $\alpha$  alone and competes with sIL6xR complex binding to mgp130 or enhances sgp103 binding to sIL6xR complex. In certain embodiments, a binding domain specific for an IL6xR comprises (i) a V<sub>H</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% identical to the amino acid sequence of a V<sub>H</sub> domain found in any one of SEQ ID NOS:435-496; or (ii) a V<sub>L</sub> domain having an amino acid sequence that is at least 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or 100% identical to the amino acid sequence of a V<sub>L</sub> domain found in any one of SEQ ID NOS:373-434; or (iii) both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii); or both a V<sub>H</sub> domain of (i) and a V<sub>L</sub> domain of (ii) wherein the V<sub>H</sub> and V<sub>L</sub> are from the same reference sequence. In one embodiment, such V<sub>H</sub> and V<sub>L</sub> domains can form exemplary binding domain TRU6-1019 (*see* SEQ ID NOS:453 and 391, respectively).

**[00160]** In still further embodiments, an IL6 antagonist binding domain, which binds to the IL6xR with a higher affinity than IL6 or IL6R $\alpha$  or either IL6 or IL6R $\alpha$  alone, and competes with gp130 for binding to the sIL6xR complex or enhances sgp130 binding to sIL6xR complex, comprises V<sub>H</sub> and V<sub>L</sub> domains comprising framework regions and CDR1, CDR2 and CDR3 regions, wherein (a) the V<sub>H</sub> domain comprises the amino acid sequence of a heavy chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:435-496; or (b) the V<sub>L</sub> domain comprises the amino acid sequence of a light chain CDR1, CDR2, and CDR3 found in any one of SEQ ID NOS:373-434; or (c) the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b); or the binding domain comprises a V<sub>H</sub> amino acid sequence of (a) and a V<sub>L</sub> amino acid sequence of (b), wherein the V<sub>H</sub> and V<sub>L</sub> amino acid sequences are from the same reference sequence. The V<sub>L</sub> and V<sub>H</sub> domains of these multi-specific fusion proteins may be arranged in either orientation and may be separated by up to about a 5-30 amino acid linker as disclosed herein. In certain embodiments, a linker joining the V<sub>H</sub> and V<sub>L</sub> domains comprises an amino acid sequence of Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576). In certain embodiments, a multi-specific fusion protein comprising the IL6 antagonist binding domain measurably inhibits IL6 cis- and trans-signaling, preferably trans-signaling and, optionally, does not inhibit signaling of gp130 family cytokines other than IL6.

**[00161]** Exemplary structures of such multi-specific fusion proteins, referred to herein as Xceptor molecules, include N-BD-X-ED-C, N-ED-X-BD-C, N-ED1-X-ED2-C, wherein BD is an immunoglobulin-like or immunoglobulin variable region binding domain, X is an intervening domain, and ED is a receptor ectodomain, or the like. In some constructs, X can comprise an immunoglobulin constant region or sub-region disposed between the first and second binding domains. In some embodiments, a multi-specific fusion protein of this disclosure has an intervening domain (X) comprising, from amino-terminus to carboxy-terminus, a structure as follows: -L1-X-L2-, wherein L1 and L2 are each independently a linker comprising from two to about 150 amino acids; and X is an immunoglobulin constant region or sub-region. In further embodiments, the multi-specific fusion protein will have an intervening domain that is albumin, transferrin, or another serum protein binding protein, wherein the fusion protein remains primarily or substantially as a single chain polypeptide in a composition. In still further embodiments, a multi-specific fusion protein of this disclosure has the following structure: N-BD1-X-L2-BD2-C, wherein N and C represent the amino-terminus and carboxy-terminus, respectively; BD1 is a TGF $\beta$  antagonist that is at least about

90% identical to an ectodomain of TGF $\beta$ R2; -X- is -L1-CH<sub>2</sub>CH<sub>3</sub>-, wherein L1 is the first IgG1 hinge, optionally mutated by substituting the first cysteine and wherein -CH<sub>2</sub>CH<sub>3</sub>- is the CH<sub>2</sub>CH<sub>3</sub> region of an IgG1 Fc domain, optionally mutated to eliminate Fc $\gamma$ RI-III interaction while retaining FcRn interaction; L2 is a linker selected from SEQ ID NO:497-604 and 1223-1228; and BD2 is a binding domain specific for an IL6 or IL6/IL6R complex.

**[00162]** In particular embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) an IL6 antagonist comprising a heavy chain variable region with CDR1, CD2, and CDR3 amino acid sequences at least 80% to 100% identical to sequences set forth in SEQ ID NOS:435-496, respectively, and a light chain variable region with CDR1, CDR2, and CDR3 amino acid sequences at least 80% to 100% identical to sequences set forth in SEQ ID NOS:373-434, respectively, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or an IL6 antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH<sub>2</sub> and CH<sub>3</sub> comprising amino acids 276 to 489 of SEQ ID NO:625, (iii) the CH<sub>2</sub>CH<sub>3</sub> constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or an IL6 antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), the second linker is Linker 102 (SEQ ID NO:589), and a further (third) linker between the IL6 antagonist V<sub>H</sub> and V<sub>L</sub> domains is Linker 46 (SEQ ID NO:542).

**[00163]** In other embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) an IL10 antagonist comprising an amino acid sequence at least 80% to 100% identical to an amino acid sequence of SEQ ID NO:745 or to amino acids 22-401 of SEQ ID NO:745, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or an IL10 antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH<sub>2</sub> and CH<sub>3</sub>, (iii) the CH<sub>2</sub>CH<sub>3</sub> constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or an IL10 antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), and the second linker is Linker 102 (SEQ ID NO:589).

**[00164]** In further embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) a VEGF antagonist comprising an amino acid sequence at least 80% to 100% identical to an amino acid sequence of SEQ ID NO:747, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or a VEGF antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH2 and CH3, (iii) the CH2CH3 constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or a VEGF antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576) and the second linker is Linker 102 (SEQ ID NO:589).

**[00165]** In further embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) a TNF $\alpha$  antagonist comprising an amino acid sequence at least 80% to 100% identical to an amino acid sequence of SEQ ID NO:748 or 749, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or a TNF $\alpha$  antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH2 and CH3, (iii) the CH2CH3 constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or a TNF $\alpha$  antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), and the second linker is Linker 102 (SEQ ID NO:589). In specific embodiments, the multi-specific Xceptor fusion protein has an amino acid sequence of SEQ ID NO:1236.

**[00166]** In further embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) a HGF antagonist comprising a heavy chain variable region with CDR1, CD2, and CDR3 amino acid sequences at least 80% to 100% identical to sequences set forth in SEQ ID NOS:921-1078, respectively, and a light chain variable region with CDR1, CDR2, and CDR3 amino acid sequences at least 80% to 100% identical to sequences set forth in SEQ ID NOS:762-920, respectively, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or a HGF antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH2 and CH3, (iii) the

CH<sub>2</sub>CH<sub>3</sub> constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or a HGF antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), the second linker is Linker 102 (SEQ ID NO:589), and a further (third) linker between the HGF antagonist V<sub>H</sub> and V<sub>L</sub> domains is Linker 46 (SEQ ID NO:542).

**[00167]** In yet other embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) a TWEAK antagonist comprising an amino acid sequence at least 80% to 100% identical to an amino acid sequence of SEQ ID NO:761, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or a TWEAK antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH<sub>2</sub> and CH<sub>3</sub>, (iii) the CH<sub>2</sub>CH<sub>3</sub> constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or a TWEAK antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), and the second linker is Linker 102 (SEQ ID NO:589). In specific embodiments, the multi-specific Xceptor fusion protein has an amino acid sequence of SEQ ID NO:1237.

**[00168]** In yet other embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) an IGF antagonist comprising an amino acid sequence at least 80% to 100% identical to an amino acid sequence of SEQ ID NO:754-760, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-terminus, (i) a TGF $\beta$  antagonist of (a) or an IGF antagonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH<sub>2</sub> and CH<sub>3</sub>, (iii) the CH<sub>2</sub>CH<sub>3</sub> constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or an IGF antagonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), and the second linker is Linker 102 (SEQ ID NO:589).

**[00169]** In other embodiments, a multi-specific Xceptor fusion protein has (a) a TGF $\beta$  antagonist comprising an amino acid sequence at least 80% to 100% identical to a sequence as set forth in SEQ ID NO:743 or 744 and (b) a GITR agonist comprising an amino acid sequence at least 80% to 100% identical to amino acids 74-181 of SEQ ID NO:746, wherein, from amino-terminus to carboxy-terminus or from carboxy-terminus to amino-

terminus, (i) a TGF $\beta$  antagonist of (a) or a GITR agonist of (b) is fused to a first linker, (ii) the first linker is fused to an immunoglobulin heavy chain constant region of CH2 and CH3, (iii) the CH2CH3 constant region polypeptide is fused to a second linker, and (iv) the second linker is fused to a TGF $\beta$  antagonist of (a) or a GITR agonist of (b). In certain embodiments, the first linker is Linker 47 (SEQ ID NO:543) or Linker 80 (SEQ ID NO:576), and the second linker is Linker 102 (SEQ ID NO:589).

### **Making Multi-Specific Fusion Proteins**

[00170] To efficiently produce any of the binding domain polypeptides or fusion proteins described herein, a leader peptide is used to facilitate secretion of expressed polypeptides and fusion proteins. Using any of the conventional leader peptides (signal sequences) is expected to direct nascently expressed polypeptides or fusion proteins into a secretory pathway and to result in cleavage of the leader peptide from the mature polypeptide or fusion protein at or near the junction between the leader peptide and the polypeptide or fusion protein. A particular leader peptide will be chosen based on considerations known in the art, such as using sequences encoded by polynucleotides that allow the easy inclusion of restriction endonuclease cleavage sites at the beginning or end of the coding sequence for the leader peptide to facilitate molecular engineering, provided that such introduced sequences specify amino acids that either do not interfere unacceptably with any desired processing of the leader peptide from the nascently expressed protein or do not interfere unacceptably with any desired function of a polypeptide or fusion protein molecule if the leader peptide is not cleaved during maturation of the polypeptides or fusion proteins. Exemplary leader peptides of this disclosure include natural leader sequences (*i.e.*, those expressed with the native protein) or use of heterologous leader sequences, such as

H<sub>3</sub>N-MDFQVQIFSFLISASVIMSRG(X)<sub>n</sub>-CO<sub>2</sub>H, wherein X is any amino acid and n is zero to three (SEQ ID NO:1185) or H<sub>3</sub>N-MEAPAQLLFLLLLWLPDTTG-CO<sub>2</sub>H (SEQ ID NO:1186).

[00171] As noted herein, variants and derivatives of binding domains, such as ectodomains, light and heavy variable regions, and CDRs described herein, are contemplated. In one example, insertion variants are provided wherein one or more amino acid residues supplement a specific binding agent amino acid sequence. Insertions may be located at either or both termini of the protein, or may be positioned within internal regions of the specific binding agent amino acid sequence. Variant products of this disclosure also include mature specific binding agent products, *i.e.*, specific binding agent products wherein a leader or

signal sequence is removed, and the resulting protein having additional amino terminal residues. The additional amino terminal residues may be derived from another protein, or may include one or more residues that are not identifiable as being derived from a specific protein. Polypeptides with an additional methionine residue at position -1 are contemplated, as are polypeptides of this disclosure with additional methionine and lysine residues at positions -2 and -1. Variants having additional Met, Met-Lys, or Lys residues (or one or more basic residues in general) are particularly useful for enhanced recombinant protein production in bacterial host cells.

**[00172]** As used herein, "amino acids" refer to a natural (those occurring in nature) amino acid, a substituted natural amino acid, a non-natural amino acid, a substituted non-natural amino acid, or any combination thereof. The designations for natural amino acids are herein set forth as either the standard one- or three-letter code. Natural polar amino acids include asparagine (Asp or N) and glutamine (Gln or Q); as well as basic amino acids such as arginine (Arg or R), lysine (Lys or K), histidine (His or H), and derivatives thereof; and acidic amino acids such as aspartic acid (Asp or D) and glutamic acid (Glu or E), and derivatives thereof. Natural hydrophobic amino acids include tryptophan (Trp or W), phenylalanine (Phe or F), isoleucine (Ile or I), leucine (Leu or L), methionine (Met or M), valine (Val or V), and derivatives thereof; as well as other non-polar amino acids such as glycine (Gly or G), alanine (Ala or A), proline (Pro or P), and derivatives thereof. Natural amino acids of intermediate polarity include serine (Ser or S), threonine (Thr or T), tyrosine (Tyr or Y), cysteine (Cys or C), and derivatives thereof. Unless specified otherwise, any amino acid described herein may be in either the D- or L-configuration.

**[00173]** Substitution variants include those fusion proteins wherein one or more amino acid residues in an amino acid sequence are removed and replaced with alternative residues. In some embodiments, the substitutions are conservative in nature; however, this disclosure embraces substitutions that are also non-conservative. Amino acids can be classified according to physical properties and contribution to secondary and tertiary protein structure. A conservative substitution is recognized in the art as a substitution of one amino acid for another amino acid that has similar properties. Exemplary conservative substitutions are set out in Table 1 (*see* WO 97/09433, page 10, published March 13, 1997), immediately below.



**Table 1. Conservative Substitutions I**

Side Chain	Characteristic	Amino Acid
Aliphatic	Non-polar	G, A, P, I, L, V
	Polar – uncharged	S, T, M, N, Q
	Polar – charged	D, E, K, R
Aromatic		H, F, W, Y
Other		N, Q, D, E

[00174] Alternatively, conservative amino acids can be grouped as described in Lehninger (Biochemistry, Second Edition; Worth Publishers, Inc. NY:NY (1975), pp.71-77) as set out in Table 2, immediately below.

**Table 2. Conservative Substitutions II**

Side Chain	Characteristic	Amino Acid
Non-polar (hydrophobic)	Aliphatic:	A, L, I, V, P
	Aromatic	F, W
	Sulfur-containing	M
	Borderline	G
Uncharged-polar	Hydroxyl	S, T, Y
	Amides	N, Q
	Sulfhydryl	C
	Borderline	G
Positively Charged (Basic)		K, R, H
Negatively Charged (Acidic)		D, E

[00175] Variants or derivatives can also have additional amino acid residues which arise from use of specific expression systems. For example, use of commercially available vectors that express a desired polypeptide as part of a glutathione-S-transferase (GST) fusion product provides the desired polypeptide having an additional glycine residue at position -1 after cleavage of the GST component from the desired polypeptide. Variants which result from expression in other vector systems are also contemplated, including those wherein histidine tags are incorporated into the amino acid sequence, generally at the carboxy and/or amino terminus of the sequence.

[00176] Deletion variants are also contemplated wherein one or more amino acid residues in a binding domain of this disclosure are removed. Deletions can be effected at one

or both termini of the fusion protein, or from removal of one or more residues within the amino acid sequence.

[00177] In certain illustrative embodiments, fusion proteins of this disclosure are glycosylated, the pattern of glycosylation being dependent upon a variety of factors including the host cell in which the protein is expressed (if prepared in recombinant host cells) and the culture conditions.

[00178] This disclosure also provides derivatives of fusion proteins. Derivatives include specific binding domain polypeptides bearing modifications other than insertion, deletion, or substitution of amino acid residues. In certain embodiments, the modifications are covalent in nature, and include for example, chemical bonding with polymers, lipids, other organic, and inorganic moieties. Derivatives of this disclosure may be prepared to increase circulating half-life of a specific binding domain polypeptide, or may be designed to improve targeting capacity for the polypeptide to desired cells, tissues, or organs.

[00179] This disclosure further embraces fusion proteins that are covalently modified or derivatized to include one or more water-soluble polymer attachments such as polyethylene glycol, polyoxyethylene glycol, or polypropylene glycol, as described U.S. Patent NOS: 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192 and 4,179,337. Still other useful polymers known in the art include monomethoxy-polyethylene glycol, dextran, cellulose, and other carbohydrate-based polymers, poly-(N-vinyl pyrrolidone)-polyethylene glycol, propylene glycol homopolymers, a polypropylene oxide/ethylene oxide co-polymer, polyoxyethylated polyols (e.g., glycerol) and polyvinyl alcohol, as well as mixtures of these polymers. Particularly preferred are polyethylene glycol (PEG)-derivatized proteins. Water-soluble polymers may be bonded at specific positions, for example at the amino terminus of the proteins and polypeptides according to this disclosure, or randomly attached to one or more side chains of the polypeptide. The use of PEG for improving therapeutic capacities is described in US Patent No. 6,133,426.

[00180] A particular embodiment of this disclosure is an immunoglobulin or an Fc fusion protein. Such a fusion protein can have a long half-life, *e.g.*, several hours, a day or more, or even a week or more, especially if the Fc domain is capable of interacting with FcRn, the neonatal Fc receptor. The binding site for FcRn in an Fc domain is also the site at which the bacterial proteins A and G bind. The tight binding between these proteins can be used as a means to purify antibodies or fusion proteins of this disclosure by, for example, employing protein A or protein G affinity chromatography during protein purification.

**[00181]** Protein purification techniques are well known to those of skill in the art. These techniques involve, at one level, the crude fractionation of the polypeptide and non-polypeptide fractions. Further purification using chromatographic and electrophoretic techniques to achieve partial or complete purification (or purification to homogeneity) is frequently desired. Analytical methods particularly suited to the preparation of a pure fusion protein are ion-exchange chromatography; exclusion chromatography; polyacrylamide gel electrophoresis; and isoelectric focusing. Particularly efficient methods of purifying peptides are fast protein liquid chromatography and HPLC.

**[00182]** Certain aspects of the present disclosure concern the purification, and in particular embodiments, the substantial purification, of a fusion protein. The term "purified fusion protein" as used herein, is intended to refer to a composition, isolatable from other components, wherein the fusion protein is purified to any degree relative to its naturally obtainable state. A purified fusion protein therefore also refers to a fusion protein, free from the environment in which it may naturally occur.

**[00183]** Generally, "purified" will refer to a fusion protein composition that has been subjected to fractionation to remove various other components, and which composition substantially retains its expressed biological activity. Where the term "substantially purified" is used, this designation refers to a fusion binding protein composition in which the fusion protein forms the major component of the composition, such as constituting about 50%, about 60%, about 70%, about 80%, about 90%, about 95%, about 99% or more of the protein, by weight, in the composition.

**[00184]** Various methods for quantifying the degree of purification are known to those of skill in the art in light of the present disclosure. These include, for example, determining the specific binding activity of an active fraction, or assessing the amount of fusion protein in a fraction by SDS/PAGE analysis. A preferred method for assessing the purity of a protein fraction is to calculate the binding activity of the fraction, to compare it to the binding activity of the initial extract, and to thus calculate the degree of purification, herein assessed by a "-fold purification number." The actual units used to represent the amount of binding activity will, of course, be dependent upon the particular assay technique chosen to follow the purification and whether or not the expressed fusion protein exhibits a detectable binding activity.

**[00185]** Various techniques suitable for use in protein purification are well known to those of skill in the art. These include, for example, precipitation with ammonium sulfate, PEG, antibodies and the like, or by heat denaturation, followed by centrifugation;

chromatography steps such as ion exchange, gel filtration, reverse phase, hydroxylapatite, and affinity chromatography; isoelectric focusing; gel electrophoresis; and combinations of these and other techniques. As is generally known in the art, it is believed that the order of conducting the various purification steps may be changed, or that certain steps may be omitted, and still result in a suitable method for the preparation of a substantially purified protein.

[00186] There is no general requirement that the fusion protein always be provided in its most purified state. Indeed, it is contemplated that less substantially purified proteins will have utility in certain embodiments. Partial purification may be accomplished by using fewer purification steps in combination, or by utilizing different forms of the same general purification scheme. For example, it is appreciated that a cation-exchange column chromatography performed utilizing an HPLC apparatus will generally result in greater purification than the same technique utilizing a low pressure chromatography system. Methods exhibiting a lower degree of relative purification may have advantages in total recovery of protein product, or in maintaining binding activity of an expressed protein.

[00187] It is known that the migration of a polypeptide can vary, sometimes significantly, with different conditions of SDS/PAGE (Capaldi et al. (1977) Biochem. Biophys. Res. Comm. 76:425). It will therefore be appreciated that under differing electrophoresis conditions, the apparent molecular weights of purified or partially purified fusion protein expression products may vary.

### **Polynucleotides, Expression Vectors, and Host Cells**

[00188] This disclosure provides polynucleotides (isolated or purified or pure polynucleotides) encoding the multi-specific fusion protein of this disclosure, vectors (including cloning vectors and expression vectors) comprising such polynucleotides, and cells (*e.g.*, host cells) transformed or transfected with a polynucleotide or vector according to this disclosure.

[00189] In certain embodiments, a polynucleotide (DNA or RNA) encoding a binding domain of this disclosure, or a multi-specific fusion protein containing one or more such binding domains is contemplated. Expression cassettes encoding multi-specific fusion protein constructs are provided in the examples appended hereto.

[00190] The present disclosure also relates to vectors that include a polynucleotide of this disclosure and, in particular, to recombinant expression constructs. In one embodiment, this disclosure contemplates a vector comprising a polynucleotide encoding a multi-specific

fusion protein containing a TGF $\beta$  antagonist domain and an IL6 or IL6/IL6R binding domain of this disclosure, along with other polynucleotide sequences that cause or facilitate transcription, translation, and processing of such multi-specific fusion protein-encoding sequences.

[00191] Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described, for example, in Sambrook *et al.*, Molecular Cloning: A Laboratory Manual, Second Edition, Cold Spring Harbor, NY, (1989). Exemplary cloning/expression vectors include cloning vectors, shuttle vectors, and expression constructs, that may be based on plasmids, phagemids, phasmids, cosmids, viruses, artificial chromosomes, or any nucleic acid vehicle known in the art suitable for amplification, transfer, and/or expression of a polynucleotide contained therein

[00192] As used herein, "vector" means a nucleic acid molecule capable of transporting another nucleic acid to which it has been linked. Exemplary vectors include plasmids, yeast artificial chromosomes, and viral genomes. Certain vectors can autonomously replicate in a host cell, while other vectors can be integrated into the genome of a host cell and thereby are replicated with the host genome. In addition, certain vectors are referred to herein as "recombinant expression vectors" (or simply, "expression vectors"), which contain nucleic acid sequences that are operatively linked to an expression control sequence and, therefore, are capable of directing the expression of those sequences.

[00193] In certain embodiments, expression constructs are derived from plasmid vectors. Illustrative constructs include modified pNASS vector (Clontech, Palo Alto, CA), which has nucleic acid sequences encoding an ampicillin resistance gene, a polyadenylation signal and a T7 promoter site; pDEF38 and pNEF38 (CMC ICOS Biologics, Inc.), which have a CHEF1 promoter; and pD18 (Lonza), which has a CMV promoter. Other suitable mammalian expression vectors are well known (*see, e.g.*, Ausubel *et al.*, 1995; Sambrook *et al.*, *supra*; *see also, e.g.*, catalogs from Invitrogen, San Diego, CA; Novagen, Madison, WI; Pharmacia, Piscataway, NJ). Useful constructs may be prepared that include a dihydrofolate reductase (DHFR)-encoding sequence under suitable regulatory control, for promoting enhanced production levels of the fusion proteins, which levels result from gene amplification following application of an appropriate selection agent (*e.g.*, methotrexate).

[00194] Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, and a promoter derived from a highly-expressed gene to direct transcription of a downstream structural sequence, as

described above. A vector in operable linkage with a polynucleotide according to this disclosure yields a cloning or expression construct. Exemplary cloning/expression constructs contain at least one expression control element, *e.g.*, a promoter, operably linked to a polynucleotide of this disclosure. Additional expression control elements, such as enhancers, factor-specific binding sites, terminators, and ribosome binding sites are also contemplated in the vectors and cloning/expression constructs according to this disclosure. The heterologous structural sequence of the polynucleotide according to this disclosure is assembled in appropriate phase with translation initiation and termination sequences. Thus, for example, the fusion protein-encoding nucleic acids as provided herein may be included in any one of a variety of expression vector constructs as a recombinant expression construct for expressing such a protein in a host cell.

**[00195]** The appropriate DNA sequence(s) may be inserted into a vector, for example, by a variety of procedures. In general, a DNA sequence is inserted into an appropriate restriction endonuclease cleavage site(s) by procedures known in the art. Standard techniques for cloning, DNA isolation, amplification and purification, for enzymatic reactions involving DNA ligase, DNA polymerase, restriction endonucleases and the like, and various separation techniques are contemplated. A number of standard techniques are described, for example, in Ausubel *et al.* (*Current Protocols in Molecular Biology*, Greene Publ. Assoc. Inc. & John Wiley & Sons, Inc., Boston, MA, 1993); Sambrook *et al.* (*Molecular Cloning*, Second Ed., Cold Spring Harbor Laboratory, Plainview, NY, 1989); Maniatis *et al.* (*Molecular Cloning*, Cold Spring Harbor Laboratory, Plainview, NY, 1982); Glover (Ed.) (*DNA Cloning* Vol. I and II, IRL Press, Oxford, UK, 1985); Hames and Higgins (Eds.) (*Nucleic Acid Hybridization*, IRL Press, Oxford, UK, 1985); and elsewhere.

**[00196]** The DNA sequence in the expression vector is operatively linked to at least one appropriate expression control sequence (*e.g.*, a constitutive promoter or a regulated promoter) to direct mRNA synthesis. Representative examples of such expression control sequences include promoters of eukaryotic cells or their viruses, as described above. Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein-I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art, and preparation of certain particularly preferred recombinant expression constructs comprising at least one promoter or regulated promoter

operably linked to a nucleic acid encoding a protein or polypeptide according to this disclosure is described herein.

**[00197]** Variants of the polynucleotides of this disclosure are also contemplated. Variant polynucleotides are at least 90%, and preferably 95%, 99%, or 99.9% identical to one of the polynucleotides of defined sequence as described herein, or that hybridizes to one of those polynucleotides of defined sequence under stringent hybridization conditions of 0.015M sodium chloride, 0.0015M sodium citrate at about 65-68°C or 0.015M sodium chloride, 0.0015M sodium citrate, and 50% formamide at about 42°C. The polynucleotide variants retain the capacity to encode a binding domain or fusion protein thereof having the functionality described herein.

**[00198]** The term “stringent” is used to refer to conditions that are commonly understood in the art as stringent. Hybridization stringency is principally determined by temperature, ionic strength, and the concentration of denaturing agents such as formamide. Examples of stringent conditions for hybridization and washing are 0.015M sodium chloride, 0.0015M sodium citrate at about 65-68°C or 0.015M sodium chloride, 0.0015M sodium citrate, and 50% formamide at about 42°C (*see* Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual*, 2nd Ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, N.Y., 1989).

**[00199]** More stringent conditions (such as higher temperature, lower ionic strength, higher formamide, or other denaturing agent) may also be used; however, the rate of hybridization will be affected. In instances wherein hybridization of deoxyoligonucleotides is concerned, additional exemplary stringent hybridization conditions include washing in 6x SSC, 0.05% sodium pyrophosphate at 37°C (for 14-base oligonucleotides), 48°C (for 17-base oligonucleotides), 55°C (for 20-base oligonucleotides), and 60°C (for 23-base oligonucleotides).

**[00200]** A further aspect of this disclosure provides a host cell transformed or transfected with, or otherwise containing, any of the polynucleotides or vector/expression constructs of this disclosure. The polynucleotides or cloning/expression constructs of this disclosure are introduced into suitable cells using any method known in the art, including transformation, transfection and transduction. Host cells include the cells of a subject undergoing *ex vivo* cell therapy including, for example, *ex vivo* gene therapy. Eukaryotic host cells contemplated as an aspect of this disclosure when harboring a polynucleotide, vector, or protein according to this disclosure include, in addition to a subject's own cells (*e.g.*, a human patient's own cells), VERO cells, HeLa cells, Chinese hamster ovary (CHO)

cell lines (including modified CHO cells capable of modifying the glycosylation pattern of expressed multivalent binding molecules, *see* US Patent Application Publication No. 2003/0115614), COS cells (such as COS-7), W138, BHK, HepG2, 3T3, RIN, MDCK, A549, PC12, K562, HEK293 cells, HepG2 cells, N cells, 3T3 cells, *Spodoptera frugiperda* cells (*e.g.*, Sf9 cells), *Saccharomyces cerevisiae* cells, and any other eukaryotic cell known in the art to be useful in expressing, and optionally isolating, a protein or peptide according to this disclosure. Also contemplated are prokaryotic cells, including *Escherichia coli*, *Bacillus subtilis*, *Salmonella typhimurium*, a Streptomycete, or any prokaryotic cell known in the art to be suitable for expressing, and optionally isolating, a protein or peptide according to this disclosure. In isolating protein or peptide from prokaryotic cells, in particular, it is contemplated that techniques known in the art for extracting protein from inclusion bodies may be used. The selection of an appropriate host is within the scope of those skilled in the art from the teachings herein. Host cells that glycosylate the fusion proteins of this disclosure are contemplated.

**[00201]** The term "recombinant host cell" (or simply "host cell") refers to a cell containing a recombinant expression vector. It should be understood that such terms are intended to refer not only to the particular subject cell but to the progeny of such a cell. Because certain modifications may occur in succeeding generations due to either mutation or environmental influences, such progeny may not, in fact, be identical to the parent cell, but are still included within the scope of the term "host cell" as used herein.

**[00202]** Recombinant host cells can be cultured in a conventional nutrient medium modified as appropriate for activating promoters, selecting transformants, or amplifying particular genes. The culture conditions for particular host cells selected for expression, such as temperature, pH and the like, will be readily apparent to the ordinarily skilled artisan. Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney fibroblasts, described by Gluzman (1981) Cell 23:175, and other cell lines capable of expressing a compatible vector, for example, the C127, 3T3, CHO, HeLa and BHK cell lines. Mammalian expression vectors will comprise an origin of replication, a suitable promoter and, optionally, enhancer, and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5'-flanking nontranscribed sequences, for example, as described herein regarding the preparation of multivalent binding protein expression constructs. DNA sequences derived from the SV40 splice, and polyadenylation sites may be used to provide the required nontranscribed genetic



elements. Introduction of the construct into the host cell can be effected by a variety of methods with which those skilled in the art will be familiar, including calcium phosphate transfection, DEAE-Dextran-mediated transfection, or electroporation (Davis *et al.* (1986) Basic Methods in Molecular Biology).

[00203] In one embodiment, a host cell is transduced by a recombinant viral construct directing the expression of a protein or polypeptide according to this disclosure. The transduced host cell produces viral particles containing expressed protein or polypeptide derived from portions of a host cell membrane incorporated by the viral particles during viral budding.

### **Compositions and Methods of Use**

[00204] To treat human or non-human mammals suffering a disease state associated with TGF $\beta$ , IL6, IL10, GITR, VEGF, TNF, HGF, TWEAK, IGF1 or IGF2 dysregulation, a multi-specific fusion protein of this disclosure is administered to the subject in an amount that is effective to ameliorate symptoms of the disease state following a course of one or more administrations. Being polypeptides, the multi-specific fusion proteins of this disclosure can be suspended or dissolved in a pharmaceutically acceptable diluent, optionally including a stabilizer of other pharmaceutically acceptable excipients, which can be used for intravenous administration by injection or infusion, as more fully discussed below.

[00205] A pharmaceutically effective dose is that dose required to prevent, inhibit the occurrence of, or treat (alleviate a symptom to some extent, preferably all symptoms of) a disease state. The pharmaceutically effective dose depends on the type of disease, the composition used, the route of administration, the type of subject being treated, the physical characteristics of the specific subject under consideration for treatment, concurrent medication, and other factors that those skilled in the medical arts will recognize. For example, an amount between 0.1 mg/kg and 100 mg/kg body weight (which can be administered as a single dose, or in multiple doses given hourly, daily, weekly, monthly, or any combination thereof that is an appropriate interval) of active ingredient may be administered depending on the potency of a binding domain polypeptide or multi-specific protein fusion of this disclosure.

[00206] In certain aspects, compositions of fusion proteins are provided by this disclosure. Pharmaceutical compositions of this disclosure generally comprise one or more type of binding domain or fusion protein in combination with a pharmaceutically acceptable carrier, excipient, or diluent. Such carriers will be nontoxic to recipients at the dosages and

concentrations employed. Pharmaceutically acceptable carriers for therapeutic use are well known in the pharmaceutical art, and are described, for example, in Remington's Pharmaceutical Sciences, Mack Publishing Co. (A.R. Gennaro (Ed.) 1985). For example, sterile saline and phosphate buffered saline at physiological pH may be used. Preservatives, stabilizers, dyes and the like may be provided in the pharmaceutical composition. For example, sodium benzoate, sorbic acid, or esters of *p*-hydroxybenzoic acid may be added as preservatives. *Id.* at 1449. In addition, antioxidants and suspending agents may be used. *Id.* The compounds of the present invention may be used in either the free base or salt forms, with both forms being considered as being within the scope of the present invention.

**[00207]** Pharmaceutical compositions may also contain diluents such as buffers; antioxidants such as ascorbic acid, low molecular weight (less than about 10 residues) polypeptides, proteins, amino acids, carbohydrates (*e.g.*, glucose, sucrose, or dextrans), chelating agents (*e.g.*, EDTA), glutathione or other stabilizers or excipients. Neutral buffered saline or saline mixed with nonspecific serum albumin are exemplary appropriate diluents. Preferably, product is formulated as a lyophilizate using appropriate excipient solutions as diluents.

**[00208]** Compositions of this disclosure can be used to treat disease states in human and non-human mammals that are a result of or associated with TGF $\beta$  or IL6 dysregulation. Increased production or activity of TGF $\beta$  has been implicated in various disease processes, including tumorigenesis, angiogenesis, metastasis, metastatic migration, and epithelial and mesenchymal cancers (*see, e.g.*, Oft et al. (1998) Curr. Biol. 8:1243; Pardali & Mousaka (2007) Biochim. Biophys. Acta 1775:21). In addition, TGF $\beta$  signal transduction has been associated with angiogenesis and the development of vascular disorders (Bertolino et al. (2005) Chest 128:585S).

**[00209]** It has been suggested that IL10 may play a key role in the occurrence of lymphocytic diseases (US Patent No. 5,639,600) and that IL10 may increase proliferation of non-Hodgkin's lymphoma cells (Voorzanger et al. (1996) Cancer Res. 56:5499). More recently it has been proposed that TGF $\beta$  and IL10 work together to ensure a controlled inflammatory response (Li & Flavell, (2008) Immunity 28:468). It has been suggested that tumor-expressed GITRL mediates immunosubversion in humans (Baltz et al. (2007) FASEB J. 21:2442). Overexpression of VEGF and TGF $\beta$  has been associated with the development of cervical cancer (Baritaki et al. (2007) Int. J. Oncol. 31:69). TNF $\alpha$  has been associated with the development of renal cell carcinoma (Harrison et al. (2007) J. Clin. Oncol. 25:4542-

9). TGF $\beta$  has been shown to promote HGF-dependent invasion of squamous carcinoma cells (Lewis et al. (2004) Br. J. Cancer, 90:822), and HGF has been shown to stimulate cell growth and enhance expression of TGF $\alpha$  in human pancreatic cancer cells (Ohba et al. (1999) J. Gastroenterol. 34:498-504). In addition, TGF $\beta$  and HGF have been shown to stimulate the invasiveness of gastric cancer cells (Inoue et al., (197) Jpn. J. Cancer Res. 88:152). IGF1R has been identified in the treatment of cancers, including sarcomas (Scotlandi & Picci (2008) Curr. Opin. Oncol. 20:419-27; Yuen & Macaulay (2008) Expert Opin. Ther. Targets 12:589-603).

**[00210]** IL-6 trans-signaling has been implicated in malignancies, such as colon cancer, while IL6 cis-signaling has been implicated in malignancies including hormone-independent prostate cancer, B-cell proliferative disorders such as B cell non-Hodgkin's lymphoma, and advanced cancers of kidney, breast, colon, lung, brain, and other tissues (*see, e.g.,* Sansone *et al.* (2007) J. Clin Invest. 117:3988). Thus, multi-specific fusion proteins of this disclosure are useful in treating various TGF $\beta$  related autoimmune disorders (such as systemic lupus erythematosus (SLE) or rheumatoid arthritis), Alzheimer's disease or hyperproliferative diseases or malignant disorders, including polycystic kidney disease, lung cancer, colon cancer, urothelial cancer, bladder cancer, renal cell cancer, breast cancer, ovarian cancer, Rhabdomyosarcoma, Ewing's sarcoma, osteosarcoma, neuroblastoma, head & neck cancer, melanoma, glioblastoma, pancreatic cancer, or hepatocarcinoma, or the like.

**[00211]** "Pharmaceutically acceptable salt" refers to a salt of a binding domain polypeptide or fusion protein of this disclosure that is pharmaceutically acceptable and that possesses the desired pharmacological activity of the parent compound. Such salts include the following: (1) acid addition salts, formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like; or formed with organic acids such as acetic acid, propionic acid, hexanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, 3-(4-hydroxybenzoyl)benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, 1,2-ethane-disulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, 4-chlorobenzenesulfonic acid, 2-naphthalenesulfonic acid, 4-toluenesulfonic acid, camphorsulfonic acid, 4-methylbicyclo[2.2.2]-oct-2-ene-1-carboxylic acid, glucoheptonic acid, 3-phenylpropionic acid, trimethylacetic acid, tertiary butylacetic acid, lauryl sulfuric acid, gluconic acid, glutamic acid, hydroxynaphthoic acid, salicylic acid, stearic acid,

muconic acid, and the like; or (2) salts formed when an acidic proton present in the parent compound either is replaced by a metal ion, *e.g.*, an alkali metal ion, an alkaline earth ion, or an aluminum ion; or coordinates with an organic base such as ethanolamine, diethanolamine, triethanolamine, N-methylglucamine, or the like.

[00212] In particular illustrative embodiments, a polypeptide or fusion protein of this disclosure is administered intravenously by, for example, bolus injection or infusion. Routes of administration in addition to intravenous include oral, topical, parenteral (*e.g.*, sublingually or buccally), sublingual, rectal, vaginal, and intranasal. The term parenteral as used herein includes subcutaneous injections, intravenous, intramuscular, intrasternal, intracavernous, intrathecal, intrameatal, intraurethral injection, perispinal or infusion techniques. The pharmaceutical composition is formulated so as to allow the active ingredients contained therein to be bioavailable upon administration of the composition to a patient. Compositions that will be administered to a patient take the form of one or more dosage units, where for example, a tablet may be a single dosage unit, and a container of one or more compounds of this disclosure in aerosol form may hold a plurality of dosage units.

[00213] For oral administration, an excipient and/or binder may be present, such as sucrose, kaolin, glycerin, starch dextrins, cyclodextrins, sodium alginate, ethyl cellulose, and carboxy methylcellulose. Sweetening agents, preservatives, dye/colorant, flavor enhancer, or any combination thereof may optionally be present. A coating shell may also optionally be used.

[00214] In a composition intended to be administered by injection, one or more of a surfactant, preservative, wetting agent, dispersing agent, suspending agent, buffer, stabilizer, isotonic agent, or any combination thereof may optionally be included.

[00215] For nucleic acid-based formulations, or for formulations comprising expression products according to this disclosure, about 0.01  $\mu\text{g/kg}$  to about 100  $\text{mg/kg}$  body weight will be administered, for example, by the intradermal, subcutaneous, intramuscular, or intravenous route, or by any route known in the art to be suitable under a given set of circumstances. A preferred dosage, for example, is about 1  $\mu\text{g/kg}$  to about 20  $\text{mg/kg}$ , with about 5  $\mu\text{g/kg}$  to about 10  $\text{mg/kg}$  particularly preferred. It will be evident to those skilled in the art that the number and frequency of administration will be dependent upon the response of the host.

[00216] The pharmaceutical compositions of this disclosure may be in any form that allows for administration to a patient, such as, for example, in the form of a solid, liquid, or

gas (aerosol). The composition may be in the form of a liquid, *e.g.*, an elixir, syrup, solution, emulsion or suspension, for administration by any route described herein.

**[00217]** A liquid pharmaceutical composition as used herein, whether in the form of a solution, suspension or other like form, may include one or more of the following components: sterile diluents such as water for injection, saline solution (*e.g.*, physiological saline), Ringer's solution, isotonic sodium chloride, fixed oils such as synthetic mono- or diglycerides that may serve as the solvent or suspending medium, polyethylene glycols, glycerin, propylene glycol or other solvents; antibacterial agents such as benzyl alcohol or methyl paraben; antioxidants such as ascorbic acid or sodium bisulfite; buffers such as acetates, citrates or phosphates; chelating agents such as ethylenediaminetetraacetic acid; and agents for the adjustment of tonicity such as sodium, chloride, or dextrose. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic. Physiological saline is a preferred additive. An injectable pharmaceutical composition is preferably sterile.

**[00218]** It may also be desirable to include other components in the preparation, such as delivery vehicles including aluminum salts, water-in-oil emulsions, biodegradable oil vehicles, oil-in-water emulsions, biodegradable microcapsules, and liposomes. Examples of adjuvants for use in such vehicles include N-acetylmuramyl-L-alanine-D-isoglutamine (MDP), lipopolysaccharides (LPS), glucan, IL-12, GM-CSF,  $\gamma$ -interferon, and IL-15.

**[00219]** While any suitable carrier known to those of ordinary skill in the art may be employed in the pharmaceutical compositions of this disclosure, the type of carrier will vary depending on the mode of administration and whether a sustained release is desired. For parenteral administration, the carrier may comprise water, saline, alcohol, a fat, a wax, a buffer, or any combination thereof. For oral administration, any of the above carriers or a solid carrier, such as mannitol, lactose, starch, magnesium stearate, sodium saccharine, talcum, cellulose, glucose, sucrose, magnesium carbonate, or any combination thereof, may be employed.

**[00220]** Also contemplated is the administration of multi-specific fusion protein compositions of this disclosure in combination with a second agent. A second agent may be one accepted in the art as a standard treatment for a particular disease state, such as inflammation, autoimmunity, and cancer. Exemplary second agents contemplated include cytokines, growth factors, steroids, NSAIDs, DMARDs, chemotherapeutics, radiotherapeutics, or other active and ancillary agents.

[00221] This disclosure contemplates a dosage unit comprising a pharmaceutical composition of this disclosure. Such dosage units include, for example, a single-dose or a multi-dose vial or syringe, including a two-compartment vial or syringe, one comprising the pharmaceutical composition of this disclosure in lyophilized form and the other a diluent for reconstitution. A multi-dose dosage unit can also be, *e.g.*, a bag or tube for connection to an intravenous infusion device.

[00222] This disclosure also contemplates a kit comprising a pharmaceutical composition in a unit dose or multi-dose container, *e.g.*, a vial, and a set of instructions for administering the composition to patients suffering a disorder as described herein.

[00223] All U.S. patents, U.S. patent application publications, U.S. patent applications, foreign patents, foreign patent applications, non-patent publications, tables, sequences, webpages, or the like referred to in this specification, are incorporated herein by reference, in their entirety. The following examples are intended to illustrate, but not limit, this disclosure.

## **EXAMPLES**

### **Xceptor Sequences**

[00224] Exemplary IL6 antagonist variable region ( $V_L$  and  $V_H$ ) binding sequences (SEQ ID NO:373-496) are disclosed herein. Also disclosed are amino acid sequences and nucleic acid expression cassettes for exemplary Xceptor fusion proteins comprising a TGF $\beta$ R2 ectodomain and an anti-IL6xR binding domain. Xceptor fusion proteins having a TGF $\beta$ R2 ectodomain at the amino-terminus and an anti-IL6xR binding domain at the carboxy terminus, are referred to herein as TRU(XB6)-1019.1 and TRU(XB6)-1019.2 (amino acid sequences provided in SEQ ID NO:737 and 738, respectively, with the corresponding nucleotide sequences being provided in SEQ ID NO:741 and 742, respectively). The Xceptor fusion proteins in the reverse orientation – that is, having an anti-IL6xR binding domain at the amino-terminus and a TGF $\beta$ R2 ectodomain at the carboxy terminus, are referred to herein as TRU(X6B)-1019.1 and TRU(X6B)-1019.2 (amino acid sequences provided in SEQ ID NO:735 and 736, respectively, with the corresponding nucleotide sequences being provided in SEQ ID NO:739 and 740, respectively).

**Activity Examples**

[00225] Various Xceptor fusion proteins described herein were tested for activity as described below. Abbreviations used in the following examples include the following terms, except where indicated otherwise:

PBS-T: PBS, pH 7.2-7.4 and 0.1% Tween®20

Working buffer: PBS-T with 1% BSA

Blocking buffer (PBS-T with 3% BSA

**EXAMPLE 1****TGFβ BINDING DOMAINS**

[00226] A phage library of Fab binding domains is screened for binding domains specific for either a TGFβ or an IL6xR complex essentially as described by Hoet *et al.* (2005) Nature Biotechnol. 23:344. The binding domains are cloned by PCR amplification. Briefly, the VL and VH regions from the Fab library clones are amplified using PCR SuperMix (Invitrogen, San Diego, CA) and appropriate primers that create the G<sub>4</sub>S linker via overlap, with an initial anneal at 56°C for 9 cycles, then 62 °C for an additional 20 cycles. The PCR products are separated on an agarose gel and purified using a Qiagen (Chatsworth, CA) PCR Purification column. The second round sewing reaction involves mixing a molar equivalent of VL and VH products with Expand buffer and water, denaturing at 95 °C for 5 sec, then cooling slowly to room temperature. To amplify, a mix of dNTPs is added with Expand enzyme and incubated at 72 °C for 10 sec. The outside primers are added (5' VH and 3' VL) and the mix is cycled 35 times with an anneal at 62°C and a 45 min extension reaction. The resulting 750 basepair product is gel purified, digested with EcoRI and NotI, and cloned in plasmid pD28 (for more details, see US Patent Application Publication No. 2005/0136049 and PCT Application Publication No. WO 2007/146968).

**EXAMPLE 2****XCEPTOR BINDING TO IL6 AND HYPER IL6 BY ELISA**

[00227] Hyper-IL6 (HIL6 or IL6xR), recombinant human IL6 (rhIL6), and human soluble IL6R binding activity was examined for exemplary Xceptors TRU(XT6)-1002, 1019, 1025, 1042, 1058, and TRU(X6T)-1019 (SEQ ID NO:608, 625, 631, 648, 664 and 670,

respectively), substantially as follows. Each of these Xceptors includes a TNFRSF1B ectodomain and an anti-IL6xR binding domain.

#### *HIL6 and IL6 Binding*

[00228] Added to each well of a 96-well plate was 100 µl goat anti-human IgG-Fc (Jackson ImmunoResearch, West Grove, PA) from a 2 µg/ml solution in PBS, pH 7.2-7.4. The plate was covered, and incubated overnight at 4°C. After washing four times with PBS (pH 7.2-7.4) and 0.1% Tween®20 (PBS-T), 250 µl Blocking buffer (PBS-T with 3% BSA or 10% normal goat serum) was added to each well, the plate was covered, and incubated at room temperature for 2 hours (or at 4°C overnight). After washing the plate three times with PBS-T, added in duplicate wells to the anti-human IgG-Fc coated plate was 100 µl / well Xceptor TNFRSF1B::anti-HIL6 samples and human gp130-Fc chimera (R&D Systems, Minneapolis, MN) serially diluted three-fold in Working buffer starting at 300 ng/ml, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, added in duplicate wells was 100 µl/well human Hyper IL-6 or recombinant human IL-6 from a 150 pM solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, 100 µl/well anti-human IL-6-biotin (R&D Systems) from a 150 ng/ml solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, 100 µl per well horse radish peroxidase-conjugated streptavidin (Zymed, San Francisco, CA) diluted 1:4,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate six times with PBS-T, 100 µl per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce, Rockford, IL) was added for about 3 to 5 minutes and then the reaction was stopped with 50 µl Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

#### *sIL6R Binding*

[00229] Added to each well of a 96-well plate was 100 µl goat anti-human IgG-Fc (ICN Pharmaceuticals, Costa Mesa, CA) from a 2 µg/ml solution in PBS, pH 7.2-7.4. The plates were covered, and incubated overnight at 4°C. After washing four times with PBS-T, 250 µl Blocking buffer (PBS-T with 3% BSA or 10% normal goat serum) was added to each well, the plate was covered, and incubated at room temperature for 2 hours (or at 4°C overnight). After washing the plate three times with PBS-T, added in duplicate wells to the anti-human IgG-Fc coated plate was 100 µl/well Xceptor TNFRSF1B::anti-HIL6 samples,



positive control anti-human IL-6R (R&D Systems, Minneapolis, MN) and negative controls human IgG or human gp130-Fc chimera (R&D Systems), each serially diluted three-fold in Working buffer starting at 300 ng/ml, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, added in duplicate wells was 100 µl/well recombinant human sIL-6R (R&D Systems) from a 75 pM solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, added 100 µl/well anti-human IL-6R-biotin (R&D Systems) from a 100 ng/ml solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, 100 µl per well horse radish peroxidase-conjugated streptavidin (Zymed, San Francisco, CA) diluted 1:4,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate six times with PBS-T, 100 µl per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce, Rockford, IL) was added for about 3 to 5 minutes and then the reaction was stopped with 50 µl Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

[00230] The data in Figures 1A-1C demonstrate that all Xceptor fusion proteins, whether the TNFRSF1B ectodomain was on the amino- or carboxy terminus of the fusion protein molecules, can bind HIL6. Furthermore, these assays show that the Xceptor proteins have specificity for the IL6xR complex because only two of the Xceptors bind rhIL6 (Figure 1B) and none bind sIL6R (Figure 1C). In related studies, the xceptor TRU(XT6)-1002 and the SMIP TRU(S6)-1002 were found to cross-react with IL6 from the non-human primate *Mucaca mulatta*.

### EXAMPLE 3

#### XCEPTOR BINDING TO TNF- $\alpha$ BY ELISA

[00231] TNF- $\alpha$  binding activity was examined for Xceptors TRU(XT6)-1002, 1042, 1058, 1019, and TRU(X6T)-1019 (SEQ ID NO:608, 648, 664, 625 and 670, respectively), substantially as follows.

[00232] Added to each well of a 96-well plate was 100 µl goat anti-human IgG-Fc (ICN Pharmaceuticals, Costa Mesa, CA) from a 2 µg/ml solution in PBS, pH 7.2-7.4. The plate was covered, and incubated overnight at 4°C. After washing four times with PBS-T, 250 µl Blocking buffer was added to each well, the plate was covered, and incubated at room

temperature for 2 hours (or at 4°C overnight). After washing the plate three times with PBS-T, added in duplicate wells to the anti-human IgG-Fc coated plate was 100 µl/well Xceptor TNFRSF1B::anti-HIL6 samples, positive controls Enbrel® (etanercept) and recombinant human TNFR2 (TNFRSF1B)-Fc chimera (R&D Systems, Minneapolis, MN), and negative controls human IgG or human gp130-Fc chimera (R&D Systems), each serially diluted three-fold in Working buffer starting at 300 ng / ml, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, added in duplicate wells was 100 µl/well recombinant human TNF-α (R&D Systems) from a 2 ng/ml solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, added 100 µl/well anti-human TNF-α-biotin (R&D Systems) from a 200 ng/ml solution in Working buffer, the plate was covered, and incubated at room temperature for about 1 to 2 hours. After washing the plate five times with PBS-T, 100 µl per well horse radish peroxidase-conjugated streptavidin (Jackson ImmunoResearch, West Grove, PA) diluted 1:1,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate six times with PBS-T, 100 µl per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce, Rockford, IL) was added for about 3 to 5 minutes and then the reaction was stopped with 50 µl Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

[00233] The data in Figure 2 shows that all Xceptor fusion proteins tested can bind TNF-α, whether the TNFRSF1B ectodomain was on the amino- or carboxy terminus of the fusion protein.

#### EXAMPLE 4

##### XCEPTOR DUAL LIGAND BINDING BY ELISA

[00234] Concurrent binding to TNF-α and to IL6xR complex was examined for Xceptor fusion protein TRU(XT6)-1006 (SEQ ID NO:612), substantially as follows.

[00235] Added to each well of a 96-well plate was 100 µl human HIL-6 solution (5 µg/ml in PBS, pH 7.2-7.4). The plate was covered, and incubated overnight at 4°C. After washing four times with PBS-T, then 250 µl Blocking buffer was added to each well, the plate was covered, and incubated at room temperature for 2 hours (or at 4°C overnight). After washing the plate three times with PBS-T, added in duplicate wells to the HIL-6 coated plate was 100 µl/well Xceptor TNFRSF1B::HIL6 samples serially diluted three-fold in

Working buffer starting at 300 ng / ml. Negative controls included human gp130-Fc chimera (R&D Systems, Minneapolis, MN), Enbrel® (etanercept), and Working buffer only. The plate was covered and incubated at room temperature for 1.5 hours. After washing the plate five times with PBS-T, 100 µl per well recombinant human TNF- $\alpha$  (R&D Systems, Minneapolis, MN) to 2 ng / ml in Working buffer was added, the plate was covered, and incubated at room temperature for 1.5 hr. After washing the plate five times with PBS-T, 100 µl per well anti-human TNF- $\alpha$ -biotin (R&D Systems) to 200 ng/ml in Working buffer was added, the plate was covered, and incubated at room temperature for 1.5 hr. After washing the plate five times with PBS-T, 100 µl per well horse radish peroxidase-conjugated streptavidin (Jackson ImmunoResearch, West Grove, PA) diluted 1:1000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate six times with PBS-T, 100 µl per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce, Rockford, IL) was added for 3-5 minutes and then the reaction was stopped with 50 µl Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

**[00236]** The data in Figure 3 demonstrates that Xceptor proteins can bind two ligands simultaneously (in this case TNF- $\alpha$  and Hyper IL6).

## EXAMPLE 5

### XCEPTOR BLOCKING OF HYPER IL6 BINDING TO GP130 BY ELISA

**[00237]** Blocking of Hyper IL6 (IL6xR) binding to soluble gp130 receptor by Xceptor fusion proteins TRU(XT6)-1004, 1006, 1007, 1008, 1013, and 1019 (SEQ ID NO:610, 612, 613, 614, 619 and 625, respectively), was examined substantially as follows.

**[00238]** Added to each well of a 96-well plate was 100 µl human gp130-Fc chimera (R&D Systems, Minneapolis, MN) from of 0.25 – 0.5 µg/ml solution in PBS, pH 7.2-7.4. The plates were covered, and incubated overnight at 4°C. After washing four times with PBS-T, 250 µl Blocking buffer (PBS-T with 3% BSA or 10% normal goat serum) was added to each well, the plate was covered, and incubated at room temperature for 2 hours (or at 4°C overnight). Serial five-fold dilutions in Working buffer starting at 50 µg/ml were made of the following samples: Xceptor TNFRSF1B::anti-HIL6 samples, positive controls human gp130-Fc chimera (R&D Systems) and anti-human IL-6R (R&D Systems), and negative controls anti-human IL-6 (R&D Systems), human IgG or Enbrel® (etanercept). Equal volumes of the serially diluted Xceptor samples were mixed with Hyper IL-6 (final Hyper IL-6 concentration

of 2.5 ng/ml) and incubated at room temperature for 1 hour. After washing the plate three times with PBS-T, added in duplicate wells to the human gp130-Fc coated plate was 100  $\mu$ l/well of the serially dilutions of Xceptor / HIL6 mixtures, human gp130-Fc chimera, anti-human IL-6R, anti-human IL-6, human IgG, and Enbrel® (etanercept), the plate was covered, and incubated at room temperature for about 1.5 hours. After washing the plate five times with PBS-T, 100  $\mu$ l per well horse radish peroxidase-conjugated anti-mouse IgG-Fc (Pierce, Rockford, IL) diluted 1:10,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 1 hour. After washing the plate six times with PBS-T, 100  $\mu$ l per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce) was added for about 5 to 15 minutes and then the reaction was stopped with 50  $\mu$ l Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

[00239] The data in Figure 4 demonstrate that Xceptor proteins comprising an anti-IL6xR binding domain can block soluble gp130 from binding to HIL6.

## EXAMPLE 6

### XCEPTOR BLOCKING OF IL6 AND HYPER IL6 INDUCED CELL PROLIFERATION

[00240] Blocking of IL6 or Hyper IL6 (IL6xR) induced cell proliferation of TF-1 cells was examined for Xceptor fusion proteins TRU(XT6)-1011, 1014, 1025, 1026, 1002, and TRU(X6T)-1019 (SEQ ID NO:617, 620, 631, 632, 608 and 670, respectively), substantially as follows.

[00241] Added to each well of a 96-well flat bottom plate were  $0.3 \times 10^6$  TF-1 cells (human erythroleukemia cells) in the fresh growth medium (10% FBS-RPMI 1640; 2mM L-glutamine; 100 units/ml penicillin; 100  $\mu$ g/ml streptomycin; 10 mM HEPES; 1mM sodium pyruvate; and 2 ng/ml Hu GM-CSF) one day before use in proliferation assay. The cells were then harvested and washed twice with assay medium (same as growth medium except without GM-CSF, cytokine-free), then resuspended at  $1 \times 10^5$  cells/ml in assay medium. For blocking IL-6 activity, serial dilutions of a TNFSFR1B::anti-HIL-6 Xceptor of interest or antibody was pre-incubated with a fixed concentration of recombinant human IL-6 (rhIL-6) (R&D Systems, Minneapolis, MN) or hyper IL-6 (HIL-6) in 96-well plates for 1 hour at 37°C, 5%CO<sub>2</sub>. Controls used included human IgG; human gp130-Fc chimera (R&D Systems); anti-hIL-6 antibody (R&D Systems); and anti-hIL-6R antibody (R&D Systems). After the pre-incubation period,  $1 \times 10^4$  cells (in 100  $\mu$ l) was added to each well. The final assay mixture, in a total volume of 200  $\mu$ L/well, containing TNFSFR1B::HIL-6, rhIL-6, or

HIL-6 and cells was incubated at 37°C, 5%CO<sub>2</sub> for 72 hours. During the last 4-6 hours of culture, <sup>3</sup>H-thymidine (20 µCi/ml in assay medium, 25 µL/well) was added. The cells were harvested onto UniFilter-96 GF/c plates and incorporated <sup>3</sup>H-Thymidine was determined using TopCount reader (Packard). The data are presented as the Mean of cpm ± SD of triplicates. The percentage of blocking = 100 - (test cpm – control cpm / maximum cpm - control cpm)\*100.

[00242] The data in Figure 5A and Figure 5B demonstrate that all Xceptor proteins, whether the TNFRSF1B ectodomain was on the amino- or carboxy terminus of the fusion protein molecules, can block cell proliferation induced by IL6 or Hyper IL6, respectively, or both.

### EXAMPLE 7

#### XCEPTOR BLOCKING OF TNF- $\alpha$ BINDING TO TNFR BY ELISA

[00243] Blocking of TNF- $\alpha$  binding to TNF receptor by Xceptor fusion proteins TRU(XT6)-1004, 1006, 1007, 1008, 1013, and 1019 (SEQ ID NO:610, 612, 613, 614, 619 and 625, respectively) was examined substantially as follows.

[00244] Added to each well of a 96-well plate was 100 µL recombinant human TNFR2-Fc chimera (R&D Systems, Minneapolis, MN) from of 0.25 – 0.5 µg/ml solution in PBS, pH 7.2-7.4. The plates were covered, and incubated overnight at 4°C. After washing four times with PBS-T, 250 µL Blocking buffer (PBS-T with 3% BSA or 10% normal goat serum) was added to each well, the plate was covered, and incubated at room temperature for 2 hours (or at 4°C overnight). Serial five-fold dilutions in Working buffer starting at 50 to 250 µM were made of the following samples: Xceptor TNFRSF1B::anti-HIL6 samples, positive controls Enbrel® (etanercept) and anti-TNF- $\alpha$  (R&D Systems), and negative controls human gp130-Fc chimera (R&D Systems) and human IgG. Equal volumes of the serially diluted Xceptor samples were mixed with TNF- $\alpha$  (final TNF- $\alpha$  concentration of 2.5 ng/ml) and incubated at room temperature for 1 hour. After washing the plate three times with PBS-T, added in duplicate wells to the recombinant human TNFR2-Fc coated plate was 100 µl/well of the serially dilutions of Xceptor / TNF- $\alpha$  mixture, Enbrel® (etanercept), anti-TNF- $\alpha$ , human gp130-Fc chimera, and human IgG, the plate was covered, and incubated at room temperature for about 1.5 hours. After washing the plate five times with PBS-T, 100 µL per well of anti-human TNF- $\alpha$ -biotin (R&D Systems) from a 200 ng/ml solution in

Working buffer was added, the plate was covered, and incubated at room temperature for 1 to 2 hours. After washing the plate five times with PBS-T, 100  $\mu$ L per well horse radish peroxidase-conjugated streptavidin (Jackson ImmunoResearch, West Grove, PA) diluted 1:1,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate six times with PBS-T, 100  $\mu$ L per well 3,3',5,5'-tetramethylbenzidine (TMB) substrate solution (Pierce, Rockford, IL) was added for about 3 to 5 minutes and then the reaction was stopped with 50  $\mu$ L Stop buffer (1N H<sub>2</sub>SO<sub>4</sub>) per well. The absorbance of each well was read at 450 nm.

[00245] The data in Figure 6 show that Xceptor proteins blocked TNF- $\alpha$  binding to TNF receptor, which was approximately equivalent to blocking by TNFR-Fc.

### EXAMPLE 8

#### XCEPTOR BLOCKING OF TNF- $\alpha$ INDUCED CELL KILLING

[00246] Blocking of TNF- $\alpha$  induced killing of L929 cells was examined for Xceptor fusion proteins TRU(XT6)-1011, 1014, 1025, 1026, 1002, and TRU(X6T)-1019 (SEQ ID NO:617, 620, 631, 632, 608 and 670, respectively), substantially as follows.

[00247] A suspension of L929 mouse fibroblast cells (ATCC, Manassas, VA) was prepared at a density of  $2 \times 10^5$  cells/ml in culture medium (10% FBS-RPMI 1640; 2 mM L-glutamine; 100 units/ml penicillin; 100  $\mu$ g/ml streptomycin; and 10 mM HEPES), then 100  $\mu$ L was added to each well of a 96-well flat bottom black plate and incubated overnight at 37°C, 5% CO<sub>2</sub> in a humidified incubator. Xceptor TNFRSF1B::anti-HIL6 samples serially diluted in assay medium (same as culture medium but supplemented with 2% FBS) were mixed with an equal volume of recombinant human TNF- $\alpha$  (rhTNF- $\alpha$ ; R&D Systems, Minneapolis, MN), and incubated at 37°C, 5% CO<sub>2</sub> in a humidified incubator for 1 hour. Positive controls (*i.e.*, those agents that block TNF- $\alpha$  induced killing of L929 cells) included Enbrel® (etanercept), rhTNFR2-Fc chimera (R&D Systems, Minneapolis, MN), and anti-TNF- $\alpha$  antibody (R&D Systems, Minneapolis, MN). Negative controls included assay medium alone (no TNF- $\alpha$  added) and antibody hIgG (with TNF- $\alpha$  added). To analyze TNF- $\alpha$  activity, culture medium was removed from the L929 cells and then each well received 50  $\mu$ L of a TNF- $\alpha$ /Xceptor or control mixture, and 50  $\mu$ L actinomycin D (Sigma-Aldrich, St. Louis, MO) (from a freshly prepared working solution of 4  $\mu$ g/ml). The cells were then incubated for 24 hrs at 37°C, 5% CO<sub>2</sub> in a humidified incubator. To measure cell viability, added to each well was 100  $\mu$ L ATPlite 1 Step Reagent (PerkinElmer, Waltham, MA) according to the manufacturer's

instructions, shaken for two minutes, and then luminescence is measured using a TopCount reader (Packard).

[00248] The data in Figure 9 demonstrate that all Xceptor proteins, whether the TNFRSF1B ectodomain was on the amino- or carboxy terminus of the fusion protein molecules, can block TNF- $\alpha$  induced cell killing in this assay.

## EXAMPLE 9

### XCEPTOR BINDING TO TGF $\beta$ BY ELISA

[00249] TGF $\beta$  binding activity was examined for Xceptors X6B and XB6, substantially as follows.

[00250] Added to each well of a 96-well plate was 100  $\mu$ l goat anti-human IgG-Fc (ICN Pharmaceuticals, Costa Mesa, CA) from a 2  $\mu$ g/ml solution in PBS, pH 7.2-7.4. The plate was covered, and incubated overnight at 4°C. After washing four times with PBS-T, 250  $\mu$ l Blocking buffer (PBS-T with 10% NGS) was added to each well, the plate was covered, and incubated at room temperature for 2 hours. After washing the plate three times with PBS-T, added in duplicate wells to the anti-human IgG-Fc coated plate was 100  $\mu$ l/well Xceptor TGF $\beta$ R2::anti-HIL6 samples, positive control recombinant human TGF $\beta$ RII-Fc chimera (R&D Systems, Minneapolis, MN), and negative control recombinant human TNFR2 (TNFRSF1B)-Fc chimera (R&D Systems), each diluted to 300ng/ml in Working buffer (PBS-T with 1% BSA). The plate was covered, and incubated at room temperature for about 1 hour. After washing the plate five times with Wash buffer (PBS/0.1% Tween 20 (PBS-T)), added was 100  $\mu$ l/well TGF $\beta$ -1 ligand (R&D Systems), serially diluted two-fold in Working buffer starting at 4 ng/ml. The plate was covered, and incubated at room temperature for 1 hour. After washing the plate five times with Wash buffer, added was 100  $\mu$ l/well biotinylated anti-TGF $\beta$ -1 (R&D Systems) from a 200 ng/ml solution in Working buffer. The plate was covered and incubated at room temperature for 1 hour. After washing the plate five times with Wash buffer, 100  $\mu$ l per well horse radish peroxidase-conjugated streptavidin (Pierce Rockford, IL) diluted 1:20,000 in Working buffer was added, the plate was covered, and incubated at room temperature for 30 minutes. After washing the plate five times with Wash buffer, 100  $\mu$ l per well QuantaBlu Fluorogenic Peroxidase Substrate solution (Pierce, Rockford, IL; prepared by mixing 9 ml substrate solution with 1 ml peroxide solution) was added, and the plate was covered and incubated at room temperature for 20

min. The reaction was stopped with 50  $\mu$ l QuantaBlu Stop Solution per well. The absorbance of each well was read at 325 nm.

[00251] The data in Figure 8 shows that the Xceptor fusion proteins tested bound TGF $\beta$ -1, whether the TGF $\beta$ R2 ectodomain was on the amino- or carboxy terminus of the fusion protein.

## EXAMPLE 10

### EXPRESSION OF XCEPTOR FUSION PROTEINS

[00252] Expression of certain of the Xceptor fusion proteins disclosed herein in 293 cells was performed using the FreeStyle™ 293 Expression System (Invitrogen, Carlsbad, CA) according to the manufacturer's instructions.

[00253] For each 30 ml transfection,  $3 \times 10^7$  cells in 28 ml of FreeStyle™ 293 Expression Medium were used. On the day of transfection, a small aliquot of the cell suspension was transferred to a microcentrifuge tube, and the viability and the amount of cell clumping determined using the trypan blue dye exclusion method. The suspension was vigorously vortexed for 45 seconds to break up cell clumps and total cell counts determined using a Coulter Counter or a hemacytometer. The viability of the cells was over 90%. A shaker flask containing the required cells was placed in a 37°C incubator on an orbital shaker.

[00254] For each transfection sample, lipid-DNA complexes were prepared as follows. 30  $\mu$ g of plasmid DNA was diluted in Opti-MEM® I to a total volume of 1 ml and mixed gently. 60  $\mu$ l of 293fectin™ was diluted in Opti-MEM® I to a total volume of 1 ml, mixed gently, and incubated for 5 minutes at room temperature. After the 5 minute incubation, the diluted DNA was added to the diluted 293fectin™ to obtain a total volume of 2 ml and mixed gently. The resulting solution was incubated for 20-30 minutes at room temperature to allow DNA- 293fectin™ complexes to form.

[00255] While the DNA-293fectin™ complexes were incubating, the cell suspension was removed from the incubator and the appropriate volume of cell suspension was placed in a sterile, disposable 125 ml Erlenmeyer shaker flasks. Fresh, pre-warmed FreeStyle™ 293 Expression Medium was added up to a total volume of 28 ml for a 30 ml transfection.

[00256] After the DNA-293fectin™ complex incubation was complete, 2 ml of DNA-293fectin™ complex was added to the shaker flasks. 2 ml of Opti-MEM® I was added to the negative control flask, instead of DNA-293fectin™ complex. Each flask contained a total volume of 30 ml, with a final cell density of approximately  $1 \times 10^6$  viable cells/ml. The



cells were incubated in a 37°C incubator with a humidified atmosphere of 8% CO<sub>2</sub> in air on an orbital shaker rotating at 125 rpm. Cells were harvested at approximately 7 days post-transfection and assayed for recombinant protein expression.

[00257] Xceptor molecules having a TNFRSF1B ectodomain and a TGFβRII ectodomain were expressed in 293 cells as described above.

## EXAMPLE 11

### XCEPTOR BINDING TO LIGANDS BY ELISA

[00258] The ability of xceptor molecules comprising a TNFRSF1B ectodomain and either a TWEAKR ectodomain, an OPG ectodomain, a TGFβRII ectodomain or an IL7R ectodomain to bind to the ligands TWEAK, RANKL, TGFβ or IL7, respectively, was examined substantially as follows.

[00259] Mouse and human ligands (R&D Systems, Minnesota, MN) were added to wells of a 96-well plate at a concentration of 1μg/ml in PBS (100 μL/well). Plates were incubated at 4°C overnight. After washing five times with PBS-T, 250 μL Blocking Buffer (PBS-T with 3% BSA) was added to each well, and the plate covered and incubated at room temperature (RT) for 2 hours. Serial three fold dilutions of xceptors were made in Working Buffer (PBS-T with 1% BSA) starting at 300ng/ml. As a negative control, an irrelevant xceptor was used. The plate was incubated at RT for 1 hour. After washing five times with PBS-T, 100 μL per well of HRP-conjugated anti-human IgG-Fc (1:5000 in Working buffer) was added, the plate covered, and incubated at RT for 1 hour. After washing five times with PBS-T, 100 μL of Quant-Blu substrate (Pierce, Rockford, IL) was added to each well. The plate was incubated at RT for 10-30 minutes, and fluorescence measured at 325/420nm.

[00260] The results are shown in Table 3 below. The binding of the TNFRxTGFβRII to mouse TGFβ was not tested, however it is noted that mouse and human TGFβ are 99% identical.

**Table 3. Xceptor binding to Ligands**

TNFR x R	Ligand	Mouse ligand binding	Human ligand binding
TNFR x TWEAKR	TWEAK	+++	+++
TNFR x OPG	RANKL	+++	+++
TNFR x TGFβRII	TGFβ	homologous	+++
TNFR x IL7R	IL7	ND	+

ND = Not Done

## EXAMPLE 12

### XCEPTOR BLOCKING OF TGF $\beta$ -1-INDUCED INHIBITION OF CELL PROLIFERATION

[00261] Blocking of TGF $\beta$ -1 induced inhibition of IL-4 proliferation of HT2 cells was examined for the Xceptor of SEQ ID NO:1236 using the method described by Tsang et al. (Tsang, M. et al. (1995) Cytokine 7:389).

[00262] Briefly, in a 96 well plate, Xceptor TNFR::TGF $\beta$ RII samples were serially diluted in culture medium (RPMI, 10% FCS, 0.05 mM beta-mercaptoethanol) containing 1 ng/ml of human TGF $\beta$ -1; 100  $\mu$ l per well. The plate was incubated at 37° C, 5% CO<sub>2</sub> in a humidified incubator for 1.5 hours. Negative controls included an irrelevant xceptor protein (with TGF $\beta$ -1 added) and culture medium (with and without TGF $\beta$ -1 added). The positive control was a recombinant TGF $\beta$ RII-Fc chimera (R&D Systems, Minneapolis, MN). Following incubation, 1x10<sup>4</sup> HT2 cells (ATCC, Manassas, VA) in 100  $\mu$ l of culture medium containing 15 ng/ml mIL4 (R&D Systems, Minneapolis, MN) was added to each well. The plate was then incubated at 37°C, 5% CO<sub>2</sub> in a humidified incubator for 72 hours.

[00263] To analyze TGF $\beta$ -1 activity by measuring cell viability, 100  $\mu$ l of culture medium was removed from each well and replaced with 10  $\mu$ L WST-8 reagent (Dojindo Molecular Technologies, Rockville, MD). The plate was incubated for 2 hours at 37°C, and absorbance for each well read at 450nm.

[00264] The data in Figure 9 shows that the xceptor protein blocked TGF $\beta$ -1 inhibition of IL4-mediated proliferation of HT2 cells.

## EXAMPLE 13

### SPECIFICITY OF BINDING TO HYPER IL6 AND NOT OTHER GP130 CYTOKINES

[00265] The effect of Xceptor fusion proteins on induction of TF-1 cell proliferation by IL6 and the gp130 cytokines IL-11, leukemia inhibitory factor (LIF), oncostatin M (OSM) and cardiotrophin-1 (CT-1) was examined substantially as follows.

[00266] Added to each well of a 96-well flat bottom plate was 0.3x10<sup>6</sup> TF-1 cells (human erythroleukemia cells) in fresh growth medium (10% FBS-RPMI 1640, 2mM L-glutamine, 100 units/ml penicillin, 100  $\mu$ g/ml streptomycin, 10 mM HEPES, 1mM sodium pyruvate and 2 ng/ml Hu GM-CSF) one day before use in the proliferation assay. The cells

were harvested and washed twice with assay medium (same as growth medium except without GM-CSF, cytokine-free), then resuspended at  $1 \times 10^5$  cells/ml in assay medium. For examining blocking of LIF, OSM, and CT-1 activity, serial dilutions of a TNFSFR1B::anti-HIL-6 xceptors TRU(XT6)-1002 (SEQ ID NO:608), TRU(XT6)-1019 (SEQ ID NO:625), TRU(XT6)-1022 (SEQ ID NO:628), and TRU(XT6)-1025 (SEQ ID NO:631) were pre-incubated with a fixed concentration of each gp130 cytokine individually or hyper IL-6 (HIL-6) in 96-well plates for 1 hour at 37°C, 5% CO<sub>2</sub>. After the pre-incubation period,  $1 \times 10^4$  cells (in 100 µl) were added to each well. The final assay mixture, in a total volume of 200 µL/well, containing TNFSFR1B::HIL-6, gp130 cytokine or HIL-6 and cells, was incubated at 37°C, 5% CO<sub>2</sub> for 72 hours. During the last 4-6 hours of culture, <sup>3</sup>H-thymidine (20 µCi/ml in assay medium, 25 µL/well) was added. The cells were harvested onto UniFilter-96 GF/c plates and incorporated <sup>3</sup>H-Thymidine was determined using TopCount reader (Packard). The percentage of blocking =  $100 - (\text{test cpm} - \text{control cpm} / \text{maximum cpm} - \text{control cpm}) * 100$ .

[00267] The results showed that the xceptor blocked IL6 activity but not IL-11, LIF, OSM or CT-1 (data not shown), and therefore bound to hyper IL6 but had no effect on the other gp130 cytokines tested.

#### EXAMPLE 14

##### SMIP AND XCEPTOR BINDING TO IL6R ON LIVER CELLS

[00268] The ability of TRU(S6)-1002, TRU(XT6)-1019 and the anti-IL6 antibody hu-PM1 to bind to IL6R on the liver-derived HepG2 cells was examined as follows.

[00269] HepG2 cells were washed in FACS Buffer and adjusted to  $2 \times 10^6$  cells/mL in FACS Buffer (PBS + 3% FBS). To wells of a 96-well plate were added 50 µL of this solution ( $10^5$  cells/well). The plates were held at 37°C until ready to add diluted test molecules. Serial dilutions of the test molecules were prepared in FACS Buffer to give a 2X working stock which was diluted to 1X when added to cells. The diluted test molecules were added to cells (50 µL/well) and the cells incubated for 20 min on ice. Whole IgG was used as a control. The cells were then washed two times with FACS Buffer and resuspended in phycoerythrin-conjugated goat anti-human antibody (Jackson Labs; diluted 1:200 in FACS Buffer). After being incubated for 20 min on ice in the dark, the cells were washed two times with FACS buffer, resuspended in 200ul PBS and read on a LSRII™ flow cytometer (BD Biosciences, San Jose, CA).

[00270] As shown in Fig. 10, TRU(S6)-1002 and TRU(XT6)-1029 showed essentially no binding to HepG2 cells.

### EXAMPLE 15

#### SMIP AND XCEPTOR BLOCKING OF IL-6 AND TNF ACTIVITY IN MICE

[00271] The ability of SMIP and Xceptor fusion proteins disclosed herein to block IL-6 or TNF-induced production of serum amyloid A (SAA) protein in mice was examined as described below. SAA is one of the major acute-phase proteins in humans and mice. Prolonged elevation of plasma SAA levels is found in chronic inflammation and leads to amyloidosis which affects the liver, kidney and spleen (Rienhoff et al., (1990) Mol. Biol. Med. 7:287). Both IL-6 and TNF have been shown to induce SAA when administered alone (Benigni et al., (1996) Blood 87:1851; Ramadori et al., (1988) Eur. J. Immunol. 18:1259).

##### (a) Blocking of hyperIL-6 activity

[00272] Female BALB/C mice were injected retro-orbitally with 0.2 ml PBS, or Enbrel® (200 ug), TRU(S6)-1002 (200 ug) or TRU(XT6)-1002 (300 ug or 500ug) in PBS. One hour later, the mice were injected IP with 0.2 ml PBS or 2 µg human hyper-IL6 in PBS. Mouse serum was collected at 2 hours and 24 hours after the IP injection. The serum concentration of SAA was determined by ELISA, and concentration of sgp130 was determined by a Luminex-based mouse soluble receptor assay. As shown in Figs. 11 and 12, TRU(S6)-1002 and TRU(XT6)-1002 blocked hyperIL6-induced expression of both sgp130 and SAA.

##### (b) Blocking of TNF activity

[00273] Female BALB/C mice were injected retro-orbitally with 0.2 ml PBS, or Enbrel® (200 µg), TRU(S6)-1002 (200 µg) or TRU(XT6)-1002 (300 µg) in PBS. One hour later, the mice were injected IP with 0.2 ml PBS or 0.5 ug mouse TNF-α in PBS. Mouse serum was collected at 2 hours and 24 hours after the IP injection. The serum concentration of SAA was determined by ELISA, and concentration of sgp130 was determined by a Luminex-based mouse soluble receptor assay. As shown in Figs. 13A and B, the Xceptor TRU(XT6)-1002 blocked TNFα-induced expression of SAA, with the level of SAA observed at 2 hours post-injection being similar to that seen with Enbrel®.

**EXAMPLE 16**  
**XCEPTOR ACTIVITY *IN VIVO***

[00274] The therapeutic efficacy of Xceptor molecules described herein is examined in animal models of disease as described below.

**(a) Multiple Myeloma**

[00275] The activity of Xceptor molecules is examined in at least one of two well characterized mouse models of multiple myeloma, namely the 5T2 multiple myeloma (5T2MM) model and the 5T33 multiple myeloma (5T33MM) model. In the 5T33 model, mice are treated with Xceptors from the time of injection of tumor cells (prophylactic mode). In the 5T2MM model, mice are treated from the onset of the disease (therapeutic mode). The effect of treatment on tumor development and angiogenesis is assessed in both models, with bone studies also being performed in the 5T2MM model.

[00276] The 5TMM murine model of myeloma was initially developed by Radl et al. (J. Immunol. (1979) 122:609; *see also* Radl et al. Am. J. Pathol. (1988) 132:593; Radl J. Immunol. Today (1990) 11:234). Its clinical characteristics resemble the human disease closely: the tumor cells are located in the bone marrow, the serum paraprotein concentration is a measure of disease development, neovascularization is increased in both the 5T2MM and 5T33MM models (Van Valckenborgh et al., Am. J. Pathol. (1988) 132:593), and in certain lines a clear osteolytic bone disease develops. The 5T2MM model includes moderate tumor growth and the development of osteolytic bone lesions. These lesions are associated with a decrease in cancellous bone volume, decreased bone mineral density and increased numbers of osteoclasts (Croucher et al., Blood (2001) 98:3534). The 5T33MM model has a more rapid tumor take and, in addition to the bone marrow, tumor cells also grow in the liver (Vanderkerken et al., Br. J. Cancer (1997) 76:451).

[00277] The 5T2 and 5T33MM models have been extensively characterized. Specific monoclonal antibodies have been raised against the idiotype of both 5T2 and 5T33MM allowing the detection, with great sensitivity, of the serum paraprotein by ELISA, and the specific staining of the tumor cells both by FACS analysis and immunostaining of histological sections (Vanderkerken et al., Br. J. Cancer (1997) 76:451). The sequence analysis of the VH gene enables the detection of cells by RT-PCR and Northern blot analysis (Zhu et al., Immunol. (1998) 93:162). The 5TMM models, which can be used for both *in vitro* and *in vivo* experiments, generate a typical MM disease and different methods are

available to assess tumor load in the bone marrow, serum paraprotein concentrations, bone marrow angiogenesis (by measuring the microvessel density) and osteolytic bone lesions (by a combination of radiography, densitometry and histomorphometry). The investigation of these latter parameters allow the use of the 5TMM models in a preclinical setting and study of the growth and biology of the myeloma cells in a complete syngeneic microenvironment. Both molecules targeting the MM cells themselves and molecules targeting the bone marrow microenvironment can be studied. Specifically, while the 5T33MM model can be used to study both the microenvironment and the MM cells themselves, the 5T2MM model can also be used to study the myeloma associated bone disease.

[00278] To study the prophylactic efficacy of the Xceptor molecules disclosed herein, C57BL/KaLwRij mice are injected with  $2 \times 10^6$  5T33 MM cells and with Xceptor on day 0. Mice are sacrificed at day 28 and tumor development is assessed by determining serum paraprotein concentration and the percentage of tumor cells on isolated bone marrow cells (determined by flow cytometry with anti-idiotypic antibodies or by cytosmears). The weight of spleen and liver is determined and these organs are fixed in 4% formaldehyde for further analysis. Bone samples are fixed for further processing including CD31 immunostaining on paraffin sections and quantification of microvessel density.

[00279] To study the therapeutic efficacy of the Xceptor molecules disclosed herein, mice are injected with 5T2MM cells on day 0, and Xceptor is administered following the onset of disease, as determined by the presence of detectable levels of serum paraprotein. Mice are sacrificed approximately five weeks following administration of Xceptor, and tumor development is assessed as described above for the prophylactic study. In addition, bone analysis is performed using X-rays to determine the number of bone lesions and trabecular bone area, and TRAP staining to assess the number of osteoclasts.

#### **(b) Rheumatoid Arthritis**

[00280] The therapeutic efficacy of the Xceptor molecules disclosed herein is examined in at least one of two murine models of rheumatoid arthritis (RA), namely the collagen induced arthritis (CIA) and glucose-6-phosphate isomerase (G6PI) models. Each of these models has been shown by others to be useful for predicting efficacy of certain classes of therapeutic drugs in RA (*see* Holmdahl (2000) Arthritis Res. 2:169; Holmdahl (2006) Immunol. Lett. 103:86; Holmdahl (2007) Methods Mol. Med. 136:185; McDevitt (2000) Arthritis Res. 2:85; Kamradt and Schubert (2005) Arthritis Res. Ther. 7:20).

(i) CIA Model

[00281] The CIA model is the best characterized mouse model of arthritis in terms of its pathogenesis and immunological basis. In addition, it is the most widely used model of RA and, although not perfect for predicting the ability of drugs to inhibit disease in patients, is considered by many to be the model of choice when investigating potential new therapeutics for RA (Jirholt, J. et al. (2001) *Arthritis Res.* 3:87-97; Van den Berg, W.B. (2002) *Curr. Rheumatol. Rep.* 4:232-239; Rosloniec, E. (2003) *Collagen-Induced Arthritis*. In *Current Protocols in Immunology*, eds. Coligan et al., John Wiley & Sons, Inc, Hoboken, NJ).

[00282] In the CIA model, arthritis is induced by immunization of male DBA/1 mice with collagen II (CII) in Complete Freund's Adjuvant (CFA). Specifically, mice are injected intradermally/ subcutaneously with CII in CFA on Day -21 and boosted with CII in Incomplete Freund's Adjuvant (IFA) on Day 0. Mice develop clinical signs of arthritis within days of the boost with CII/IFA. A subset of mice (0% to 10%) immunized with CII/CFA develop signs of arthritis on or around Day 0 without a boost and are excluded from the experiments. In some CIA experiments, the boost is omitted and mice are instead treated with Xceptor or control starting 21 days after immunization with CII/CFA (i.e. the day of first treatment is Day 0).

[00283] Mice are treated with Xceptor, vehicle (PBS), or negative or positive control in a preventative and/or therapeutic regimen. Preventative treatment starts on Day 0 and continues through the peak of disease in control (untreated) mice. Therapeutic treatment starts when the majority of mice show mild signs of arthritis. Enbrel<sup>®</sup>, which has been shown to have good efficacy in both the CIA and G6PI-induced models of arthritis, is used as a positive control. Data collected in every experiment includes clinical scores and cumulative incidence of arthritis. Clinical signs of arthritis in the CIA model are scored using a scale from 0 to 4 as shown in Table 4 below:

**Table 4.**

<b>Score</b>	<b>Observations</b>
0	No apparent swelling or redness
1	Swelling/redness in one to three digits
2	Redness and/or swelling in more than three digits, mild swelling extending into the paw, swollen or red ankle, or mild swelling/redness of forepaw
3	Swollen paw with mild to moderate redness
4	Extreme redness and swelling in entire paw

(ii) G6PI Model

[00284] In the G6PI model, arthritis is induced by immunization of DBA/1 mice with G6PI in adjuvant (Kamradt, T. and D. Schubert (2005) *Arthritis Res. Ther.* 7:20-28; Schubert, D. et al. (2004) *J. Immunol.* 172:4503-4509; Bockermann, R. et al. (2005) *Arthritis Res. Ther.* 7:R1316-1324; Iwanami, K. et al. (2008) *Arthritis Rheum.* 58:754-763; Matsumoto, I. et al. (2008) *Arthritis Res. Ther.* 10:R66). G6PI is an enzyme present in virtually all cells in the body and it is not known why immunization induces a joint specific disease. A number of agents, such as CTLA4-Ig, TNF antagonists (e.g. Enbrel®) and anti-IL6 receptor monoclonal antibody, have been shown to inhibit development of arthritis in the G6PI model.

[00285] Male DBA/1 mice are immunized with G6PI in Complete Freund's Adjuvant (CFA) in order to induce arthritis. Specifically, mice are injected intradermally/subcutaneously with G6PI in CFA on Day 0 and develop clinical signs of arthritis within days of the immunization. As with the CIA model discussed above, mice are treated with Xceptor, vehicle (PBS), or negative or positive control in a preventative and/or therapeutic regimen. Preventative treatment starts on Day 0 and continues through the peak of disease in control mice. Therapeutic treatment starts when the majority of mice show mild signs of arthritis. Enbrel®, which has been shown to have good efficacy in both the CIA and G6PI-induced models of arthritis, is used as a positive control. Data collected in every experiment includes clinical scores and cumulative incidence of arthritis. Clinical signs of arthritis in the G6PI model are scored using a scale similar to that employed for the CIA model.

(c) Polycystic Kidney Disease

[00286] The efficacy of an xceptor fusion protein (preferably containing a TNF antagonist and a TGF $\beta$  antagonist, as disclosed herein) in the treatment of polycystic kidney disease is tested in murine models as described in Gattone et al., *Nat. Med.* (2003) 9:1323-6; Torres et al. *Nat. Med.* (2004) 10:363-4; Wang et al. *J. Am. Soc. Nephrol.* (2005) 16:846-851; and Wilson (2008) *Curr. Top. Dev. Biol.* 84:311-50.

[00287] SEQ ID NOS:1-1255 are set out in the attached Sequence Listing. The codes for nucleotide sequences used in the attached Sequence Listing, including the symbol "n," conform to WIPO Standard ST.25 (1998), Appendix 2, Table 1.



CLAIMS

We claim:

1. A multi-specific fusion protein having one of the following structures from amino terminus to carboxy terminus:

- (a) BD-ID-ED;
- (b) ED-ID-BD; or
- (c) ED1-ID-ED2

wherein:

ED is a TGF $\beta$  antagonist and ED1 and ED2 are different antagonists wherein ED1 or ED2 is a TGF $\beta$  antagonist;

ID is an intervening domain; and

BD is a TNF antagonist, IL6 antagonist, IL10 antagonist, VEGF antagonist, HGF antagonist, IGF antagonist, or a GITR agonist.

2. The multi-specific fusion protein of claim 1, wherein the binding domain is an immunoglobulin variable binding domain.

3. The multi-specific fusion protein of claim 1 or 2, wherein the ectodomain is a receptor ligand binding domain.

4. The multi-specific fusion protein of any of the preceding claims, wherein the intervening domain has the following structure:

-L1-CH2CH3-,

wherein:

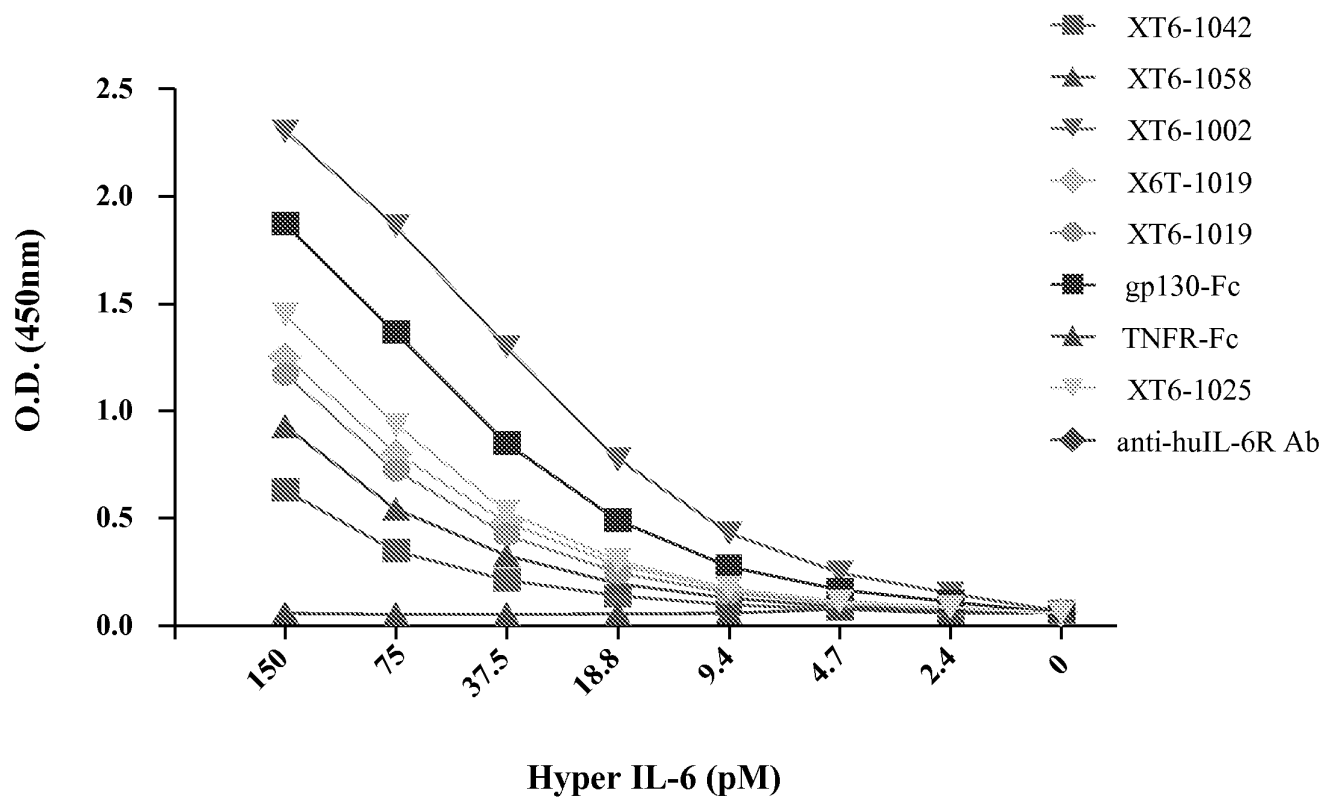
L1 is an immunoglobulin hinge linker, optionally an IgG1 hinge having the first cysteine substituted with a different amino acid;

-CH2CH3- is the CH2CH3 region of an IgG1 Fc domain, optionally mutated to eliminate Fc $\gamma$ RI-III interaction while retaining FcRn interaction.

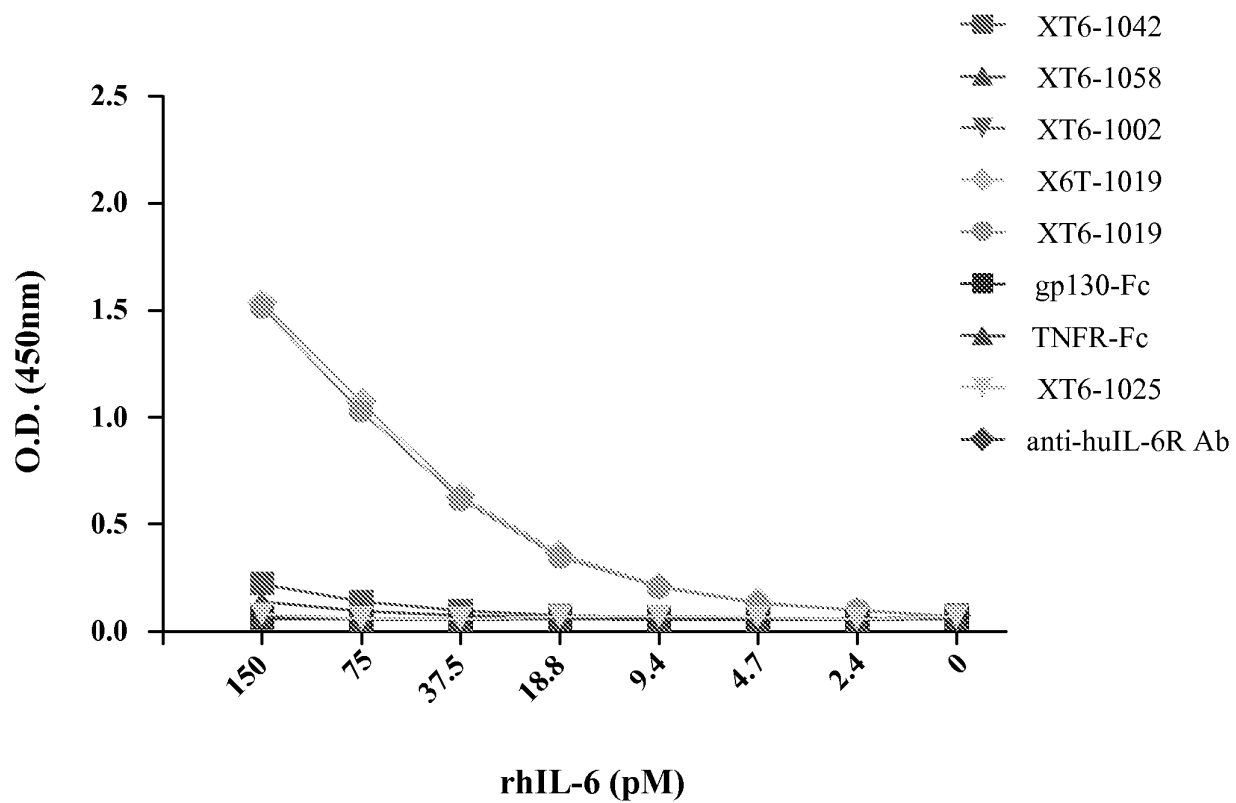
5. The multi-specific fusion protein of any of the preceding claims, wherein the BD is connected to the intervening domain by a first linker and the ED is connected to the intervening domain by a second linker and wherein the first and second linkers may be the same or different.

6. The multi-specific fusion protein of claim 5, wherein the first and second linkers are selected from SEQ ID NO:497-604 and 1223-1228, optionally wherein the first linker is SEQ ID NO:576 and the second linker is SEQ ID NO:1223.
7. The multi-specific fusion protein of any of the preceding claims, comprising an amino acid sequence as set forth in any one of SEQ ID NOS:735-742.
8. A composition comprising one or more multi-specific fusion proteins according to any of the preceding claims and a pharmaceutically acceptable carrier, diluent, or excipient.
9. A composition of claim 8 wherein the multi-specific fusion protein exists as a dimer or a multimer in the composition.
10. A polynucleotide encoding a multi-specific fusion protein according to any one of claims 1-7.
11. An expression vector comprising a polynucleotide according to claim 10 operably linked to an expression control sequence.
12. A host cell comprising an expression vector according to claim 11.
13. A method for treating a subject with a malignant condition comprising the administration of a therapeutically effective amount of a multi-specific fusion protein or composition thereof of any of the preceding claims.
14. The method of claim 13 wherein the malignant condition is breast cancer, renal cell carcinoma, melanoma or prostate cancer.

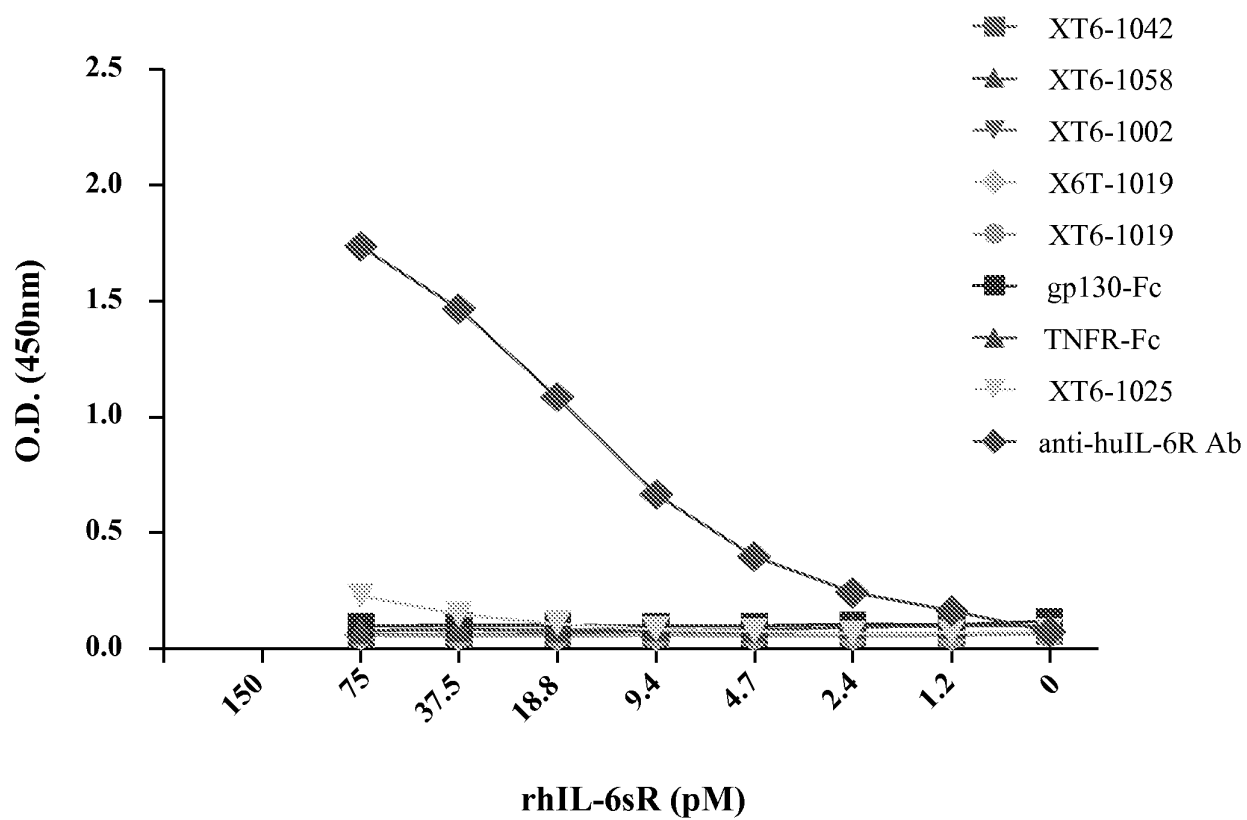
1/16

*Fig. 1A*

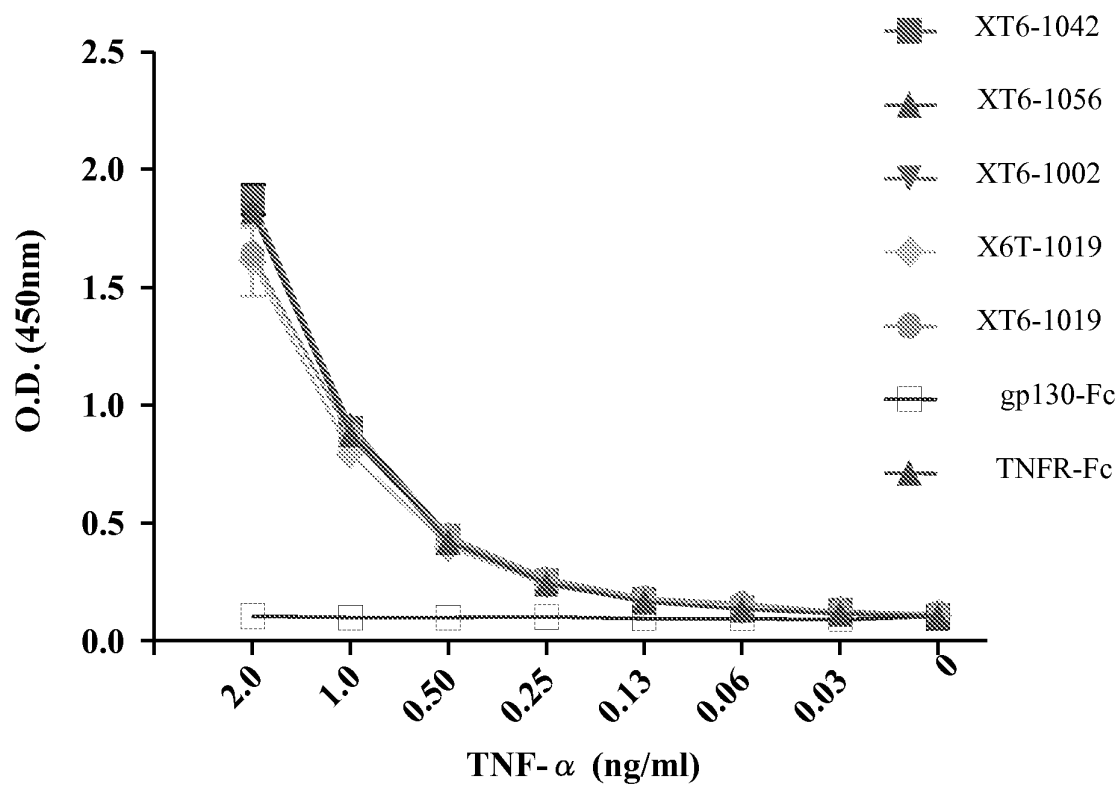
2/16

***Fig. 1B***

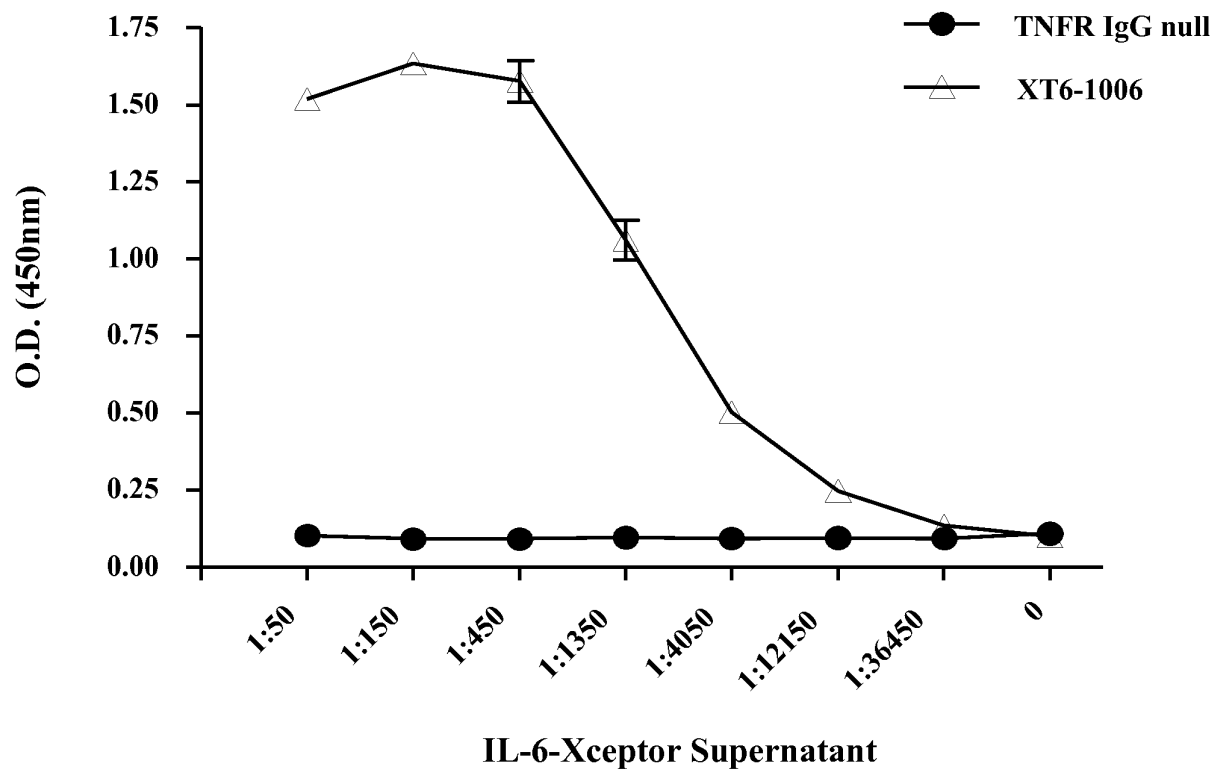
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*Fig. 1C*

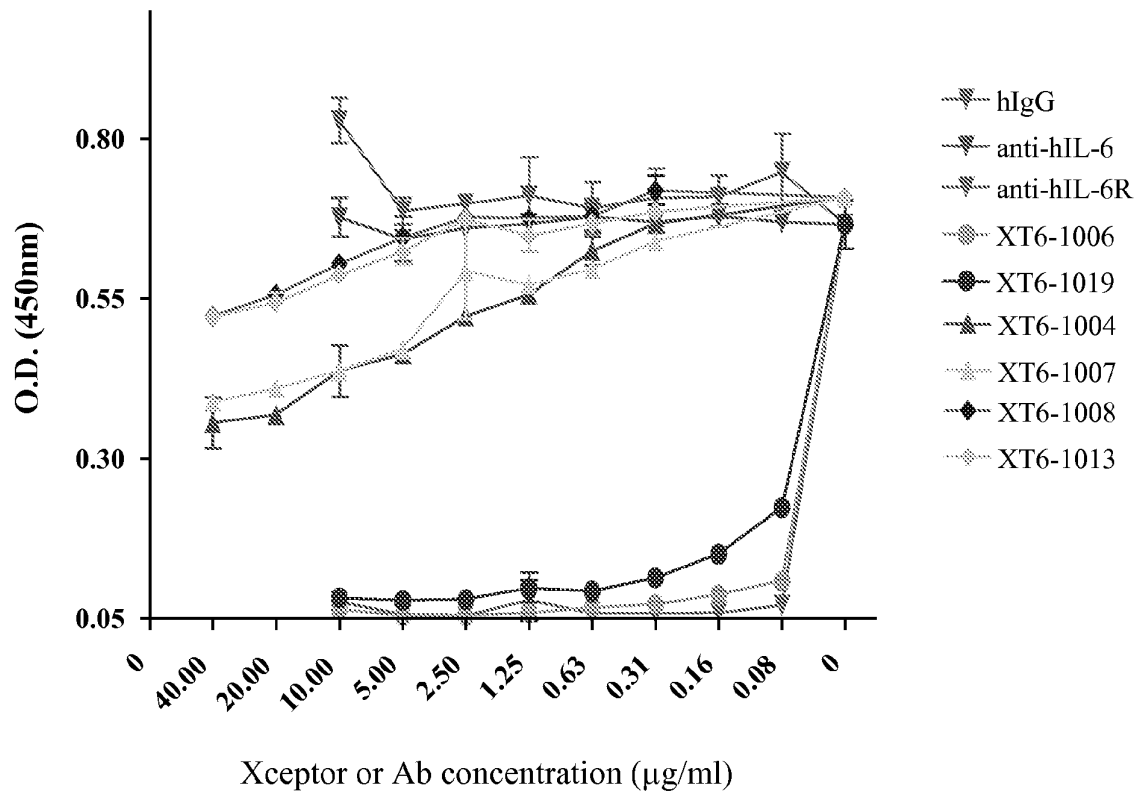
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*Fig. 2*

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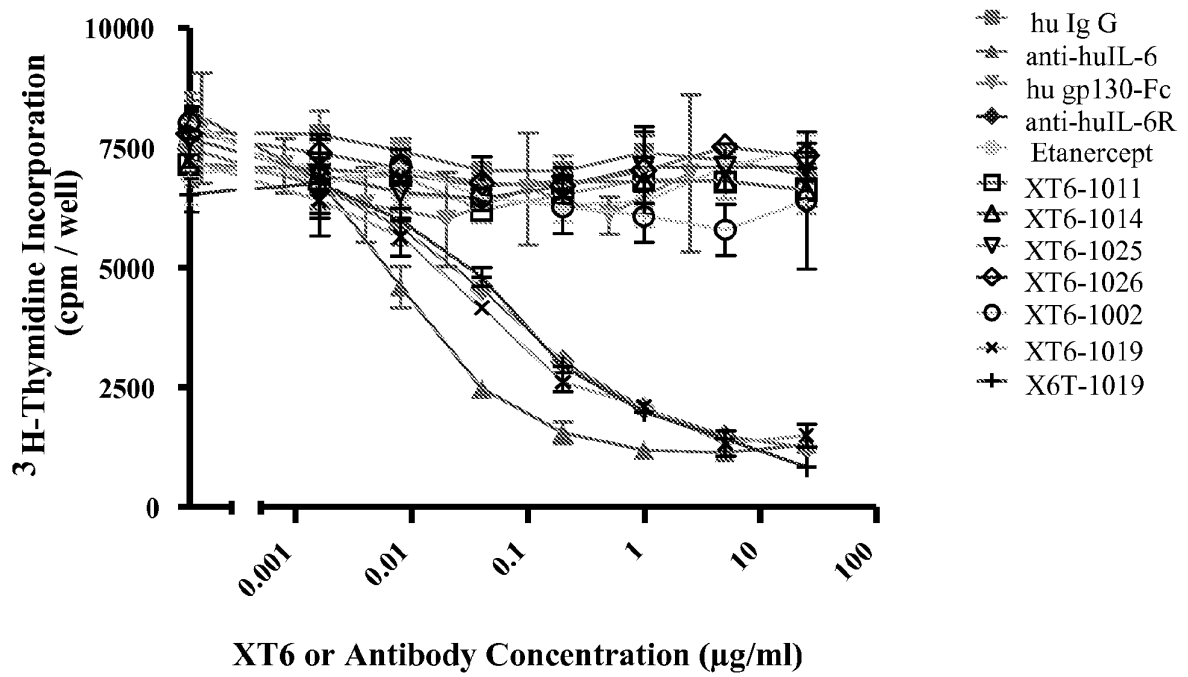
*Fig. 3*

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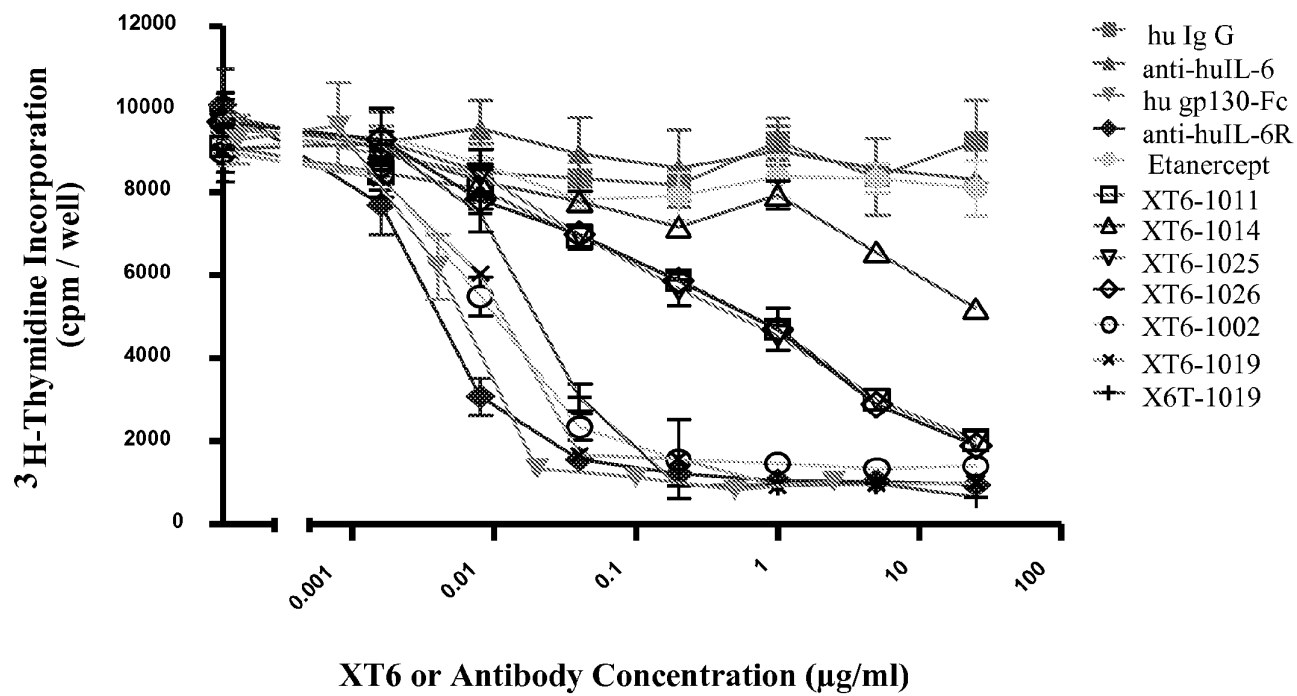
**Fig. 4**



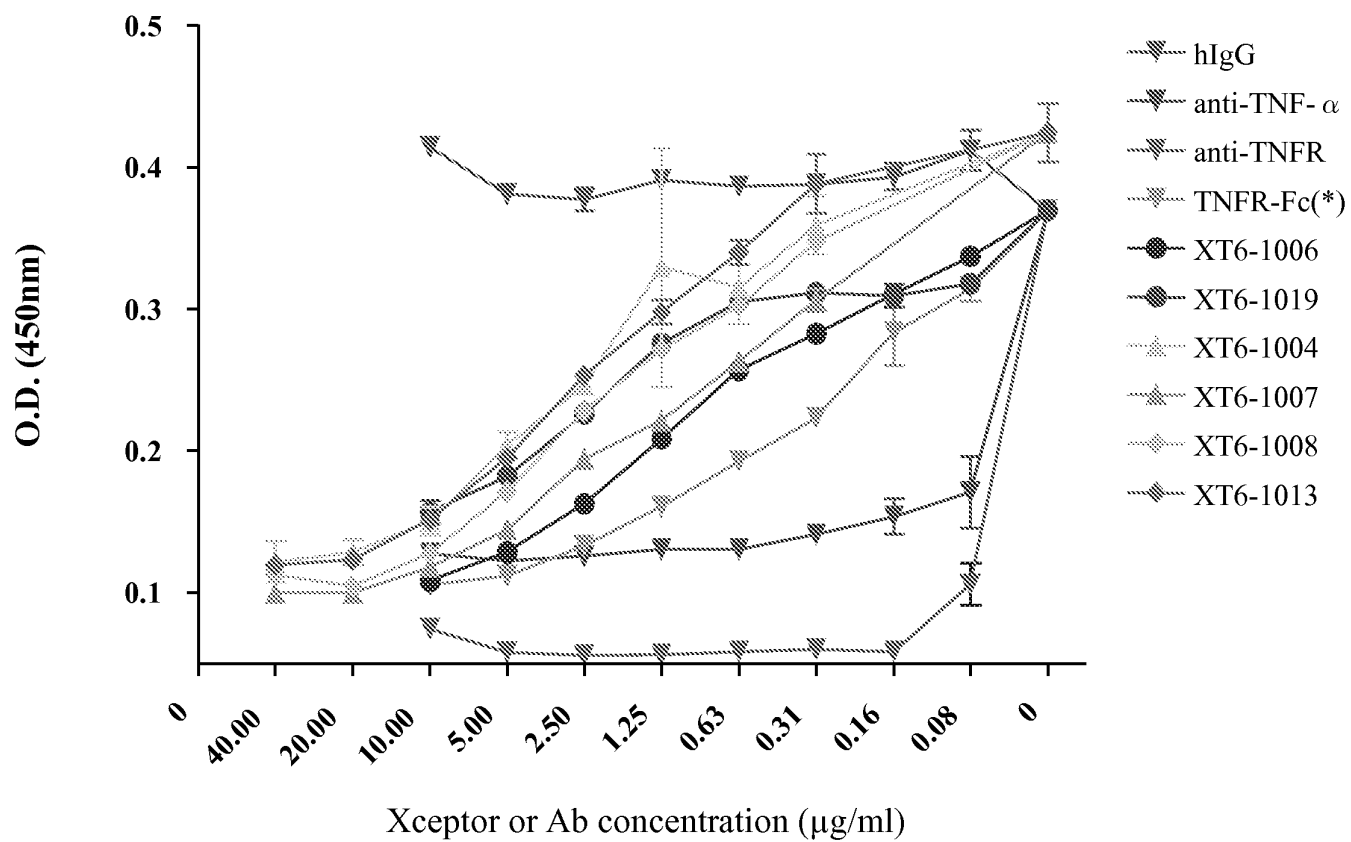
7/16

*Fig. 5A*

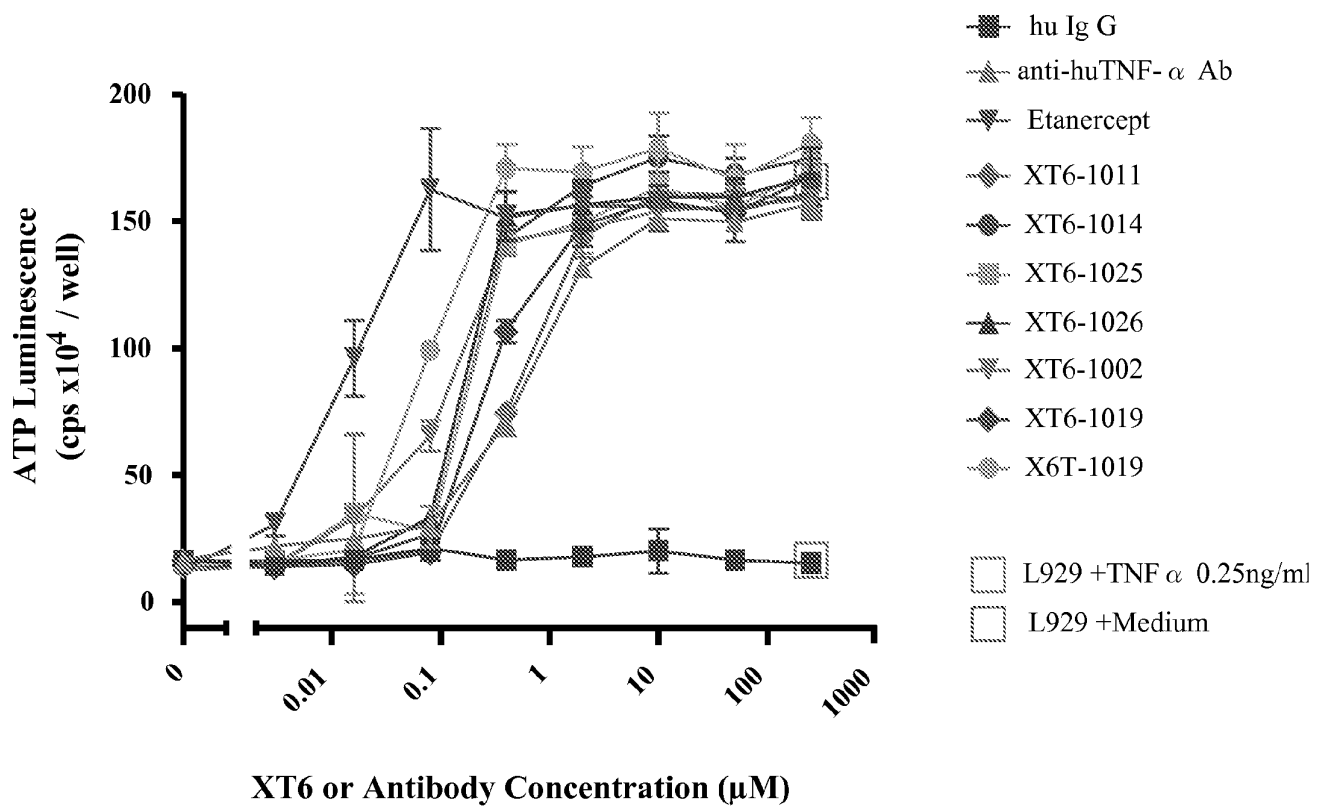
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*Fig. 5B*

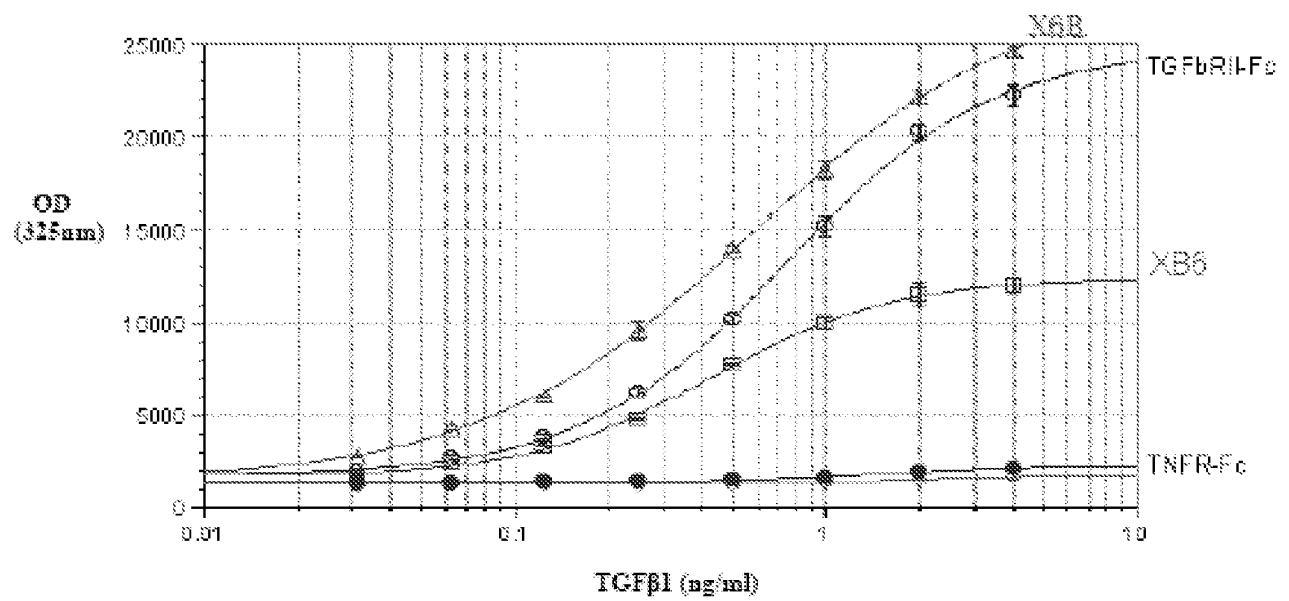
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**Fig. 6**

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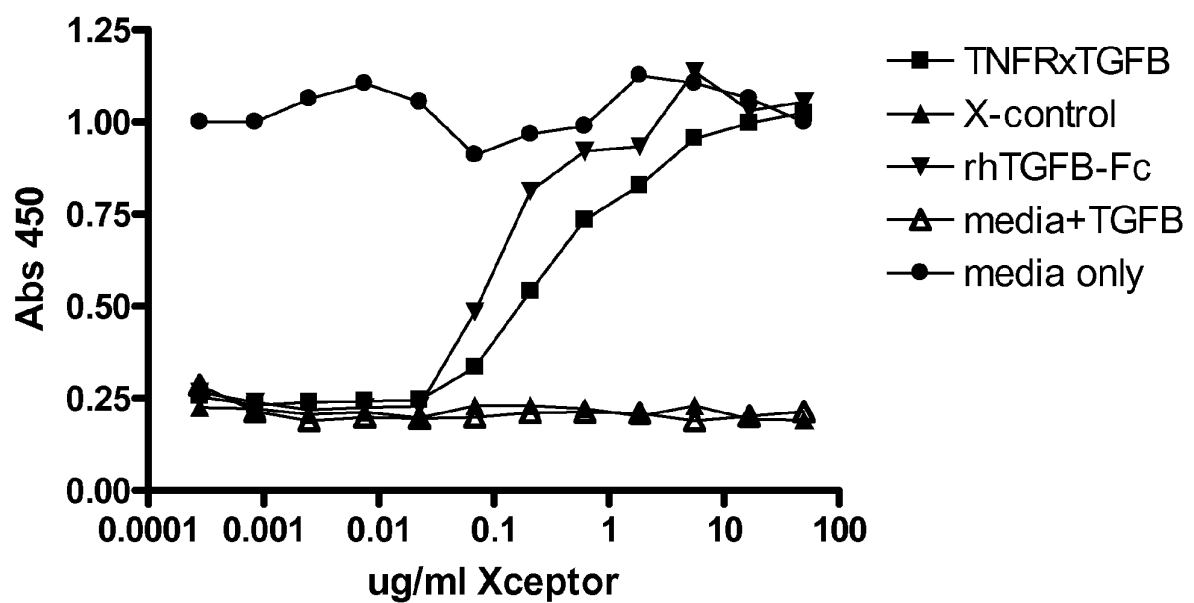
*Fig. 7*

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*Fig. 8*

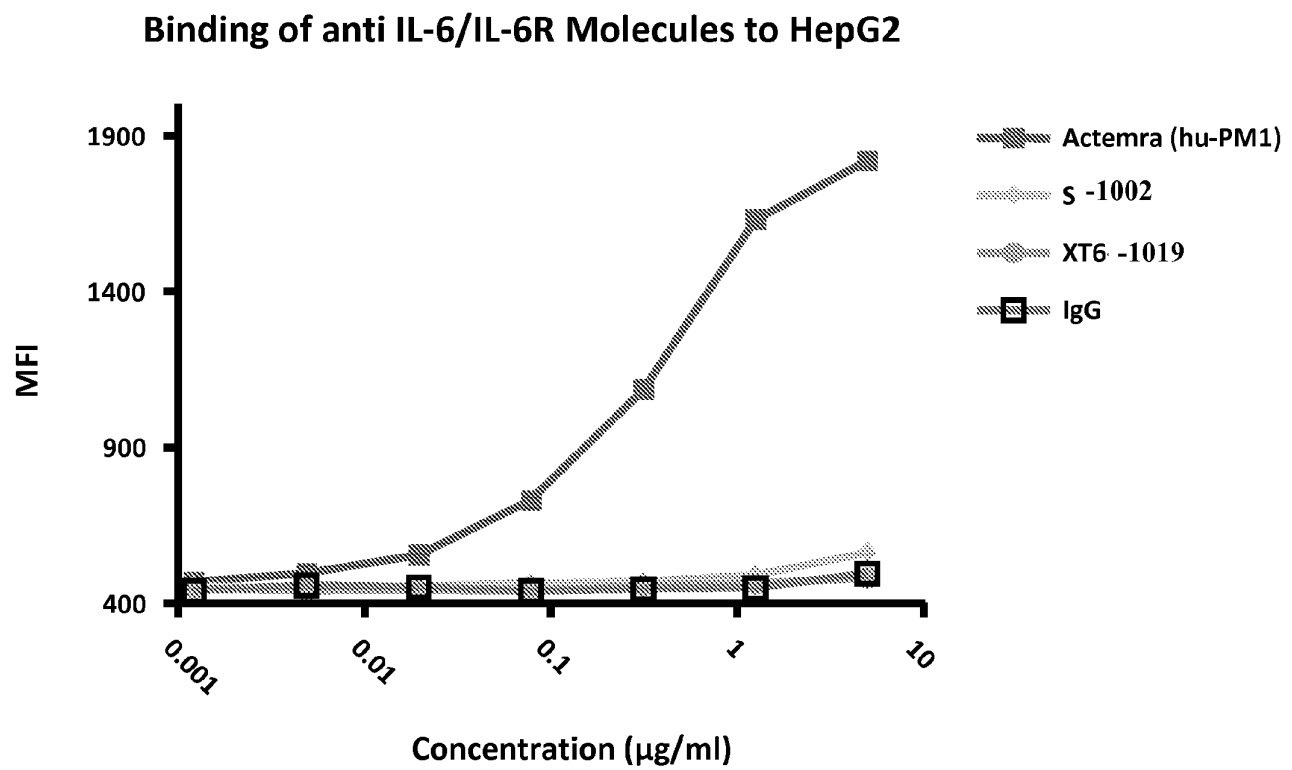
12/16

**TNFRxTGFB $\beta$ RII Xceptor:**  
blocking TGFB-1 inhibition of IL4 proliferation of HT2



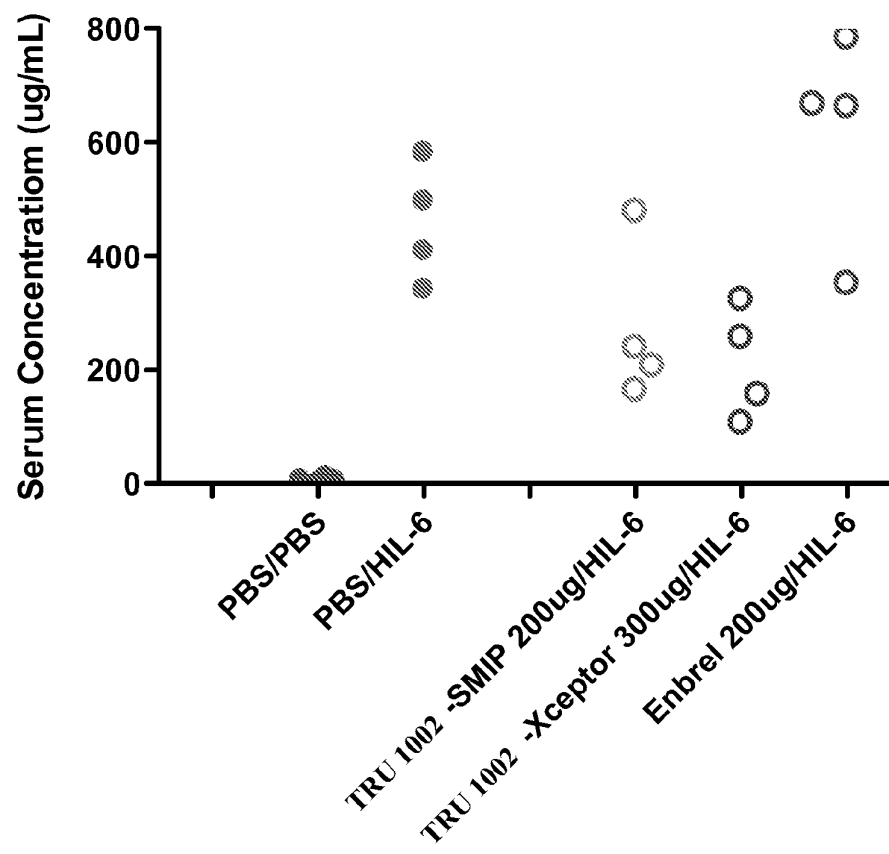
*Fig. 9*

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*Fig. 10*

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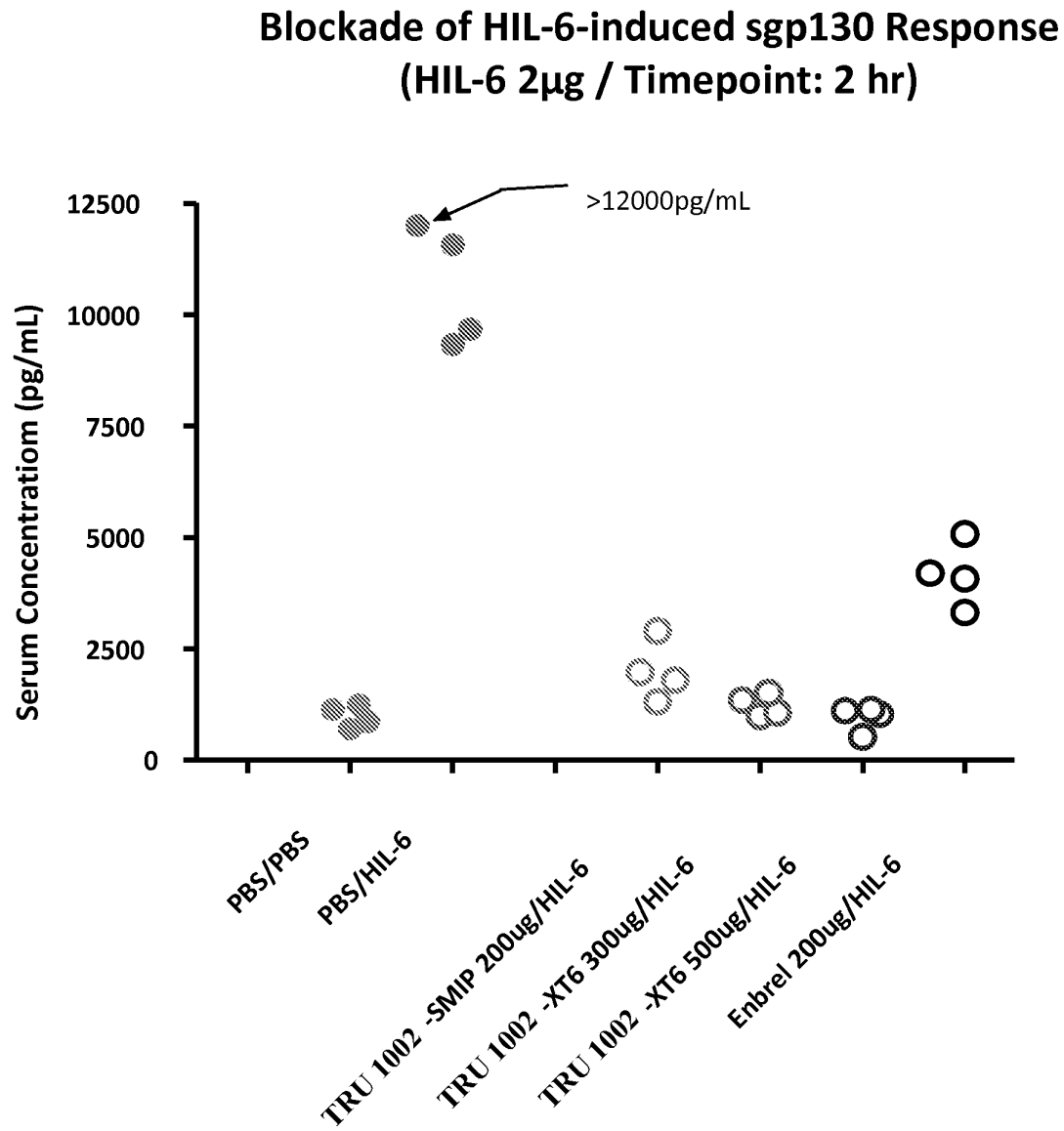
**Blockade of HIL-6-induced Mouse SAA Response**  
**HIL-6 2ug /Timepoint: 24 hr**



***Fig. 11***

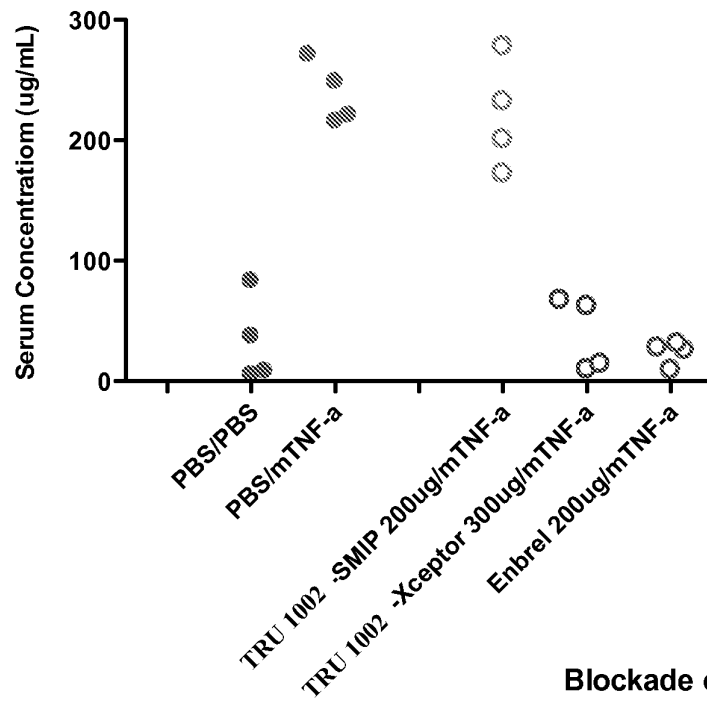


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*Fig. 12*

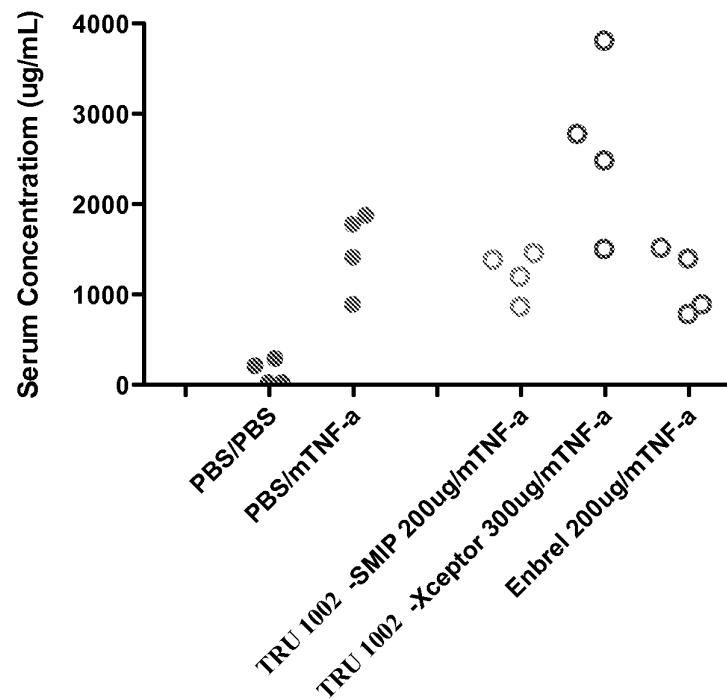
16/16

**Blockade of mTNFa-induced Mouse SAA Response**  
(mTNFa 0.5ug /Timepoint: 2 hr)



*Fig. 13A*

**Blockade of mTNF-a-induced Mouse SAA Response**  
(mTNF-a 0.5ug /Timepoint: 24 hr)



*Fig. 13B*

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<400> 198

Asp His Tyr Gly Asp Tyr Ala Phe Asp Tyr  
1 5 10

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<400> 199

Leu Tyr Glu Met Val  
1 5

<210> 200

<211> 17

<212> PRT

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<220>

<223> Made in a lab

<400> 200

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

<210> 201

<211> 17

<212> PRT

<213> Artificial Sequence

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<223> Made in a lab

<400> 201

Asp Leu Ala Val Ala Gly Trp Asp Tyr Tyr Tyr Tyr Tyr Gly Met Asp  
1 5 10 15  
Val

<210> 202

<211> 5

<212> PRT

<213> Artificial Sequence

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<223> Made in a lab

<400> 202

Gln Tyr Asn Met Ile  
1 5

<210> 203

<211> 16  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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 Trp Ile Ser Ser Ser Gly Gly Thr Glu Tyr Ala Asp Ser Val Lys Gly  
 1 5 10 15

<210> 204  
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 <212> PRT  
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<400> 204  
 Asp Arg Gly Tyr Gly Ser Gly Ser Tyr Gly Ala Tyr Asp Ala Phe Asp  
 1 5 10 15  
 Ile

<210> 205  
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 <212> PRT  
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<400> 205  
 His Tyr Ser Met Gly  
 1 5

<210> 206  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 206  
 Ser Ile Tyr Pro Ser Gly Gly Ser Thr Leu Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 207  
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 <212> PRT  
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<220>

<223> Made in a lab

<400> 207

Trp Gly Val Gly Ala Thr Phe Asp Tyr  
1 5

<210> 208

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 208

His Tyr Met Met Ala  
1 5

<210> 209

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 209

Tyr Ile Tyr Pro Ser Gly Gly Trp Thr Ser Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 210

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 210

Phe Asp Tyr Thr Ile Gly Phe Asp Phe  
1 5

<210> 211

<211> 5

<212> PRT

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<223> Made in a lab

<400> 211

His Tyr Gly Met Thr  
1 5

<210> 212

<211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 212  
 Gly Ile Arg Ser Ser Gly Gly Val Thr Asn Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 213  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 213  
 Glu Gly Ser Gly Trp Ser Lys Ala Phe Asp Ile  
 1 5 10

<210> 214  
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<220>  
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<400> 214  
 His Tyr Asn Met Arg  
 1 5

<210> 215  
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 <212> PRT  
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<220>  
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<400> 215  
 Ser Ile Ser Pro Ser Gly Gly Phe Thr Gly Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 216  
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<220>



<223> Made in a lab

<400> 216

Gly Ile Asn Tyr Tyr Asp Ser Ser Gly Tyr Tyr Pro Pro Val Gly Met  
 1 5 10 15  
 Asp Val

<210> 217

<211> 5

<212> PRT

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<400> 217

Trp Tyr Asn Met Leu  
 1 5

<210> 218

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 218

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

<210> 219

<211> 14

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 219

Asp Arg Gly Gly Ser Pro Phe Arg Pro Asp Ala Phe Asp Ile  
 1 5 10

<210> 220

<211> 5

<212> PRT

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<223> Made in a lab

<400> 220

Glu Tyr Asp Met Leu  
 1 5

<210> 221  
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<220>  
 <223> Made in a lab

<400> 221  
 Ser Ile Trp Pro Ser Gly Gly Phe Thr Gln Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 222  
 <211> 16  
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 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 222  
 Asn Tyr Tyr Asp Phe Trp Ser Gly Pro Tyr Tyr Tyr Gly Met Asp Val  
 1 5 10 15

<210> 223  
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 <223> Made in a lab

<400> 223  
 Gly Tyr Val Met Gly  
 1 5

<210> 224  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 224  
 Ser Ile Ser Pro Ser Gly Gly Tyr Thr Leu Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 225  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 225

Ala	Phe	Ser	Thr	Arg	Trp	Tyr	Trp	Gly	Ala	Phe	Asp	Ile
1				5					10			

<210> 226

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 226

Ile	Tyr	His	Met	Asn
1			5	

<210> 227

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 227

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

Gly

<210> 228

<211> 16

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 228

Ser	Tyr	Arg	Ala	Ala	Gly	Trp	Val	Asp	Tyr	Tyr	Tyr	Gly	Met	Asp	Val
1				5					10					15	

<210> 229

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<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 229

Leu	Tyr	Asp	Met	His
1			5	

<210> 230  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 230  
 Arg Ile Tyr Ser Ser Gly Gly Thr Thr Gln Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 231  
 <211> 12  
 <212> PRT  
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<220>  
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<400> 231  
 Asp Pro Gly Tyr Gly Ser His His Ser Phe Asp Tyr  
 1 5 10

<210> 232  
 <211> 5  
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<220>  
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<400> 232  
 Ser Tyr Tyr Met Thr  
 1 5

<210> 233  
 <211> 17  
 <212> PRT  
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<220>  
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<400> 233  
 Gly Ile Tyr Ser Ser Gly Gly Pro Thr Gln Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 234  
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 <212> PRT  
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<220>

<223> Made in a lab

<400> 234

Ala Gly Gly Asp Ala Phe Asp Ile

1

5

<210> 235

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 235

Ser Tyr Glu Met Phe

1

5

<210> 236

<211> 17

<212> PRT

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<220>

<223> Made in a lab

<400> 236

Ser Ile Ser Pro Ser Gly Gly Tyr Thr Tyr Tyr Ala Asp Ser Val Lys

1

5

10

15

Gly

<210> 237

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 237

Met Thr Thr Ser Gly Phe His Leu Ile

1

5

<210> 238

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 238

Asp Tyr Pro Met Gln

1

5

<210> 239  
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 <212> PRT  
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<220>  
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<400> 239  
 Trp Ile Gly Pro Ser Gly Gly Trp Thr Val Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 240  
 <211> 8  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 240  
 Asp Asp Gly Ile Ala Gly Phe Leu  
 1 5

<210> 241  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 241  
 Trp Tyr Leu Met His  
 1 5

<210> 242  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 242  
 Gly Ile Trp Pro Ser Gly Gly His Thr Leu Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 243  
 <211> 16  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 243

Glu	Pro	Leu	Leu	Trp	Phe	Gly	Glu	Leu	Ser	Tyr	Asn	Trp	Phe	Asp	Pro
1				5					10					15	

<210> 244

<211> 6

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 244

Tyr	Tyr	Glu	Asn	Met	Ala
1				5	

<210> 245

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 245

Gly	Ile	Tyr	Pro	Ser	Gly	Gly	Leu	Thr	Tyr	Tyr	Ala	Asp	Ser	Val	Lys
1				5					10					15	

Gly

<210> 246

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 246

Ser	Arg	Arg	Tyr	Tyr	Asp	Ser	Ser	Asp	Ala	Phe	Asp	Ile
1				5					10			

<210> 247

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 247

Thr	Tyr	Val	Met	Tyr
1				5

<210> 248  
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 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 248  
 Gly Ile Gly Pro Ser Gly Gly Asn Thr Phe Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 249  
 <211> 14  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 249  
 Asp Pro Gly Asp Phe Trp Ser Gly Tyr Tyr Gly Met Asp Val  
 1 5 10

<210> 250  
 <211> 5  
 <212> PRT  
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<220>  
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<400> 250  
 Phe Tyr Gly Met Ala  
 1 5

<210> 251  
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 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 251  
 Ser Ile Tyr Pro Ser Gly Gly Tyr Thr Asp Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 252  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence



<220>

<223> Made in a lab

<400> 252

Ser Ala Gly Gly Trp Ile Gly Gly Gly Ala Phe Asp Ile  
1 5 10

<210> 253

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 253

Lys Tyr Pro Met Met  
1 5

<210> 254

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 254

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

<210> 255

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 255

Tyr Gly Ser Gly Ser Tyr Tyr Leu Tyr Tyr Tyr Tyr Tyr Tyr Met Asp  
1 5 10 15  
Val

<210> 256

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 256

Trp Tyr Ser Met Trp  
1 5

<210> 257  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 257  
Tyr Ile Val Pro Ser Gly Gly Glu Thr Ile Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 258  
<211> 16  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 258  
Asn Leu Gly Glu Gly Phe Trp Ser Asp Tyr Tyr Pro Pro Leu Asp Tyr  
1 5 10 15

<210> 259  
<211> 5  
<212> PRT  
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<220>  
<223> Made in a lab

<400> 259  
His Tyr Gly Met His  
1 5

<210> 260  
<211> 15  
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<213> Artificial Sequence

<220>  
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<400> 260  
Ser Ile Tyr Pro Gly Met Thr Gln Tyr Ala Asp Ser Val Lys Gly  
1 5 10 15

<210> 261  
<211> 11  
<212> PRT  
<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 261

Asp Arg Gly Ser Gly Ile Asp Ala Phe Asp Ile  
1 5 10

<210> 262

<211> 5

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<220>

<223> Made in a lab

<400> 262

Trp Tyr Asp Met Leu  
1 5

<210> 263

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 263

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

<210> 264

<211> 16

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 264

Thr Arg Ser Met Tyr Ser Ser Ser Trp Tyr Gly Ala Pro Pro Thr His  
1 5 10 15

<210> 265

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 265

Trp Tyr Lys Met His  
1 5

<210> 266  
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 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 266  
 Gly Ile Ser Ser Ser Gly Gly Leu Thr Lys Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 267  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 267  
 Glu Arg Arg Gly Asp Gly Gly Ala Phe Asp Ile  
 1 5 10

<210> 268  
 <211> 5  
 <212> PRT  
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<220>  
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<400> 268  
 Glu Tyr Thr Met Tyr  
 1 5

<210> 269  
 <211> 17  
 <212> PRT  
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<220>  
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<400> 269  
 Tyr Ile Ser Pro Ser Gly Gly Thr Thr Asn Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 270  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 270

Gly Ala Trp Gly Asp Ile Tyr Tyr Tyr Gly Met Asp Val  
1 5 10

<210> 271

<211> 5

<212> PRT

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<220>

<223> Made in a lab

<400> 271

Ser Tyr Trp Met His  
1 5

<210> 272

<211> 17

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<220>

<223> Made in a lab

<400> 272

Ser Ile Tyr Pro Ser Gly Gly Ser Thr Glu Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 273

<211> 6

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 273

Gly Ile Trp Phe Asp Pro  
1 5

<210> 274

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 274

Arg Tyr Gly Met Met  
1 5

<210> 275  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 275  
 Tyr Ile Ser Ser Ser Gly Gly Phe Thr Arg Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 276  
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 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 276  
 Val Gly Gly Tyr Ser Tyr Gly Pro His Phe Asp Phe  
 1 5 10

<210> 277  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 277  
 Trp Tyr His Met Ile  
 1 5

<210> 278  
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 <213> Artificial Sequence

<220>  
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<400> 278  
 Trp Ile Ser Pro Ser Gly Gly Phe Thr Lys Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 279  
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 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 279

Tyr Asp Ser Arg Ala Ala Ala Gly Thr Asn Ala Phe Asp Ile  
1 5 10

<210> 280

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 280

Pro Tyr Lys Met Val  
1 5

<210> 281

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 281

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

<210> 282

<211> 14

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 282

Gly Gly Tyr Gly Trp Ser Tyr Tyr Tyr Tyr Gly Met Asp Val  
1 5 10

<210> 283

<211> 5

<212> PRT

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<220>

<223> Made in a lab

<400> 283

Asn Tyr Trp Met Tyr  
1 5

<210> 284  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 284  
 Val Ile Ser Ser Ser Gly Gly His Thr Phe Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 285  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 285  
 Asp Tyr Glu Gly Gly Ser Asn Asp Tyr  
 1 5

<210> 286  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 286  
 Asn Tyr Ala Met Ser  
 1 5

<210> 287  
 <211> 17  
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 <213> Artificial Sequence

<220>  
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<400> 287  
 Ser Ile Tyr Ser Ser Gly Gly Tyr Thr Ala Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 288  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence



<220>

<223> Made in a lab

<400> 288

Val Pro His Val Phe Arg Gly Glu Leu Asp Tyr  
1 5 10

<210> 289

<211> 5

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<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 289

Phe Tyr Thr Met Trp  
1 5

<210> 290

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 290

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

<210> 291

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 291

Pro Asp Ser Tyr Gly Tyr Leu Tyr Tyr Gly Met Asp Val  
1 5 10

<210> 292

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 292

Trp Tyr Pro Met Glu  
1 5

<210> 293  
 <211> 17  
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<220>  
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<400> 293  
 Gly Ile Gly Pro Ser Gly Gly Gln Thr Thr Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 294  
 <211> 6  
 <212> PRT  
 <213> Artificial Sequence

<220>  
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<400> 294  
 Gly Ser Tyr Ser Phe Ile  
 1 5

<210> 295  
 <211> 5  
 <212> PRT  
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<220>  
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<400> 295  
 Leu Tyr Lys Met Ala  
 1 5

<210> 296  
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 <212> PRT  
 <213> Artificial Sequence

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<400> 296  
 Val Ile Gly Ser Ser Gly Gly Arg Thr Pro Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 297  
 <211> 16  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 297

Ala	Pro	Leu	Ser	Gly	Trp	Phe	Gly	Gln	Ala	His	Asp	Ala	Phe	Asp	Ile
1				5				10						15	

<210> 298

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 298

Thr	Tyr	Glu	Met	Asn
1			5	

<210> 299

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 299

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

Gly

<210> 300

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 300

Gly	Pro	Tyr	Ser	Tyr	Gly	His	Asp	Tyr	Gly	Met	Asp	Val
1				5				10				

<210> 301

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 301

Val	Tyr	Pro	Met	His
1			5	

<210> 302  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 302  
 Ser Ile Tyr Ser Ser Gly Gly Phe Thr Met Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 303  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 303  
 Glu Gly Val Ala Asp Ala Phe Asp Ile  
 1 5

<210> 304  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 304  
 Ser Tyr Thr Met His  
 1 5

<210> 305  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 305  
 Ser Ile Ser Pro Ser Gly Gly Met Thr Phe Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 306  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 306

Thr	Tyr	Asp	Phe	Trp	Ser	Gly	Tyr	Phe	Asp	Tyr
1				5					10	

<210> 307

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 307

Thr	Tyr	Glu	Met	Gly
1			5	

<210> 308

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 308

Val	Ile	Trp	Ser	Ser	Gly	Gly	His	Thr	Trp	Tyr	Ala	Asp	Ser	Val	Lys
1				5					10					15	

Gly

<210> 309

<211> 16

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 309

Ser	Asn	Gln	Gly	Asp	Phe	Trp	Ser	Gly	Tyr	Pro	Phe	Ala	Phe	Asp	Ile
1				5					10					15	

<210> 310

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 310

Asn	Tyr	Asn	Met	Tyr
1			5	

<210> 311  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 311  
 Tyr Ile Ser Pro Ser Gly Gly Met Thr Lys Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 312  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 312  
 Arg Gly Val Leu Gly Tyr Tyr Gly Met Asp Val  
 1 5 10

<210> 313  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 313  
 Leu Tyr Ser Met Asn  
 1 5

<210> 314  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 314  
 Ser Ile Ser Ser Ser Gly Gly Ala Thr Leu Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 315  
 <211> 14  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 315

Asp Leu Ile Ser Ser Gly Tyr Tyr Pro Asp Ala Phe Asp Ile  
1 5 10

<210> 316

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 316

Gln Tyr Met Met His  
1 5

<210> 317

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 317

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

<210> 318

<211> 19

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 318

Pro Leu Ala Leu Gly Tyr Asp Phe Trp Ser Gly Tyr Gln Ala Ala Gly  
1 5 10 15  
Phe Asp Tyr

<210> 319

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 319

Trp Tyr Ser Met Val  
1 5

<210> 320  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 320  
Ser Ile Val Pro Ser Gly Gly Leu Thr Trp Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 321  
<211> 10  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 321  
Asp Val Tyr Gly Asp Tyr Tyr Phe Asp Asn  
1 5 10

<210> 322  
<211> 5  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 322  
Tyr Tyr Asn Met Thr  
1 5

<210> 323  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 323  
Ser Ile Ser Pro Ser Gly Gly Arg Thr Gly Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 324  
<211> 10



<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 324  
 Val Ser Gly Phe Tyr Ala Ala Phe Asp Tyr  
 1 5 10

<210> 325  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 325  
 Trp Tyr Trp Met Thr  
 1 5

<210> 326  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 326  
 Ser Ile Ser Pro Ser Gly Gly His Thr Ser Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 327  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 327  
 Ser Ser Gly Trp Tyr Glu Asp Tyr Phe Asp Tyr  
 1 5 10

<210> 328  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 328

Pro Tyr Glu Met Ala  
1 5

<210> 329  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 329  
Val Ile Gly Pro Ser Gly Gly Tyr Thr Glu Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 330  
<211> 14  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 330  
Gly Tyr Asp Phe Trp Ser Gly Tyr Tyr Asp Ala Phe Asp Ile  
1 5 10

<210> 331  
<211> 5  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 331  
Ser Tyr Phe Met Gly  
1 5

<210> 332  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 332  
Ser Ile Trp Pro Ser Gly Gly Asn Thr Thr Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 333  
<211> 10

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 333  
 His Val Gly Trp Gly Arg Tyr Phe Asp Tyr  
 1 5 10

<210> 334  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 334  
 Ala Tyr Arg Met Ile  
 1 5

<210> 335  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 335  
 Tyr Ile Ser Ser Ser Gly Gly Arg Thr Asp Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 336  
 <211> 15  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 336  
 Gly Gly Leu Arg Tyr Phe Asp Trp Leu Ala Pro Ser Met Asp Tyr  
 1 5 10 15

<210> 337  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 337

Ala Tyr Ser Met Val  
1 5

<210> 338  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 338  
Tyr Ile Tyr Pro Ser Gly Gly Ile Thr Thr Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 339  
<211> 7  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 339  
Glu Gly Gln Val Phe Asp Ile  
1 5

<210> 340  
<211> 5  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 340  
Asn Tyr Ala Met Gly  
1 5

<210> 341  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 341  
Arg Ile Val Pro Ser Gly Gly Met Thr Ser Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 342  
<211> 9

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 342  
 Asp Asp Phe Trp Ser Gly Met Asp Val  
 1 5

<210> 343  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 343  
 Leu Tyr Asn Met Gln  
 1 5

<210> 344  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 344  
 Gly Ile Ser Leu Ser Gly Gly Lys Thr Phe Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 345  
 <211> 12  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 345  
 Asp Tyr Gly Val Ala Thr Leu Asp Ala Phe Asp Ile  
 1 5 10

<210> 346  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 346

Ala Tyr Thr Met His  
1 5

<210> 347  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 347  
Ser Ile Tyr Pro Ser Gly Gly Thr Thr Pro Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 348  
<211> 18  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 348  
Val Val Asn Ile Asp Phe Trp Ser Gly Tyr Asn Met Arg Ser Ala Phe  
1 5 10 15  
Asp Ile

<210> 349  
<211> 5  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 349  
Asp Tyr Leu Met Gly  
1 5

<210> 350  
<211> 17  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 350  
Val Ile Ser Ser Ser Gly Gly Pro Thr Ala Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 351  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 351  
 Val Gly Leu Asp Tyr Gly Ile Leu Gly Ala Phe Asp Ile  
 1 5 10

<210> 352  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 352  
 Glu Tyr Gly Met Ser  
 1 5

<210> 353  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 353  
 Ser Ile Arg Ser Ser Gly Gly Trp Thr Lys Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 354  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 354  
 Gly Leu Gly Ala Thr Ser Gly Glu Phe Asp Tyr  
 1 5 10

<210> 355  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 355

Lys Tyr Asp Met Trp  
1 5

<210> 356

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 356

Ser Ile Trp Pro Ser Gly Gly Trp Thr Ser Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 357

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 357

Ser Arg Gly Ser Pro Trp Tyr Gly Asp Phe Asp His  
1 5 10

<210> 358

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 358

Met Tyr Thr Met His  
1 5

<210> 359

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 359

Trp Ile Ser Pro Ser Gly Gly Trp Thr Lys Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly



<210> 360  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 360  
 Ser Pro Trp Gly Gly Pro Phe Asp Tyr  
 1 5

<210> 361  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 361  
 Gln Tyr Phe Met Met  
 1 5

<210> 362  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 362  
 Ser Ile Ser Pro Ser Gly Gly Tyr Thr Gln Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 363  
 <211> 14  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 363  
 Thr Gly Asp Tyr Thr Arg Tyr Tyr Ser Tyr Gly Met Asp Val  
 1 5 10

<210> 364  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 364

Gly Tyr Ser Met Ala  
1 5

<210> 365

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 365

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

<210> 366

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 366

Asp Gln Ser Phe Ser Tyr Asp Ser Ser Leu Asp Ala Phe Asp Ile  
1 5 10 15

<210> 367

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 367

Leu Tyr Gly Met His  
1 5

<210> 368

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 368

Ser Ile Val Pro Ser Gly Gly Leu Thr Arg Tyr Ala Asp Ser Val Lys  
1 5 10 15  
Gly

<210> 369  
 <211> 15  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 369  
 Leu Ala Tyr Tyr Asp Phe Trp Ser Gly Arg Asp Ala Phe Asp Ile  
 1 5 10 15

<210> 370  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 370  
 His Tyr Gly Met Asp  
 1 5

<210> 371  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 371  
 Ser Ile Tyr Ser Ser Gly Gly Tyr Thr Leu Tyr Ala Asp Ser Val Lys  
 1 5 10 15  
 Gly

<210> 372  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 372  
 Arg Thr Trp Ala Asp Ala Phe Asp Val  
 1 5

<210> 373  
 <211> 105  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 373

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

&lt;210&gt; 374

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 374

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

&lt;210&gt; 375

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 375

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Leu	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser
			20					25					30		
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Glu	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Ala	Ala	Ser	Ala	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser

50		55		60
Gly Ser Gly Leu Gly Thr Val Phe Thr Leu Thr Ile Thr Ser Leu Gln				
65		70		75
Pro Glu Asp Ser Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ser Pro Pro				80
	85		90	95
Val Thr Phe Gly Gly Gly Thr Lys Val Asp Ile Lys				
100		105		

<210> 376  
 <211> 107  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 376
Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr Leu Ser Ala Ser Val
1 5 10 15
Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser
20 25 30
Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu
35 40 45
Ile Tyr Lys Ala Ser Ser Leu Glu Ser Gly Val Pro Ser Arg Phe Ser
50 55 60
Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln
65 70 75 80
Pro Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ser Tyr Trp
85 90 95
Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
100 105

<210> 377  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 377
Gln Asp Ile Gln Met Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro
1 5 10 15
Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser
20 25 30
Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
35 40 45
Ile Tyr Asp Ala Ser Asn Arg Ala Thr Gly Ile Pro Ala Arg Phe Ser
50 55 60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65 70 75 80
Pro Glu Asp Ser Ala Val Tyr Tyr Cys Gln Gln Arg Ser Asn Trp Pro
85 90 95
Pro Phe Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
100 105

<210> 378

<211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 378  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Gln Thr Ser Gln Ser Ile Asp Arg  
 20 25 30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Ala Gly Glu Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Phe Ala Ser Tyr Tyr Cys Gln Gln Ser Tyr Arg Thr Pro  
 85 90 95  
 Arg Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 379  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 379  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro  
 1 5 10 15  
 Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser  
 20 25 30  
 Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu  
 35 40 45  
 Leu Ile Tyr Gly Ala Ser Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe  
 50 55 60  
 Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu  
 65 70 75 80  
 Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser  
 85 90 95  
 Pro Leu Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
 100 105 110

<210> 380  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 380  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val

1				5					10					15			
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser		
			20					25					30				
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu		
		35					40					45					
Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser		
	50					55					60						
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln		
65					70					75					80		
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Thr	Pro		
			85					90						95			
Arg	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys						
			100					105									

<210> 381  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 381																	
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ala	Thr	Leu	Ser	Leu	Ser	Pro		
1				5				10					15				
Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Thr		
		20					25					30					
Ser	Leu	Ala	Trp	Tyr	Gln	Gln	Ile	Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu		
		35				40					45						
Met	Tyr	Asp	Ala	Ser	Lys	Arg	Ala	Ser	Gly	Ile	Pro	Ala	Arg	Phe	Ser		
	50				55						60						
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Glu		
65					70					75					80		
Pro	Glu	Asp	Ser	Ala	Val	Tyr	Tyr	Cys	Gln	Leu	Arg	Ile	Asn	Trp	Pro		
			85					90						95			
Pro	Glu	Phe	Thr	Phe	Gly	Pro	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			100					105					110				

<210> 382  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 382																	
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala	Ser	Val		
1				5				10					15				
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Arg	Gly	Val	Ile	Thr		
		20					25					30					
Trp	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Arg	Val	Pro	Ser	Pro	Leu		
		35				40					45						
Ile	Phe	Gly	Ala	Ser	Thr	Leu	Gln	Thr	Gly	Val	Pro	Ser	Arg	Phe	Ser		
	50				55					60							
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Asn	Asp	Leu	Gln		
65					70					75					80		

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Thr His Ser Phe Pro  
                   85                  90                  95  
 Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys  
                   100                  105

<210> 383  
 <211> 106  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 383  
 Gln Ser Ala Leu Thr Gln Pro Pro Ser Val Ser Val Ser Pro Gly Gln  
   1                  5                  10                  15  
 Thr Ala Arg Ile Thr Cys Ser Gly Asp Lys Leu Gly Glu Lys Tyr Val  
                   20                  25                  30  
 Ser Trp Tyr Gln Gln Arg Ser Gly Gln Ser Pro Leu Leu Val Ile Ser  
                   35                  40                  45  
 Gln Asp Thr Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser  
                   50                  55                  60  
 Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Ala Gln Ala Ala  
  65                  70                  75                  80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Thr Trp Asp Ser Asp Thr Tyr Val  
                   85                  90                  95  
 Phe Gly Ser Gly Ser Thr Val Thr Val Leu  
                   100                  105

<210> 384  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 384  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
   1                  5                  10                  15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser  
                   20                  25                  30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
                   35                  40                  45  
 Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
                   50                  55                  60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln  
  65                  70                  75                  80  
 Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ser Thr Leu  
                   85                  90                  95  
 Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
                   100                  105

<210> 385  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence



&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 385

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

&lt;210&gt; 386

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 386

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

&lt;210&gt; 387

&lt;211&gt; 109

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 387

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Thr	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Met	Ser	Asn
			20					25					30		

Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
           35                                  40                                  45  
 Ile Tyr Asp Val Phe Thr Leu Lys Ser Gly Val Pro Ser Arg Phe Ser  
           50                                  55                                  60  
 Gly Ser Arg Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65                                  70                                  75                                  80  
 Pro Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Ser Asp Tyr Ser  
                                   85                                  90                                  95  
 Gly Ile Thr Phe Gly Pro Gly Thr Glu Val Asp Ile Arg  
                                   100                                  105

<210> 388

<211> 111

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 388

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

<210> 389

<211> 108

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 389

Gln Asp Ile Gln Met Thr Gln Ser Pro Ala Thr Leu Ser Ala Ser Val  
 1                                  5                                  10                                  15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser  
                                   20                                  25                                  30  
 Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
           35                                  40                                  45  
 Ile Tyr Lys Ala Ser Gly Leu Glu Ser Gly Val Pro Ser Arg Phe Ser  
           50                                  55                                  60  
 Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65                                  70                                  75                                  80  
 Pro Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ser Tyr Pro  
                                   85                                  90                                  95  
 Phe Thr Phe Gly Pro Gly Thr Lys Val Asp Ile Lys

100

105

<210> 390  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 390  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Leu Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Gln Ala Ser Arg Asp Ile Ser Ile  
 20 25 30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Val Pro Lys Leu Leu  
 35 40 45  
 Ile Phe Gly Ala Ser Thr Leu Arg Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Thr Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Val Ala Thr Tyr Tyr Cys Gln Lys Tyr Asn Ser Ala Pro  
 85 90 95  
 Tyr Thr Phe Gly Pro Gly Thr Lys Val Glu Ile Arg  
 100 105

<210> 391  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 391  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Val Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser  
 20 25 30  
 Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Ser Phe Pro  
 85 90 95  
 Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 392  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 392

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

&lt;210&gt; 393

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 393

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

&lt;210&gt; 394

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 394

Gln	Ser	Glu	Leu	Thr	Gln	Pro	Pro	Ser	Ala	Ser	Gly	Thr	Pro	Gly	Gln
1				5					10					15	
Arg	Val	Thr	Ile	Ser	Cys	Ser	Gly	Ser	Ser	Ser	Asn	Ile	Gly	Ser	Asn
			20					25					30		
Thr	Val	Asn	Trp	Tyr	Gln	Gln	Leu	Pro	Gly	Thr	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Ser	Asn	Asn	Gln	Arg	Pro	Ser	Gly	Val	Pro	Asp	Arg	Phe	Ser

50		55		60
Gly Ser Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln				
65		70		75
Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu				
	85		90	95
Asn Gly Tyr Val Phe Gly Ile Gly Thr Lys Val Thr Val Leu				
	100		105	110

<210> 395  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 395
Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1 5 10 15
Gly Asp Ser Val Thr Ile Ala Cys Arg Ala Ser Gln Ser Ile Ser Thr
20 25 30
Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Lys Leu Leu
35 40 45
Ile Tyr Glu Ala Thr Asn Leu Gln Ser Gly Val Pro Ser Arg Phe Ser
50 55 60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln
65 70 75 80
Pro Glu Asp Val Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Asn Phe Pro
85 90 95
Phe Thr Phe Gly Pro Gly Thr Lys Val Asp Ile Lys
100 105

<210> 396  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 396
Gln Ser Val Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln
1 5 10 15
Arg Val Thr Ile Ser Cys Ser Gly Ser Ser Ser Asn Ile Gly Gly Asn
20 25 30
Ala Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Val
35 40 45
Leu Tyr Ser Ser Asn His Arg Pro Ser Gly Val Pro Asp Arg Phe Ser
50 55 60
Gly Ser Arg Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln
65 70 75 80
Ser Gln Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95
Asn Gly Pro Leu Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

<210> 397

<211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 397  
 Gln Ser Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ser Pro Gly Gln  
 1 5 10 15  
 Thr Ala Thr Ile Ser Cys Ser Gly Glu Lys Leu Gly Asp Arg Tyr Ala  
 20 25 30  
 Trp Trp Tyr Gln Gln Lys Ala Gly Gln Ser Pro Val Leu Val Ile Tyr  
 35 40 45  
 Gln Asp Thr Lys Arg Pro Ser Gly Val Pro Asp Arg Phe Ser Gly Ser  
 50 55 60  
 Asn Ser Gly Thr Ser Ala Ser Leu Val Ile Ser Gly Leu Gln Ser Glu  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu Lys Gly  
 85 90 95  
 Val Val Phe Gly Gly Gly Ser Lys Leu Thr Val Leu  
 100 105

<210> 398  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 398  
 Gln Ser Glu Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln  
 1 5 10 15  
 Arg Val Thr Ile Ser Cys Ser Gly Arg Ser Ser Asn Ile Gly Ser Asn  
 20 25 30  
 Ser Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Ser Asn Asn His Arg Pro Ser Gly Val Pro Asp Arg Phe Ser  
 50 55 60  
 Gly Ser Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln  
 65 70 75 80  
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu  
 85 90 95  
 Tyr Gln Gly Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu  
 100 105 110

<210> 399  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 399  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val

1				5					10					15			
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser		
			20					25					30				
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu		
		35					40					45					
Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Lys	Phe	Ser		
	50					55					60						
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Leu		
65					70					75					80		
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Glu	Tyr	Phe	Pro		
			85						90					95			
Pro	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys						
			100					105									

<210> 400  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 400																	
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Asp	Ser	Leu	Ser	Leu	Ser	Pro		
1				5				10					15				
Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys	Arg	Ala	Ser	Gln	Ser	Val	Ser	Ser		
		20					25					30					
Ser	Tyr	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Arg	Leu		
		35				40					45						
Leu	Ile	Tyr	Gly	Ala	Ser	Ser	Arg	Ala	Thr	Gly	Ile	Pro	Asp	Arg	Phe		
	50					55					60						
Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Arg	Leu		
65					70					75					80		
Glu	Pro	Glu	Asp	Phe	Ala	Val	Tyr	Tyr	Cys	Gln	Gln	Tyr	Gly	Ser	Ser		
			85						90					95			
Ser	Leu	Thr	Phe	Gly	Gly	Gly	Thr	Glu	Val	Glu	Ile	Lys					
			100					105									

<210> 401  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 401																	
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ala	Thr	Leu	Ser	Ala	Ser	Val		
1				5				10					15				
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ser	Ser	Gln	Arg	Ile	Ser	Gly		
		20					25					30					
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Gln	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu		
		35				40					45						
Ile	Tyr	Ala	Ala	Ser	Thr	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Ile	Ser		
	50					55					60						
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Ser	Leu	Ile	Ile	Ser	Ser	Leu	Gln		
65					70					75					80		

Pro	Glu	Asp	Val	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Pro	Pro
				85					90					95	
Phe	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			100					105							

<210> 402  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 402															
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Thr	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Thr
			20					25					30		
Trp	Leu	Ala	Trp	Phe	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Lys	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln
65					70					75					80
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Lys	Ser	Tyr	Pro
				85					90					95	
Phe	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			100					105							

<210> 403  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 403															
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Ser	Val	Thr	Ile	Thr	Cys	Gln	Ala	Ser	Gln	Asp	Ile	Arg	Ser
			20					25					30		
Tyr	Leu	Asn	Trp	Phe	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Asp	Thr	Val	Asn	Leu	Gln	Thr	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Tyr	Ser	Phe	Thr	Ile	Asn	Asn	Leu	Gln
65					70					75					80
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Glu	Tyr	Leu	Pro
				85					90					95	
Ile	Thr	Phe	Gly	Gln	Gly	Thr	Arg	Leu	Glu	Ile	Lys				
			100					105							

<210> 404  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence



&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 404

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

&lt;210&gt; 405

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 405

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

&lt;210&gt; 406

&lt;211&gt; 109

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 406

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Ala	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Asp	Thr
			20					25					30		

Tyr Leu Asn Trp Tyr Gln His Lys Pro Gly Lys Ala Pro Lys Leu Leu  
           35                          40                          45  
 Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
           50                          55                          60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Gly Leu Gln  
 65                          70                          75                          80  
 Pro Glu Asp Phe Ala Ser Tyr Phe Cys Gln Gln Ser Tyr Ser Ser Pro  
                           85                          90                          95  
 Gly Ile Thr Phe Gly Gly Gly Thr Arg Val Glu Ile Lys  
                           100                          105

<210> 407  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 407  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1                          5                          10                          15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Gly Thr  
                           20                          25                          30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Asn Leu Leu  
           35                          40                          45  
 Ile Tyr Gly Thr Ser Ser Leu Gln Arg Gly Val Pro Ser Arg Phe Ser  
 50                          55                          60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Thr Leu Gln  
 65                          70                          75                          80  
 Pro Glu Asp Phe Val His Tyr Tyr Cys Gln Gln Ser Tyr Thr Ser Pro  
                           85                          90                          95  
 Pro Thr Phe Gly Gln Gly Thr Lys Val Glu Val Lys  
                           100                          105

<210> 408  
 <211> 106  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 408  
 Gln Tyr Glu Leu Thr Gln Ser Pro Ser Val Ser Val Ser Pro Gly Gln  
 1                          5                          10                          15  
 Thr Ala Ser Ile Pro Cys Ser Gly Asp Gln Leu Gly Gly Lys Phe Thr  
                           20                          25                          30  
 Ser Trp Tyr Gln Gln Lys Ala Gly Gln Ser Pro Leu Leu Val Ile Tyr  
           35                          40                          45  
 Gln Asp Thr Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser  
 50                          55                          60  
 Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Thr Gly Thr Gln Ala Met  
 65                          70                          75                          80  
 Asp Glu Ala Asp Tyr Tyr Cys Leu Val Trp Asp Ser Asn Thr Tyr Val  
                           85                          90                          95  
 Phe Gly Pro Gly Thr Lys Val Thr Val Leu

100

105

<210> 409  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 409  
 Gln Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln  
 1 5 10 15  
 Thr Ala Arg Ile Thr Cys Gly Gly Asn Asn Ile Gly Ser Lys Ser Val  
 20 25 30  
 His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Val Tyr  
 35 40 45  
 Asp Asp Ser Asp Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser  
 50 55 60  
 Asn Ser Gly Asn Thr Ala Thr Leu Thr Val Ser Arg Val Glu Ala Gly  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Val Trp Asp Ser Ser Gly Gly Leu  
 85 90 95  
 Gln Val Phe Gly Thr Gly Thr Lys Val Thr Val Leu  
 100 105

<210> 410  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 410  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser  
 20 25 30  
 Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Lys Ala Ser Asn Leu Glu Ser Glu Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ala Tyr Pro  
 85 90 95  
 Phe Ile Phe Gly Pro Gly Thr Lys Val Asp Ile Lys  
 100 105

<210> 411  
 <211> 107  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 411

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

&lt;210&gt; 412

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 412

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

&lt;210&gt; 413

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 413

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Ile	Ser	Ser
			20					25					30		
Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser

50		55		60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln				
65		70		75
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Ser Phe Pro				80
	85		90	95
Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys				
	100		105	

<210> 414  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 414
Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
1 5 10 15
Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser
20 25 30
Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu
35 40 45
Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser
50 55 60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln
65 70 75 80
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ser Thr Pro
85 90 95
Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile Lys
100 105

<210> 415  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 415
Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro
1 5 10 15
Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser
20 25 30
Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu
35 40 45
Leu Ile Tyr Gly Ala Ser Asn Arg Val Thr Gly Ile Pro Asp Arg Phe
50 55 60
Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu
65 70 75 80
Glu Pro Glu Asp Ser Ala Val Tyr Tyr Cys His Gln Tyr Gly Ser Gln
85 90 95
Tyr Thr Phe Gly Pro Gly Thr Lys Leu Glu Ile Lys
100 105

<210> 416

<211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 416  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Ser Cys Gln Ala Ser Gln His Ile Ser Asn  
 20 25 30  
 Phe Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Asp Thr Ser Asn Leu Arg Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Asp Phe Thr Phe Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Ile Ala Thr Tyr Tyr Cys Gln Gln Tyr Asp Asn Phe Pro  
 85 90 95  
 Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 417  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 417  
 Gln Ser Glu Leu Thr Gln Pro Pro Ser Val Ser Glu Ala Pro Gly Gln  
 1 5 10 15  
 Arg Val Thr Ile Ser Cys Ser Gly Ser Thr Ser Asn Ile Gly Asn Asn  
 20 25 30  
 Ala Val Ile Trp Tyr Gln Gln Leu Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Tyr Asp Asp Leu Leu Pro Ser Gly Val Ser Asp Arg Phe Ser  
 50 55 60  
 Gly Ser Lys Ser Gly Thr Ser Gly Ser Leu Ala Ile Ser Gly Leu Gln  
 65 70 75 80  
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ser Trp Asp Asp Asn Leu  
 85 90 95  
 Asn Gly Pro Leu Phe Gly Gly Gly Thr Lys Leu Thr Val Leu  
 100 105 110

<210> 418  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 418  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr Leu Ser Ala Ser Val

1				5					10					15			
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser		
			20					25					30				
Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu		
		35					40					45					
Ile	Tyr	Gln	Ala	Ser	Thr	Leu	Glu	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser		
	50					55					60						
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln		
65					70					75					80		
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ala	Asn	Ser	Phe	Pro		
			85						90					95			
Ile	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Asn						
			100					105									

<210> 419  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 419																	
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val		
1				5				10					15				
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Gln	Ala	Ser	Gln	Asp	Ile	Thr	Asn		
			20				25					30					
Tyr	Leu	Asn	Trp	Tyr	Leu	His	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu		
		35				40					45						
Ile	Tyr	Asp	Ala	Ser	Asn	Leu	Gln	Thr	Gly	Val	Pro	Ser	Arg	Phe	Ser		
	50				55					60							
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Phe	Thr	Ile	Thr	Ser	Leu	Gln		
65					70				75						80		
Pro	Glu	Asp	Phe	Gly	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Asp	Thr	Leu	His		
			85					90						95			
Pro	Ser	Phe	Gly	Pro	Gly	Thr	Thr	Val	Asp	Ile	Lys						
			100					105									

<210> 420  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 420																	
Gln	Ser	Ala	Leu	Thr	Gln	Pro	Pro	Ser	Ala	Ser	Gly	Ser	Pro	Gly	Gln		
1				5				10					15				
Ser	Val	Thr	Ile	Ser	Cys	Thr	Gly	Thr	Ser	Ser	Asp	Val	Gly	Val	Tyr		
			20				25					30					
Asp	Ser	Val	Ser	Trp	Tyr	Gln	Gln	His	Pro	Gly	Lys	Ala	Pro	Lys	Leu		
		35				40					45						
Met	Ile	Tyr	Asp	Val	Ser	Asp	Arg	Pro	Ser	Gly	Val	Ser	Asn	Arg	Phe		
	50				55					60							
Ser	Gly	Ser	Lys	Ser	Gly	Tyr	Thr	Ala	Ser	Leu	Thr	Ile	Ser	Ala	Leu		
65					70				75						80		

Gln	Ala	Glu	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Gly	Ser	Tyr	Arg	Ala	Ser
				85					90					95	
Ser	Ser	Tyr	Val	Phe	Gly	Thr	Gly	Thr	Lys	Val	Thr	Val	Leu		
			100					105					110		

<210> 421  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 421															
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Asn	Ile	Tyr	His
			20					25					30		
Tyr	Leu	His	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Ala	Ala	Ser	Thr	Leu	Glu	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Asn	Ser	Leu	Gln
65					70					75					80
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Thr	Tyr	Asp	Thr	Pro
				85					90					95	
Leu	Thr	Phe	Gly	Gly	Gly	Ser	Lys	Val	Glu	Val	Lys				
			100					105							

<210> 422  
 <211> 107  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 422															
Gln	Ser	Ala	Leu	Thr	Gln	Pro	Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln
1				5					10					15	
Thr	Ala	Ser	Ile	Thr	Cys	Ser	Gly	Asp	Lys	Leu	Gly	Glu	Lys	Phe	Ala
			20					25					30		
Ser	Trp	Tyr	Gln	Gln	Arg	Pro	Gly	Gln	Ser	Pro	Ile	Leu	Val	Ile	Tyr
		35					40					45			
Gln	Asp	Ser	Lys	Arg	Pro	Ser	Gly	Ile	Pro	Glu	Arg	Phe	Ser	Gly	Ser
	50					55					60				
Asn	Ser	Gly	Asn	Thr	Ala	Thr	Leu	Thr	Ile	Ser	Gly	Thr	Gln	Thr	Met
65					70					75					80
Asp	Glu	Ala	Asp	Tyr	Cys	Gln	Ala	Trp	Gly	Gly	Ser	Thr	Ala	Tyr	
				85				90					95		
Val	Phe	Gly	Ser	Gly	Thr	Lys	Val	Thr	Val	Leu					
			100					105							

<210> 423  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence



&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 423

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

&lt;210&gt; 424

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 424

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

&lt;210&gt; 425

&lt;211&gt; 107

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 425

Gln	Ser	Glu	Leu	Thr	Gln	Thr	Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln
1				5					10					15	
Thr	Ala	Ser	Ile	Thr	Cys	Ser	Gly	Asn	Arg	Leu	Glu	Asp	Lys	Tyr	Val
			20					25					30		

```

His Trp Tyr Gln Gln Lys Pro Gly Gln Ser Pro Val Leu Val Leu Tyr
   35                               40               45
Gln Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
   50                               55               60
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Glu Thr Gln Thr Leu
65                               70               75               80
Asp Glu Ala Asp Tyr Tyr Cys Gln Ala Trp Asp Ser Ser Thr Arg Glu
   85                               90               95
Val Phe Gly Gly Thr Lys Leu Thr Val Leu
   100                               105

```

<210> 426  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

```

<400> 426
Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Val Ser Ala Ser Val
 1                               5               10               15
Gly Asp Thr Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Val Ser Asp
   20                               25               30
Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Asn Leu Leu
   35                               40               45
Ile Tyr Gly Ala Ser Ser Leu Gln Thr Gly Val Pro Ser Arg Phe Ser
   50                               55               60
Gly Gly Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln
65                               70               75               80
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala His Ser Phe Pro
   85                               90               95
Phe Thr Phe Gly Gly Gly Thr Arg Val Glu Ile Lys
   100                               105

```

<210> 427  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

```

<400> 427
Gln Asp Ile Gln Met Thr Gln Ser Pro Ala Thr Leu Ser Val Ser Pro
 1                               5               10               15
Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Asn Ile Asn Thr
   20                               25               30
His Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
   35                               40               45
Ile Phe Gly Ala Ser Thr Arg Ala Thr Gly Ile Pro Ala Arg Phe Ser
   50                               55               60
Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln
65                               70               75               80
Ser Glu Asp Phe Thr Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Leu
   85                               90               95
Thr Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys

```

100

105

<210> 428  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 428  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro  
 1 5 10 15  
 Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Gly  
 20 25 30  
 Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Leu Gly Gln Ala Pro Arg Leu  
 35 40 45  
 Leu Ile Tyr Gly Ala Ser Ser Arg Ala Thr Gly Thr Pro Asp Arg Phe  
 50 55 60  
 Ser Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu  
 65 70 75 80  
 Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln His Asn Ser Tyr  
 85 90 95  
 Pro Leu Thr Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr  
 100 105

<210> 429  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 429  
 Gln Ser Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Glu  
 1 5 10 15  
 Thr Ala Thr Val Thr Cys Gly Gly Ile Asn Ile Gly Ser Lys Ser Val  
 20 25 30  
 His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Val Val Met Tyr  
 35 40 45  
 Phe Asp Ser Asp Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser  
 50 55 60  
 Asn Ser Gly Asp Thr Ala Thr Leu Thr Ile Ser Arg Val Glu Ala Gly  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Val Trp Asp Ser Arg Ser Asp Gln  
 85 90 95  
 Tyr Val Phe Gly Phe Gly Thr Lys Val Thr Val Leu  
 100 105

<210> 430  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 430

```

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

```

&lt;210&gt; 431

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 431

```

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

```

&lt;210&gt; 432

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 432

```

Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val
 1           5           10           15
Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Asn Asn
          20           25           30
Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu
          35           40           45
Ile Tyr Thr Ala Ser Thr Leu His Ser Gly Val Pro Ser Arg Phe Ser

```

50		55		60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Asn Leu Gln				
65		70		75
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ala Thr Leu				
	85		90	95
Arg Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys				
100		105		

<210> 433  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 433
Gln Ser Val Leu Thr Gln Pro Ala Ser Val Ser Gly Ser Pro Gly Gln
1 5 10 15
Ser Ile Thr Ile Ser Cys Ser Gly Ser Ser Ser Asp Val Gly His Tyr
20 25 30
Asp Tyr Val Ser Trp Tyr Gln Gln His Pro Gly Lys Ala Pro Lys Leu
35 40 45
Met Ile Tyr Asp Val Ser Asn Arg Pro Ser Gly Val Ser Asn Arg Phe
50 55 60
Ser Gly Ser Lys Ser Gly Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu
65 70 75 80
Gln Ala Asp Asp Glu Ala Glu Tyr Tyr Cys Ser Ser Tyr Thr Ser Ser
85 90 95
Gly Thr Arg Val Phe Gly Thr Gly Thr Lys Val Thr Val Leu
100 105 110

<210> 434  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 434
Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr Leu Ser Ala Ser Val
1 5 10 15
Gly Asp Arg Val Thr Phe Thr Cys Arg Ala Ser Gln Ser Val Asn Asn
20 25 30
Trp Val Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu
35 40 45
Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser
50 55 60
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln
65 70 75 80
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser Tyr Ser Met Pro
85 90 95
Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
100 105

<210> 435

<211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 435  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Lys Tyr  
 20 25 30  
 Ser Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Trp Pro Ser Gly Gly Trp Thr Thr Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Gly Gly Tyr Tyr Tyr Ala Leu Asp Tyr Trp Gly Gln Gly Asn  
 100 105 110  
 Leu Val Thr Val Ser Ser  
 115

<210> 436  
 <211> 122  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 436  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Phe Tyr  
 20 25 30  
 Gln Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Gly Ile Tyr Pro Ser Gly Gly Tyr Thr Lys Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asp Leu Gly Tyr Gly Ser Ser Trp Tyr Tyr Phe Asp Tyr Trp  
 100 105 110  
 Gly Gln Gly Thr Leu Val Thr Val Ser Ser  
 115 120

<210> 437  
 <211> 117  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 437

```

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

```

<210> 438

<211> 119

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 438

```

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

```

<210> 439

<211> 126

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 439

```

Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1           5           10           15

```

```

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

```

```

<210> 440
<211> 125
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 440
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
  1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gln Tyr
   20          25          30
Asn Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
   35          40          45
Ser Trp Ile Ser Ser Ser Gly Gly Thr Glu Tyr Ala Asp Ser Val Lys
   50          55          60
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
   65          70          75          80
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
   85          90          95
Lys Asp Arg Gly Tyr Gly Ser Gly Ser Tyr Gly Ala Tyr Asp Ala Phe
  100          105          110
Asp Ile Trp Gly Gln Gly Lys Met Val Thr Val Ser Ser
  115          120          125

```

```

<210> 441
<211> 118
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 441
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
  1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr
   20          25          30
Ser Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
   35          40          45
Ser Ser Ile Tyr Pro Ser Gly Gly Ser Thr Leu Tyr Ala Asp Ser Val

```



50		55		60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr				
65		70		80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys				
	85		90	95
Ala Arg Trp Gly Val Gly Ala Thr Phe Asp Tyr Trp Gly Gln Gly Thr				
	100		105	110
Leu Val Thr Val Ser Ser				
115				

<210> 442  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 442
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
1 5 10 15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr
20 25 30
Met Met Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45
Ser Tyr Ile Tyr Pro Ser Gly Gly Trp Thr Ser Tyr Ala Asp Ser Val
50 55 60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65 70 75 80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
85 90 95
Ala Arg Phe Asp Tyr Thr Ile Gly Phe Asp Phe Trp Gly Gln Gly Thr
100 105 110
Leu Val Thr Val Ser Ser
115

<210> 443  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 443
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
1 5 10 15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr
20 25 30
Gly Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45
Ser Gly Ile Arg Ser Ser Gly Gly Val Thr Asn Tyr Ala Asp Ser Val
50 55 60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65 70 75 80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
85 90 95

Ala Arg Glu Gly Ser Gly Trp Ser Lys Ala Phe Asp Ile Trp Gly Gln  
                   100                  105                  110  
 Gly Thr Met Val Thr Val Ser Ser  
                   115                  120

<210> 444  
 <211> 127  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 444  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr  
                   20                  25                  30  
 Asn Met Arg Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Ser Ser Ile Ser Pro Ser Gly Gly Phe Thr Gly Tyr Ala Asp Ser Val  
                   50                  55                  60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
  65                  70                  75                  80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
                   85                  90                  95  
 Ala Arg Gly Ile Asn Tyr Tyr Asp Ser Ser Gly Tyr Tyr Pro Pro Val  
                   100                  105                  110  
 Gly Met Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser  
                   115                  120                  125

<210> 445  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 445  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr  
                   20                  25                  30  
 Asn Met Leu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Ser Ser Ile Ser Pro Ser Gly Gly Tyr Thr Val Tyr Ala Asp Ser Val  
                   50                  55                  60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
  65                  70                  75                  80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
                   85                  90                  95  
 Ala Arg Asp Arg Gly Gly Ser Pro Phe Arg Pro Asp Ala Phe Asp Ile  
                   100                  105                  110  
 Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser  
                   115                  120

<210> 446  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 446  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Glu Tyr  
 20 25 30  
 Asp Met Leu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Trp Pro Ser Gly Gly Phe Thr Gln Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asn Tyr Tyr Asp Phe Trp Ser Gly Pro Tyr Tyr Tyr Gly Met  
 100 105 110  
 Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser  
 115 120 125

<210> 447  
 <211> 122  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 447  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gly Tyr  
 20 25 30  
 Val Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Ser Pro Ser Gly Gly Tyr Thr Leu Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Ala Phe Ser Thr Arg Trp Tyr Gly Ala Phe Asp Ile Trp  
 100 105 110  
 Gly Gln Gly Thr Met Val Thr Val Ser Ser  
 115 120

<210> 448  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 448

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

&lt;210&gt; 449

&lt;211&gt; 121

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 449

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

&lt;210&gt; 450

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 450

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----

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

<210> 451  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 451																	
Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly		
1				5					10					15			
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr		
			20					25					30				
Glu	Met	Phe	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val		
		35					40					45					
Ser	Ser	Ile	Ser	Pro	Ser	Gly	Gly	Tyr	Thr	Tyr	Tyr	Ala	Asp	Ser	Val		
	50					55					60						
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr		
65					70				75						80		
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys		
			85					90					95				
Ala	Thr	Met	Thr	Thr	Ser	Gly	Phe	His	Leu	Ile	Trp	Gly	Gln	Gly	Thr		
			100				105						110				
Met	Val	Thr	Val	Ser	Ser												
			115														

<210> 452  
 <211> 117  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 452																	
Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly		
1				5					10					15			
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Asp	Tyr		
			20					25					30				
Pro	Met	Gln	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val		
		35					40					45					

```

Ser Trp Ile Gly Pro Ser Gly Gly Trp Thr Val Tyr Ala Asp Ser Val
 50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Asp Asp Gly Ile Ala Gly Phe Leu Trp Gly Gln Gly Thr Leu
          100          105          110
Val Thr Val Ser Ser
          115

```

<210> 453

<211> 125

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 453

```

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

```

<210> 454

<211> 123

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 454

```

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

```



<210> 457  
 <211> 126  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 457  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Lys Tyr  
 20 25 30  
 Pro Met Met Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Tyr Ile Tyr Pro Ser Gly Gly Lys Thr Ala Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Thr Tyr Gly Ser Gly Ser Tyr Tyr Leu Tyr Tyr Tyr Tyr Tyr  
 100 105 110  
 Met Asp Val Trp Gly Lys Gly Thr Thr Val Thr Val Ser Ser  
 115 120 125

<210> 458  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 458  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr  
 20 25 30  
 Ser Met Trp Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Tyr Ile Val Pro Ser Gly Gly Glu Thr Ile Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Lys Asn Leu Gly Glu Gly Phe Trp Ser Asp Tyr Tyr Pro Pro Leu  
 100 105 110  
 Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser  
 115 120 125

<210> 459  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence



&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 459

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

&lt;210&gt; 460

&lt;211&gt; 125

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 460

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

&lt;210&gt; 461

&lt;211&gt; 120

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 461

```

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

```

```

<210> 462
<211> 122
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 462
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1           5           10           15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Glu Tyr
          20           25           30
Thr Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35           40           45
Ser Tyr Ile Ser Pro Ser Gly Gly Thr Thr Asn Tyr Ala Asp Ser Val
          50           55           60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65           70           75           80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85           90           95
Ala Lys Gly Ala Trp Gly Asp Ile Tyr Tyr Tyr Gly Met Asp Val Trp
          100          105          110
Gly Gln Gly Thr Thr Val Thr Val Ser Ser
          115          120

```

```

<210> 463
<211> 115
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 463
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1           5           10           15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
          20           25           30
Trp Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val

```

		35					40				45					
Ser	Ser	Ile	Tyr	Pro	Ser	Gly	Gly	Ser	Thr	Glu	Tyr	Ala	Asp	Ser	Val	
	50					55					60					
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	
65					70				75					80		
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	
			85					90					95			
Ala	Lys	Gly	Ile	Trp	Phe	Asp	Pro	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	
			100					105					110			
Val	Ser	Ser														
		115														

<210> 464  
 <211> 121  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

<210> 465  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
                   85                  90                  95  
 Ala Arg Tyr Asp Ser Arg Ala Ala Ala Gly Thr Asn Ala Phe Asp Ile  
                   100                  105                  110  
 Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser  
                   115                  120

<210> 466  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 466  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr  
                   20                  25                  30  
 Lys Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Ser Gly Ile Ser Pro Ser Gly Gly Tyr Thr Tyr Tyr Ala Asp Ser Val  
                   50                  55                  60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
   65                  70                  75                  80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
                   85                  90                  95  
 Ala Arg Gly Gly Tyr Gly Trp Ser Tyr Tyr Tyr Tyr Gly Met Asp Val  
                   100                  105                  110  
 Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser  
                   115                  120

<210> 467  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 467  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr  
                   20                  25                  30  
 Trp Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Ser Val Ile Ser Ser Ser Gly Gly His Thr Phe Tyr Ala Asp Ser Val  
                   50                  55                  60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
   65                  70                  75                  80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
                   85                  90                  95  
 Ala Arg Asp Tyr Glu Gly Gly Ser Asn Asp Tyr Trp Gly Gln Gly Thr  
                   100                  105                  110  
 Leu Val Thr Val Ser Ser

115

<210> 468  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 468  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr  
 20 25 30  
 Ala Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Tyr Ser Ser Gly Gly Tyr Thr Ala Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Met Tyr Tyr Cys  
 85 90 95  
 Ala Arg Val Pro His Val Phe Arg Gly Glu Leu Asp Tyr Trp Gly Gln  
 100 105 110  
 Gly Thr Leu Val Thr Val Ser Ser  
 115 120

<210> 469  
 <211> 122  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 469  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Phe Tyr  
 20 25 30  
 Thr Met Trp Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Tyr Pro Ser Gly Gly Gln Thr Leu Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Pro Asp Ser Tyr Gly Tyr Leu Tyr Tyr Gly Met Asp Val Trp  
 100 105 110  
 Gly Gln Gly Thr Thr Val Thr Val Ser Ser  
 115 120

<210> 470  
 <211> 115  
 <212> PRT

<213> Artificial Sequence

 $\langle 220 \rangle$ 

<223> Made in a lab

<400> 470

[illegible]

<210> 471

<211> 125

<212> PRT

### <213> Artificial Sequence

$\langle 220 \rangle$

<223> Made in a lab

<400> 471

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

<210> 472

<211> 122

<212> PRT

### <213> Artificial Sequence

$\langle 220 \rangle$

<223> Made in a lab

&lt;400&gt; 472

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

&lt;210&gt; 473

&lt;211&gt; 112

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 473

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

&lt;210&gt; 474

&lt;211&gt; 120

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 474

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
1				5				10						15	
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr
			20					25					30		
Thr	Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
		35					40					45			

```

Ser Ser Ile Ser Pro Ser Gly Gly Met Thr Phe Tyr Ala Asp Ser Val
 50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Thr Tyr Asp Phe Trp Ser Gly Tyr Phe Asp Tyr Trp Gly Gln
          100          105          110
Gly Thr Leu Val Thr Val Ser Ser
          115          120

```

<210> 475  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

```

<400> 475
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Thr Tyr
          20          25          30
Glu Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Val Ile Trp Ser Ser Gly Gly His Thr Trp Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Ser Asn Gln Gly Asp Phe Trp Ser Gly Tyr Pro Phe Ala Phe
          100          105          110
Asp Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
          115          120          125

```

<210> 476  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

```

<400> 476
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr
          20          25          30
Asn Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Tyr Ile Ser Pro Ser Gly Gly Met Thr Lys Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys

```



				85					90					95			
Ala	Arg	Arg	Gly	Val	Leu	Gly	Tyr	Tyr	Gly	Met	Asp	Val	Trp	Gly	Gln		
			100					105					110				
Gly	Thr	Thr	Val	Thr	Val	Ser	Ser										
			115				120										

<210> 477  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

<210> 478  
 <211> 128  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

<210> 479  
 <211> 119  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 479  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr  
 20 25 30  
 Ser Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Val Pro Ser Gly Gly Leu Thr Trp Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asp Val Tyr Gly Asp Tyr Tyr Phe Asp Asn Trp Gly Gln Gly  
 100 105 110  
 Thr Leu Val Thr Val Ser Ser  
 115

<210> 480  
 <211> 119  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 480  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr  
 20 25 30  
 Asn Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Ser Pro Ser Gly Gly Arg Thr Gly Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Val Ser Gly Phe Tyr Ala Ala Phe Asp Tyr Trp Gly Gln Gly  
 100 105 110  
 Thr Leu Val Thr Val Ser Ser  
 115

<210> 481  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 481

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

&lt;210&gt; 482

&lt;211&gt; 123

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 482

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

&lt;210&gt; 483

&lt;211&gt; 119

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 483

```

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

```

```

<210> 484
<211> 124
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 484
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ala Tyr
          20          25          30
Arg Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Tyr Ile Ser Ser Ser Gly Gly Arg Thr Asp Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Thr Thr Gly Gly Leu Arg Tyr Phe Asp Trp Leu Ala Pro Ser Met Asp
          100          105          110
Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
          115          120

```

```

<210> 485
<211> 116
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 485
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ala Tyr
          20          25          30
Ser Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val

```

		35					40					45					
Ser	Tyr	Ile	Tyr	Pro	Ser	Gly	Gly	Ile	Thr	Thr	Tyr	Ala	Asp	Ser	Val		
	50					55					60						
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr		
65					70					75					80		
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys		
			85						90					95			
Ala	Arg	Glu	Gly	Gln	Val	Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Thr	Val		
		100						105					110				
Thr	Val	Ser	Ser														
		115															

<210> 486  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

<210> 487  
 <211> 121  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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

Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
			85						90					95	
Ala	Arg	Asp	Tyr	Gly	Val	Ala	Thr	Leu	Asp	Ala	Phe	Asp	Ile	Trp	Gly
			100					105					110		
Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser							
		115					120								

&lt;210&gt; 488

&lt;211&gt; 127

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 488

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

&lt;210&gt; 489

&lt;211&gt; 122

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 489

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

115

120

<210> 490  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 490  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Glu Tyr  
 20 25 30  
 Gly Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Arg Ser Ser Gly Gly Trp Thr Lys Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Lys Gly Leu Gly Ala Thr Ser Gly Glu Phe Asp Tyr Trp Gly Gln  
 100 105 110  
 Gly Thr Leu Val Thr Val Ser Ser  
 115 120

<210> 491  
 <211> 121  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 491  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Lys Tyr  
 20 25 30  
 Asp Met Trp Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Trp Pro Ser Gly Gly Trp Thr Ser Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Lys Ser Arg Gly Ser Pro Trp Tyr Gly Asp Phe Asp His Trp Gly  
 100 105 110  
 Gln Gly Thr Leu Val Thr Val Ser Ser  
 115 120

<210> 492  
 <211> 118  
 <212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 492

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

<210> 493

<211> 123

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 493

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

<210> 494

<211> 124

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab



&lt;400&gt; 494

```

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

```

&lt;210&gt; 495

&lt;211&gt; 124

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 495

```

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

```

&lt;210&gt; 496

&lt;211&gt; 118

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 496

```

Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1           5           10           15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr
          20           25           30

```

Gly	Met	Asp	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
		35					40					45			
Ser	Ser	Ile	Tyr	Ser	Ser	Gly	Gly	Tyr	Thr	Leu	Tyr	Ala	Asp	Ser	Val
		50				55					60				
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
65					70				75						80
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Thr	Tyr	Tyr	Cys
			85					90						95	
Ala	Thr	Arg	Thr	Trp	Ala	Asp	Ala	Phe	Asp	Val	Trp	Gly	Gln	Gly	Thr
			100					105					110		
Thr	Val	Thr	Val	Ser	Ser										
			115												

<210> 497  
 <211> 2  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 497  
 Asn Ser  
 1

<210> 498  
 <211> 6  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 498  
 Ser Cys Pro Pro Cys Pro  
 1 5

<210> 499  
 <211> 8  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 499  
 Gly Gly Gly Gly Ser Gly Asn Ser  
 1 5

<210> 500  
 <211> 8  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 500  
Gly Cys Pro Pro Cys Pro Asn Ser  
1 5

<210> 501  
<211> 8  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 501  
Gly Ser Pro Pro Ser Pro Asn Ser  
1 5

<210> 502  
<211> 8  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 502  
Gly Ser Pro Pro Ser Pro Asn Ser  
1 5

<210> 503  
<211> 8  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 503  
Gly Cys Pro Pro Cys Pro Asn Ser  
1 5

<210> 504  
<211> 8  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 504  
Gly Cys Pro Pro Cys Pro Asn Ser  
1 5

<210> 505  
<211> 9  
<212> PRT  
<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 505

Gly Cys Pro Pro Cys Pro Gly Asn Ser

1

5

<210> 506

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 506

Gly Cys Pro Pro Cys Pro Ala Asn Ser

1

5

<210> 507

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 507

Gly Cys Pro Pro Cys Pro Ala Asn Ser

1

5

<210> 508

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 508

Glu Glu Glu Glu Asp Glu Gly Asn Ser

1

5

<210> 509

<211> 10

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 509

Asn Tyr Gly Gly Gly Gly Ser Gly Asn Ser

1

5

10

<210> 510

<211> 10

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 510  
 Val Ser Glu Arg Pro Phe Pro Pro Asn Ser  
 1 5 10

<210> 511  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 511  
 Glu Pro Lys Ser Cys Asp Lys Thr Cys Cys Pro  
 1 5 10

<210> 512  
 <211> 11  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 512  
 Ser Gln Pro Glu Ile Val Pro Ile Ser Asn Ser  
 1 5 10

<210> 513  
 <211> 12  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 513  
 Gly Gly Gly Gly Ser Cys Pro Pro Cys Pro Asn Ser  
 1 5 10

<210> 514  
 <211> 12  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 514  
 Lys Ala Asp Phe Leu Thr Pro Ser Ile Gly Asn Ser  
 1 5 10

<210> 515  
 <211> 12  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 515  
 Gln Met Asn Ser Glu Leu Ser Val Leu Ala Asn Ser  
 1 5 10

<210> 516  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 516  
 Glu Pro Lys Ser Cys Asp Lys Thr His Thr Cys Cys Pro  
 1 5 10

<210> 517  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 517  
 Glu Pro Lys Ser Cys Asp Lys Thr Cys Pro Pro Cys Pro  
 1 5 10

<210> 518  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 518  
 Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Asn Ser  
 1 5 10

<210> 519  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 519  
 Gly Cys Pro Pro Cys Pro Gly Gly Gly Gly Ser Asn Ser  
 1 5 10

<210> 520  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 520  
 Gly Gly Gly Gly Ser Cys Pro Pro Cys Pro Gly Asn Ser  
 1 5 10

<210> 521  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 521  
 Gly Cys Pro Pro Cys Pro Gly Gly Gly Gly Ser Asn Ser  
 1 5 10

<210> 522  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 522  
 Gly Gly Gly Ala Ser Cys Pro Pro Cys Pro Gly Asn Ser  
 1 5 10

<210> 523  
 <211> 13  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 523  
 Gly Gly Gly Ala Ser Cys Pro Pro Cys Ala Gly Asn Ser  
 1 5 10

<210> 524  
 <211> 13  
 <212> PRT  
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<223> Made in a lab

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Asn Tyr Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Asn Ser  
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Glu	Pro	Lys	Ser	Cys	Asp	Lys	Cys	His	Thr	Cys	Pro	Pro	Cys	Pro
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<400> 563

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<400> 564

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<210> 574  
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<210> 575  
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<400> 575  
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<210> 576  
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<220>  
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<400> 576

Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro Cys  
 1 5 10 15  
 Pro

<210> 577

<211> 17

<212> PRT

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<400> 577

Glu Pro Lys Ser Thr Asp Lys Thr His Thr Cys Pro Pro Cys Pro Asn  
 1 5 10 15  
 Ser

<210> 578

<211> 17

<212> PRT

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<220>

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Glu Pro Lys Ser Thr Asp Lys Thr His Thr Ser Pro Pro Ser Pro Asn  
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<400> 579

Glu Pro Lys Ser Thr Asp Lys Thr His Thr Cys Pro Pro Cys Pro Asn  
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 Ser

<210> 580

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<220>  
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 Cys Pro

<210> 582  
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Pro Asn Ser

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 Pro Cys Pro Asn Ser  
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 Pro Cys Pro Asn Ser  
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<210> 600  
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 Asp Cys Pro Asn Ser  
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<400> 601  
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 His Cys Pro Asn Ser  
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			85					90				95			
Gln	Lys	Leu	Ser	Asn	Met	Glu	Asn	Arg	Leu	Lys	Pro	Phe	Phe	Thr	Cys
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ctgacgaagc	tgcaggcaca	gaaccagtgg	ctgcaggaca	tgacaactca	tctcattctg	1200
cgcagcttta	aggagttcct	gcagtccagc	ctgagggctc	ttcggcaaat	ggatcagccc	1260
agagggccca	caatcaagcc	ctgtcctcca	tgcaaatgcc	cggctccaaa	tcttcttggt	1320
ggttcacccg	tcttcacctt	ccctccaaag	atcaaggatg	tactcatgat	ctccctgagc	1380
cccatagtca	catgtgtggt	ggtggatgtg	agcgaggacg	accagatgt	ccagatcagc	1440
tggtttgtga	acaacgtgga	agtacacaca	gctcagacac	aaacccatag	agaggattac	1500
aacagtactc	tccgggtggt	cagtgccttc	cccatccagc	accaggactg	gatgagtggc	1560
aaggagttca	aatgcaaggt	caacaacaaa	gacctccag	cgcccatcga	gagaaccatc	1620
tcaaaaccca	aagggtcagt	aagagctcca	caggatatatg	tcttgctctc	accagaagaa	1680
gagatgacta	agaaacaggt	cactctgacc	tgcatggtca	cagacttcat	gcctgaagac	1740
atttacgtgg	agtggactaa	caacgggaaa	acagagctaa	actacaagaa	cactgaacca	1800
gtcctggact	ctgatggttc	ttacttcatg	tacagcaagc	tgagagtggg	aaagaagaac	1860
tggttgga	gaaatagcta	ctcctgttca	gtggtccacg	aggggtctgca	caatcaccac	1920
acgactaaga	gcttctcccg	gactccgggt	aaatgattct	agagcggccg	c	1971

&lt;210&gt; 606

<211> 647  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 606

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Ile	Ala	Arg	Gly	Val	Asp	Pro	Glu	Glu	Pro	Gln	Leu	Ser	Cys
			20					25					30		
Phe	Arg	Lys	Ser	Pro	Leu	Ser	Asn	Val	Val	Cys	Glu	Trp	Gly	Pro	Arg
		35					40					45			
Ser	Thr	Pro	Ser	Leu	Thr	Thr	Lys	Ala	Val	Leu	Leu	Val	Arg	Lys	Phe
	50					55					60				
Gln	Asn	Ser	Pro	Ala	Glu	Asp	Phe	Gln	Glu	Pro	Cys	Gln	Tyr	Ser	Gln
65					70					75					80
Glu	Ser	Gln	Lys	Phe	Ser	Cys	Gln	Leu	Ala	Val	Pro	Glu	Gly	Asp	Ser
			85						90					95	
Ser	Phe	Tyr	Ile	Val	Ser	Met	Cys	Val	Ala	Ser	Ser	Val	Gly	Ser	Lys
			100					105					110		
Phe	Ser	Lys	Thr	Gln	Thr	Phe	Gln	Gly	Cys	Gly	Ile	Leu	Gln	Pro	Asp
		115					120					125			
Pro	Pro	Ala	Asn	Ile	Thr	Val	Thr	Ala	Val	Ala	Arg	Asn	Pro	Arg	Trp
		130				135						140			
Leu	Ser	Val	Thr	Trp	Gln	Asp	Pro	His	Ser	Trp	Asn	Ser	Ser	Phe	Tyr
145					150					155					160
Arg	Leu	Arg	Phe	Glu	Leu	Arg	Tyr	Arg	Ala	Glu	Arg	Ser	Lys	Thr	Phe
			165						170					175	
Thr	Thr	Trp	Met	Val	Lys	Asp	Leu	Gln	His	His	Cys	Val	Ile	His	Asp
			180					185					190		
Ala	Trp	Ser	Gly	Leu	Arg	His	Val	Val	Gln	Leu	Arg	Ala	Gln	Glu	Glu
		195					200					205			
Phe	Gly	Gln	Gly	Glu	Trp	Ser	Glu	Trp	Ser	Pro	Glu	Ala	Met	Gly	Thr
	210					215					220				
Pro	Trp	Thr	Glu	Ser	Arg	Ser	Pro	Pro	Ala	Gly	Gly	Gly	Ser	Asp	Val
225					230					235					240
Ala	Ala	Pro	His	Arg	Gln	Pro	Leu	Thr	Ser	Ser	Glu	Arg	Ile	Asp	Lys
			245						250					255	
Gln	Ile	Arg	Tyr	Ile	Leu	Asp	Gly	Ile	Ser	Ala	Leu	Arg	Lys	Glu	Thr
			260					265					270		
Cys	Asn	Lys	Ser	Asn	Met	Cys	Glu	Ser	Ser	Lys	Glu	Ala	Leu	Ala	Glu
		275					280					285			
Asn	Asn	Leu	Asn	Leu	Pro	Lys	Met	Ala	Glu	Lys	Asp	Gly	Cys	Phe	Gln
290						295					300				
Ser	Gly	Phe	Asn	Glu	Glu	Thr	Cys	Leu	Val	Lys	Ile	Ile	Thr	Gly	Leu
305					310					315					320
Leu	Glu	Phe	Glu	Val	Tyr	Leu	Glu	Tyr	Leu	Gln	Asn	Arg	Phe	Glu	Ser
			325						330					335	
Ser	Glu	Glu	Gln	Ala	Arg	Ala	Val	Gln	Met	Ser	Thr	Lys	Val	Leu	Ile
			340					345					350		
Gln	Phe	Leu	Gln	Lys	Lys	Ala	Lys	Asn	Leu	Asp	Ala	Ile	Thr	Thr	Pro
		355					360					365			
Asp	Pro	Thr	Thr	Asn	Ala	Ser	Leu	Leu	Thr	Lys	Leu	Gln	Ala	Gln	Asn
	370					375					380				
Gln	Trp	Leu	Gln	Asp	Met	Thr	Thr	His	Leu	Ile	Leu	Arg	Ser	Phe	Lys

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385          390          395          400
Glu Phe Leu Gln Ser Ser Leu Arg Ala Leu Arg Gln Met Asp Gln Pro
          405          410          415
Arg Gly Pro Thr Ile Lys Pro Cys Pro Pro Cys Lys Cys Pro Ala Pro
          420          425          430
Asn Leu Leu Gly Gly Ser Ser Val Phe Ile Phe Pro Pro Lys Ile Lys
          435          440          445
Asp Val Leu Met Ile Ser Leu Ser Pro Ile Val Thr Cys Val Val Val
          450          455          460
Asp Val Ser Glu Asp Asp Pro Asp Val Gln Ile Ser Trp Phe Val Asn
465          470          475          480
Asn Val Glu Val His Thr Ala Gln Thr Gln Thr His Arg Glu Asp Tyr
          485          490          495
Asn Ser Thr Leu Arg Val Val Ser Ala Leu Pro Ile Gln His Gln Asp
          500          505          510
Trp Met Ser Gly Lys Glu Phe Lys Cys Lys Val Asn Asn Lys Asp Leu
          515          520          525
Pro Ala Pro Ile Glu Arg Thr Ile Ser Lys Pro Lys Gly Ser Val Arg
          530          535          540
Ala Pro Gln Val Tyr Val Leu Pro Pro Pro Glu Glu Glu Met Thr Lys
545          550          555          560
Lys Gln Val Thr Leu Thr Cys Met Val Thr Asp Phe Met Pro Glu Asp
          565          570          575
Ile Tyr Val Glu Trp Thr Asn Asn Gly Lys Thr Glu Leu Asn Tyr Lys
          580          585          590
Asn Thr Glu Pro Val Leu Asp Ser Asp Gly Ser Tyr Phe Met Tyr Ser
          595          600          605
Lys Leu Arg Val Glu Lys Lys Asn Trp Val Glu Arg Asn Ser Tyr Ser
610          615          620
Cys Ser Val Val His Glu Gly Leu His Asn His His Thr Thr Lys Ser
625          630          635          640
Phe Ser Arg Thr Pro Gly Lys
          645

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<210> 607

<211> 749

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 607

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

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Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	
		115					120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	
	130					135					140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	
145					150					155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	
			165					170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
		180						185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
	195						200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
	260							265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
	275						280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
		340						345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
	355						360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
	435							440				445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490						495		
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500						505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515					520						525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Lys	Tyr	Ser	
	530				535						540					
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550						555					560	
Ser	Ile	Trp	Pro	Ser	Gly	Gly	Trp	Thr	Thr	Tyr	Ala	Asp	Ser	Val	Lys	

				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Arg	Gly	Gly	Tyr	Tyr	Tyr	Ala	Leu	Asp	Tyr	Trp	Gly	Gln	Gly	Thr	Leu		
	610					615					620						
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly		
625					630					635					640		
Gly	Gly	Gly	Ser	Gln	Ser	Glu	Leu	Thr	Gln	Pro	Pro	Ser	Val	Ser	Val		
				645					650						655		
Ser	Pro	Gly	Gln	Thr	Ala	Thr	Ile	Thr	Cys	Ser	Gly	Glu	Lys	Leu	Gly		
			660					665					670				
Asp	Ile	Tyr	Ala	Ser	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ser	Pro	Val		
		675					680					685					
Leu	Val	Ile	Tyr	Gln	Asp	Thr	Lys	Arg	Pro	Ser	Gly	Ile	Pro	Val	Arg		
	690					695					700						
Phe	Ser	Gly	Ser	Asn	Ser	Gly	Asn	Thr	Ala	Thr	Leu	Thr	Ile	Ser	Gly		
705					710					715					720		
Thr	Gln	Ala	Met	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Gln	Ala	Trp	Asp	Ser		
				725				730						735			
Ser	Thr	Val	Phe	Gly	Gly	Gly	Thr	Lys	Leu	Thr	Val	Leu					
			740					745									

&lt;210&gt; 608

&lt;211&gt; 758

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 608

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

Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	195	200	205
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	210	215	220
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	225	230	235
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	245	250	255
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	450	455	460
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	465	470	475
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	485	490	495
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	500	505	510
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	515	520	525
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Phe	Tyr	Gln	530	535	540
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	545	550	555
Gly	Ile	Tyr	Pro	Ser	Gly	Gly	Tyr	Thr	Lys	Tyr	Ala	Asp	Ser	Val	Lys	565	570	575
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	580	585	590
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	595	600	605
Arg	Asp	Leu	Gly	Tyr	Gly	Ser	Ser	Trp	Tyr	Tyr	Phe	Asp	Tyr	Trp	Gly	610	615	620
Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	625	630	635
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Ala	Leu	Thr	Gln	Pro	Ala	640		

				645					650					655			
Ser	Val	Ser	Gly	Ser	Pro	Gly	Gln	Ser	Ile	Thr	Ile	Ser	Cys	Ser	Gly		
			660					665					670				
Thr	Asp	Ser	Asp	Val	Gly	Gly	Tyr	Asn	His	Val	Ser	Trp	Tyr	Gln	Gln		
		675					680					685					
His	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Ile	Ile	Tyr	Asp	Val	Asp	His	Arg		
	690					695					700						
Pro	Ser	Gly	Ile	Ser	Asn	Arg	Phe	Ser	Gly	Ser	Lys	Ser	Gly	Asn	Thr		
705					710					715					720		
Ala	Ser	Leu	Thr	Ile	Ser	Gly	Leu	Gln	Ala	Glu	Asp	Glu	Ala	Asp	Tyr		
			725					730						735			
Tyr	Cys	Ser	Ser	Tyr	Arg	Ser	Gly	Ser	Thr	Tyr	Val	Phe	Gly	Thr	Gly		
			740					745					750				
Thr	Lys	Val	Thr	Val	Leu												
			755														

&lt;210&gt; 609

&lt;211&gt; 751

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 609

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr		
			20					25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln		
		35					40					45					
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys		
	50					55					60						
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp		
65				70					75						80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys		
			85					90					95				
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg		
			100				105					110					
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu		
		115				120						125					
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg		
	130				135						140						
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val		
145				150					155						160		
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr		
			165					170						175			
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
			180				185						190				
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
		195				200						205					
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
	210				215						220						
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225				230					235						240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
			245					250						255			



Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	450	455	460
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	465	470	475
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	485	490	495
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	500	505	510
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	515	520	525
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Lys	Tyr	Ser	530	535	540
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	545	550	555
Gly	Ile	Ser	Pro	Ser	Gly	Tyr	Thr	Arg	Tyr	Ala	Asp	Ser	Val	Lys	Gly	565	570	575
Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	Gln	580	585	590
Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	Arg	595	600	605
Asp	Gln	Leu	Leu	Glu	Ala	Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Thr	Val	610	615	620
Thr	Val	Ser	Ser	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	625	630	635
Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Leu	Ser	Leu	Ser	645	650	655
Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	660	665	670
Ile	Ser	Ser	Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Glu	Ala	Pro	675	680	685
Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ala	Leu	Gln	Ser	Gly	Val	Pro	Ser	690	695	700
Arg	Phe	Ser	Gly	Ser	Gly	Leu	Gly	Thr	Val	Phe	Thr	Leu	Thr	Ile	Thr			

705					710					715					720
Ser	Leu	Gln	Pro	Glu	Asp	Ser	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr
				725					730					735	
Ser	Pro	Pro	Val	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys	
			740					745					750		

<210> 610  
 <211> 752  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 610

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70				75					80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
			85					90					95		
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100				105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115				120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130				135				140						
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145				150					155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
			165				170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180			185						190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
	195				200						205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210				215						220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225				230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
			245					250					255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
		260					265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275				280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290				295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310					315					320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325				330						335		

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Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val
      340      345      350
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser
      355      360      365
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys
      370      375      380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp
      385      390      395      400
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe
      405      410      415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu
      420      425      430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe
      435      440      445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
      450      455      460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
      465      470      475      480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
      485      490      495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
      500      505      510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
      515      520      525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Glu Tyr Pro
      530      535      540
Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
      545      550      555      560
Gly Ile Tyr Ser Ser Gly Gly Leu Thr Lys Tyr Ala Asp Ser Val Lys
      565      570      575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
      580      585      590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
      595      600      605
Arg Asp His Tyr Gly Asp Tyr Ala Phe Asp Tyr Trp Gly Gln Gly Thr
      610      615      620
Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser
      625      630      635      640
Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr
      645      650      655
Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser
      660      665      670
Gln Ser Ile Ser Ser Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys
      675      680      685
Ala Pro Lys Leu Leu Ile Tyr Lys Ala Ser Ser Leu Glu Ser Gly Val
      690      695      700
Pro Ser Arg Phe Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr
      705      710      715      720
Ile Ser Ser Leu Gln Pro Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln
      725      730      735
Tyr Asn Ser Tyr Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
      740      745      750

```

&lt;210&gt; 611

&lt;211&gt; 761

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 611

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
		180						185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290				295						300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	

```

Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu
      420      425      430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe
      435      440      445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
      450      455      460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465      470      475      480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
      485      490      495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
      500      505      510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
      515      520      525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr Glu
      530      535      540
Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545      550      555      560
Gly Ile Trp Pro Ser Gly Gly Trp Thr Gln Tyr Ala Asp Ser Val Lys
      565      570      575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
      580      585      590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
      595      600      605
Arg Asp Leu Ala Val Ala Gly Trp Asp Tyr Tyr Tyr Tyr Tyr Gly Met
      610      615      620
Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser Gly Gly Gly
625      630      635      640
Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly Ser Gln Asp Ile Gln
      645      650      655
Met Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala
      660      665      670
Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser Tyr Leu Ala Trp
      675      680      685
Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Asp Ala
      690      695      700
Ser Asn Arg Ala Thr Gly Ile Pro Ala Arg Phe Ser Gly Ser Gly Ser
705      710      715      720
Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu Pro Glu Asp Ser
      725      730      735
Ala Val Tyr Tyr Cys Gln Gln Arg Ser Asn Trp Pro Pro Phe Thr Phe
      740      745      750
Gly Gly Gly Thr Lys Val Glu Ile Lys
      755      760

```

<210> 612

<211> 759

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 612

```

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
  1      5      10      15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr

```

				20				25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65				70						75				80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85				90					95		
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145				150						155				160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170				175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225				230						235				240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250				255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315				320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330				335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				

```

Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
      485      490      495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
      500      505      510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
      515      520      525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gln Tyr Asn
      530      535      540
Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545      550      555      560
Trp Ile Ser Ser Ser Gly Gly Thr Glu Tyr Ala Asp Ser Val Lys Gly
      565      570      575
Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln
      580      585      590
Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Lys
      595      600      605
Asp Arg Gly Tyr Gly Ser Gly Ser Tyr Gly Ala Tyr Asp Ala Phe Asp
610      615      620
Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly
625      630      635      640
Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Gly Ser Gln Asp Ile Gln Met
      645      650      655
Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr
      660      665      670
Ile Thr Cys Gln Thr Ser Gln Ser Ile Asp Arg Tyr Leu Asn Trp Tyr
675      680      685
Gln Gln Lys Ala Gly Glu Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser
690      695      700
Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly
705      710      715      720
Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala
      725      730      735
Ser Tyr Tyr Cys Gln Gln Ser Tyr Arg Thr Pro Arg Thr Phe Gly Gln
740      745      750
Gly Thr Lys Val Glu Ile Lys
      755

```

<210> 613

<211> 754

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 613

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

```

				85				90					95			
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	
			100					105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	
		115					120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	
	130					135					140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	
145					150					155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	
			165					170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
		180						185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
		195					200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250						255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
		340						345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410						415		
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490						495		
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500						505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515					520						525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	His	Tyr	Ser	
	530					535					540					



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Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545          550          555          560
Ser Ile Tyr Pro Ser Gly Gly Ser Thr Leu Tyr Ala Asp Ser Val Lys
          565          570          575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
          580          585          590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
          595          600          605
Arg Trp Gly Val Gly Ala Thr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
          610          615          620
Val Thr Val Ser Ser Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly
625          630          635          640
Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu
          645          650          655
Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
          660          665          670
Ser Val Ser Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
          675          680          685
Ala Pro Arg Leu Leu Ile Tyr Gly Ala Ser Ser Arg Ala Thr Gly Ile
          690          695          700
Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
705          710          715          720
Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
          725          730          735
Tyr Gly Ser Ser Pro Leu Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
          740          745          750
Ile Lys

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<210> 614

<211> 752

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 614

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

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145					150					155					160		
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr		
				165					170					175			
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
				180					185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
				195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
				210					215					220			
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225					230					235					240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
				245					250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro		
				260					265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro		
				275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr		
				290					295					300			
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn		
305					310					315					320		
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg		
				325					330					335			
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val		
				340					345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser		
				355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys		
				370					375					380			
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp		
385					390					395					400		
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe		
				405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu		
				420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe		
				435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly		
				450					455					460			
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr		
465					470					475					480		
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser		
				485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
				500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
				515					520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	His	Tyr	Met		
				530					535					540			
Met	Ala	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		
Tyr	Ile</																

Arg	Phe	Asp	Tyr	Thr	Ile	Gly	Phe	Asp	Phe	Trp	Gly	Gln	Gly	Thr	Leu
610						615					620				
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly
625					630					635					640
Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu
				645					650					655	
Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln
			660					665					670		
Ser	Ile	Ser	Ser	Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala
		675					680					685			
Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro
	690					695					700				
Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile
705				710					715						720
Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser
			725					730					735		
Tyr	Ser	Thr	Pro	Arg	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys
			740				745						750		

&lt;210&gt; 615

&lt;211&gt; 756

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 615

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50				55					60					
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65				70					75					80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
			85					90					95		
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100				105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
	115					120						125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130				135						140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145				150						155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
			165					170					175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
	195					200						205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215				220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser

225					230					235				240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr
				245					250					255
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro
			260					265					270	
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe
		275					280					285		Pro
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val
	290					295				300				Thr
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe
305					310					315				Asn
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro
			325						330					Arg
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr
			340					345					350	Val
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val
		355					360					365		Ser
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala
	370					375					380			Lys
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg
385				390						395				Asp
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly
			405						410					Phe
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro
			420					425					430	Glu
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser
	435						440					445		Phe
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln
	450					455					460			Gly
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His
465					470				475					Tyr
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn
			485						490					Ser
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser
		500						505					510	Glu
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
	515					520						525		Ser
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	His	Tyr
	530				535						540			Gly
Met	Thr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
545					550				555					Ser
Gly	Ile	Arg	Ser	Ser	Gly	Gly	Val	Thr	Asn	Tyr	Ala	Asp	Ser	Val
			565						570					Lys
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
		580					585						590	Leu
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
	595					600						605		Ala
Arg	Glu	Gly	Ser	Gly	Trp	Ser	Lys	Ala	Phe	Asp	Ile	Trp	Gly	Gln
	610					615					620			Gly
Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly
625					630				635					Gly
Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro
				645					650					Ala
Thr	Leu	Ser	Leu	Ser	Pro	Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys	Arg
			660				665						670	Ala
Ser	Gln	Ser	Ile	Ser	Thr	Ser	Leu	Ala	Trp	Tyr	Gln	Gln	Ile	Pro
	675					680						685		Gly

Gln Ala Pro Arg Leu Leu Met Tyr Asp Ala Ser Lys Arg Ala Ser Gly  
 690 695 700  
 Ile Pro Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Leu Thr Leu  
 705 710 715 720  
 Thr Ile Ser Ser Leu Glu Pro Glu Asp Ser Ala Val Tyr Tyr Cys Gln  
 725 730 735  
 Leu Arg Ile Asn Trp Pro Pro Glu Phe Thr Phe Gly Pro Gly Thr Lys  
 740 745 750  
 Val Asp Ile Lys  
 755

<210> 616

<211> 761

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 616

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225 230 235 240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
 245 250 255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
 260 265 270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
 275 280 285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr

290		295		300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn				
305		310		315
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg				
	325		330	
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val				
	340		345	
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser				
	355		360	
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys				
	370		375	
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				
	405		410	
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420		425	
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435		440	
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450		455	
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				
	485		490	
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500		505	
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515		520	
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr Asn				
	530		535	
Met Arg Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Ser Ile Ser Pro Ser Gly Gly Phe Thr Gly Tyr Ala Asp Ser Val Lys				
	565		570	
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu				
	580		585	
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala				
	595		600	
Arg Gly Ile Asn Tyr Tyr Asp Ser Ser Gly Tyr Tyr Pro Pro Val Gly				
	610		615	
Met Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser Gly Gly				
625		630		635
Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile				
	645		650	
Gln Met Thr Gln Ser Pro Ser Ser Val Ser Ala Ser Val Gly Asp Arg				
	660		665	
Val Thr Ile Thr Cys Arg Ala Ser Arg Gly Val Ile Thr Trp Leu Asn				
	675		680	
Trp Tyr Gln Gln Lys Pro Gly Arg Val Pro Ser Pro Leu Ile Phe Gly				
	690		695	
Ala Ser Thr Leu Gln Thr Gly Val Pro Ser Arg Phe Ser Gly Ser Gly				
705		710		715
Ser Gly Thr Asp Phe Thr Leu Thr Ile Asn Asp Leu Gln Pro Glu Asp				
	725		730	
Phe Ala Thr Tyr Tyr Cys Gln Gln Thr His Ser Phe Pro Leu Thr Phe				
	740		745	
				750

Gly Gly Gly Thr Lys Val Glu Ile Lys  
           755                      760

<210> 617  
 <211> 755  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 617  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
   1                  5                  10                  15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
           20                  25                  30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
           35                  40                  45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
           50                  55                  60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
   65                  70                  75                  80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
           85                  90                  95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
           100                  105                  110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
           115                  120                  125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
           130                  135                  140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
   145                  150                  155                  160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
           165                  170                  175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
           180                  185                  190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
           195                  200                  205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
           210                  215                  220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
   225                  230                  235                  240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
           245                  250                  255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
           260                  265                  270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
           275                  280                  285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr  
           290                  295                  300  
 Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn  
   305                  310                  315                  320  
 Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg  
           325                  330                  335  
 Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val  
           340                  345                  350  
 Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser

		355					360					365					
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys		
	370					375					380						
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp		
385					390					395					400		
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe		
			405						410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu		
		420						425					430				
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe		
	435					440						445					
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly		
	450					455					460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr		
465					470					475					480		
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser		
			485						490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
		500						505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
	515					520					525						
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Trp	Tyr	Asn		
	530				535					540							
Met	Leu	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		
Ser	Ile	Ser	Pro	Ser	Gly	Gly	Tyr	Thr	Val	Tyr	Ala	Asp	Ser	Val	Lys		
		565						570					575				
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
	580						585						590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
	595					600					605						
Arg	Asp	Arg	Gly	Gly	Ser	Pro	Phe	Arg	Pro	Asp	Ala	Phe	Asp	Ile	Trp		
	610				615					620							
Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly		
625				630						635					640		
Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Ala	Leu	Thr	Gln	Pro		
		645						650					655				
Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln	Thr	Ala	Arg	Ile	Thr	Cys	Ser		
	660					665						670					
Gly	Asp	Lys	Leu	Gly	Glu	Lys	Tyr	Val	Ser	Trp	Tyr	Gln	Gln	Arg	Ser		
	675					680				685							
Gly	Gln	Ser	Pro	Leu	Leu	Val	Ile	Ser	Gln	Asp	Thr	Lys	Arg	Pro	Ser		
	690				695					700							
Gly	Ile	Pro	Glu	Arg	Phe	Ser	Gly	Ser	Asn	Ser	Gly	Asn	Thr	Ala	Thr		
705				710					715						720		
Leu	Thr	Ile	Ser	Gly	Ala	Gln	Ala	Ala	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys		
		725						730					735				
Gln	Thr	Trp	Asp	Ser	Asp	Thr	Tyr	Val	Phe	Gly	Ser	Gly	Ser	Thr	Val		
	740							745					750				
Thr	Val	Leu															
	755																

&lt;210&gt; 618

&lt;211&gt; 759

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence



&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 618

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
		180						185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
		340						345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
			405						410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu

			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455				460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500					505						510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515					520						525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Glu	Tyr	Asp	
	530				535				540							
Met	Leu	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550					555						560	
Ser	Ile	Trp	Pro	Ser	Gly	Gly	Phe	Thr	Gln	Tyr	Ala	Asp	Ser	Val	Lys	
			565				570							575		
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
			580				585						590			
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	
	595					600						605				
Arg	Asn	Tyr	Tyr	Asp	Phe	Trp	Ser	Gly	Pro	Tyr	Tyr	Tyr	Gly	Met	Asp	
	610				615							620				
Val	Trp	Gly	Gln	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	
625				630					635						640	
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	
			645				650						655			
Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	
			660				665						670			
Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser	Tyr	Leu	Asn	Trp	Tyr	
		675				680						685				
Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	
	690				695						700					
Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	
705				710					715						720	
Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala	
			725						730					735		
Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Thr	Leu	Trp	Thr	Phe	Gly	Gln	
			740				745						750			
Gly	Thr	Lys	Val	Glu	Ile	Lys										
	755															

&lt;210&gt; 619

&lt;211&gt; 756

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 619

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser	
1				5					10					15		
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	
			20					25					30			

Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
		260						265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465				470					475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser

				485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
			500					505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
		515					520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Gly	Tyr	Val		
	530					535					540						
Met	Gly	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		
Ser	Ile	Ser	Pro	Ser	Gly	Gly	Tyr	Thr	Leu	Tyr	Ala	Asp	Ser	Val	Lys		
				565					570						575		
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Arg	Ala	Phe	Ser	Thr	Arg	Trp	Tyr	Trp	Gly	Ala	Phe	Asp	Ile	Trp	Gly		
	610					615					620						
Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly		
625					630					635					640		
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser		
			645					650						655			
Pro	Ala	Thr	Leu	Ser	Leu	Ser	Pro	Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys		
			660					665					670				
Arg	Ala	Ser	Gln	Ser	Val	Arg	Ser	Tyr	Leu	Ala	Trp	Tyr	Gln	Gln	Lys		
	675						680					685					
Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu	Ile	Tyr	Asp	Ala	Ser	Asn	Arg	Ala		
	690					695				700							
Thr	Gly	Ile	Pro	Ala	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe		
705					710					715					720		
Thr	Leu	Thr	Ile	Thr	Ser	Leu	Gln	Pro	Glu	Asp	Ile	Ala	Thr	Tyr	Tyr		
				725					730					735			
Cys	Gln	Gln	Tyr	Asp	Asn	Leu	Pro	Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys		
			740					745					750				
Val	Glu	Ile	Lys														
		755															

&lt;210&gt; 620

&lt;211&gt; 759

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 620

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr		
			20					25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln		
		35					40					45					
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys		
	50					55					60						
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp		
65					70					75					80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys		
				85					90						95		

Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	100	105	110
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	115	120	125
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	130	135	140
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	145	150	155
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	165	170	175
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	180	185	190
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	195	200	205
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	210	215	220
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	225	230	235
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	245	250	255
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	450	455	460
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	465	470	475
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	485	490	495
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	500	505	510
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	515	520	525
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ile	Tyr	His	530	535	540
Met	Asn	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser			

545					550					555					560
Gly	Ile	Ser	Ser	Ser	Gly	Gly	Arg	Thr	Asn	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600					605			
Ser	Ser	Tyr	Arg	Ala	Ala	Gly	Trp	Val	Asp	Tyr	Tyr	Tyr	Gly	Met	Asp
	610					615					620				
Val	Trp	Gly	Gln	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly
625					630					635					640
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met
			645					650					655		
Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Phe	Val	Gly	Asp	Arg	Val	Thr
			660					665					670		
Ile	Thr	Cys	Arg	Thr	Ser	Gln	Ser	Val	Ala	Thr	Tyr	Val	Asn	Trp	Tyr
		675					680					685			
Gln	Gln	Lys	Pro	Gly	Glu	Gly	Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser
	690					695					700				
Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Arg	Gly	Ser	Gly	Ser	Gly
705				710					715						720
Thr	Asp	Phe	Thr	Leu	Thr	Ile	Gly	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala
			725					730					735		
Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Val	Pro	Ile	Thr	Phe	Gly	Gln
			740					745					750		
Gly	Thr	Arg	Leu	Asp	Ile	Lys									
		755													

&lt;210&gt; 621

&lt;211&gt; 756

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 621

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

Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465				470					475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515					520					525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Leu	Tyr	Asp
	530					535					540				
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545				550						555					560
Arg	Ile	Tyr	Ser	Ser	Gly	Gly	Thr	Thr	Gln	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Met	Tyr	Tyr	Cys	Ala
		595					600					605			
Arg	Asp	Pro	Gly	Tyr	Gly	Ser	His	His	Ser	Phe	Asp	Tyr	Trp	Gly	Gln

610		615		620
Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly				
625		630		635
Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro				
	645		650	655
Ser Thr Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg				
	660		665	670
Ala Ser Gln Ser Met Ser Asn Trp Leu Ala Trp Tyr Gln Gln Lys Pro				
	675		680	685
Gly Lys Ala Pro Lys Leu Leu Ile Tyr Asp Val Phe Thr Leu Lys Ser				
	690		695	700
Gly Val Pro Ser Arg Phe Ser Gly Ser Arg Ser Gly Thr Glu Phe Thr				
705		710		715
Leu Thr Ile Ser Ser Leu Gln Pro Asp Asp Phe Ala Thr Tyr Tyr Cys				
	725		730	735
Gln Gln Tyr Ser Asp Tyr Ser Gly Ile Thr Phe Gly Pro Gly Thr Glu				
	740		745	750
Val Asp Ile Arg				
	755			

<210> 622  
 <211> 754  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 622
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
1 5 10 15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
20 25 30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
35 40 45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
50 55 60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
65 70 75 80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
85 90 95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
100 105 110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
115 120 125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
130 135 140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
145 150 155 160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
165 170 175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
180 185 190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
195 200 205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser
210 215 220



Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245						250					255
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330						335
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
		515					520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	Tyr
	530					535					540				
Met	Thr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Gly	Ile	Tyr	Ser	Ser	Gly	Gly	Pro	Thr	Gln	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600					605			
Arg	Ala	Gly	Gly	Asp	Ala	Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Met	Val
	610					615					620				
Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly
625					630					635					640
Gly	Gly	Ser	Gln	Ser	Val	Leu	Thr	Gln	Pro	Pro	Ser	Ala	Ser	Gly	Thr
				645					650					655	
Pro	Gly	Gln	Arg	Val	Thr	Ile	Ser	Cys	Ser	Gly	Ser	Ser	Ser	Asn	Ile
			660					665					670		
Gly	Ser	Lys	Val	Val	Tyr	Trp	Tyr	Gln	Gln	Val	Pro	Gly	Thr	Ala	Pro

	675					680						685					
Lys	Leu	Leu	Ile	Tyr	Arg	Asn	Asp	Gln	Arg	Pro	Ser	Gly	Val	Pro	Asp		
	690					695					700						
Arg	Phe	Ser	Gly	Ser	Lys	Ser	Gly	Thr	Ser	Ala	Ser	Leu	Ala	Ile	Ser		
705					710					715					720		
Gly	Leu	Arg	Ser	Glu	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Ala	Ala	Trp	Asp		
				725					730					735			
Asp	Ser	Leu	Ser	Gly	Arg	Trp	Val	Phe	Gly	Gly	Gly	Thr	Lys	Leu	Thr		
			740					745					750				
Val	Leu																

<210> 623  
 <211> 752  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 623

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr		
			20					25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln		
		35					40					45					
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys		
	50					55					60						
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp		
65				70					75						80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys		
				85					90					95			
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg		
			100					105					110				
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu		
		115					120					125					
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg		
	130					135				140							
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val		
145				150					155						160		
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr		
				165					170					175			
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
		180						185					190				
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
		195					200					205					
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
	210					215					220						
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225					230					235					240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
				245				250						255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro		
		260						265					270				
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro		
		275					280						285				

Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
		515					520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	Glu
	530					535					540				
Met	Phe	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Ser	Pro	Ser	Gly	Gly	Tyr	Thr	Tyr	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600				605				
Thr	Met	Thr	Thr	Ser	Gly	Phe	His	Leu	Ile	Trp	Gly	Gln	Gly	Thr	Met
	610					615					620				
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly
625					630					635					640
Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ala	Thr	Leu
				645					650					655	
Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln
			660					665					670		
Ser	Ile	Ser	Ser	Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala
		675					680					685			
Pro	Lys	Leu	Leu	Ile	Tyr	Lys	Ala	Ser	Gly	Leu	Glu	Ser	Gly	Val	Pro
	690					695					700				
Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Glu	Phe	Thr	Leu	Thr	Ile
705					710					715					720
Ser	Ser	Leu	Gln	Pro	Asp	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr
				725					730					735	
Asn	Ser	Tyr	Pro	Phe	Thr	Phe	Gly	Pro	Gly	Thr	Lys	Val	Asp	Ile	Lys

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740
745
750

<210> 624
<211> 751
<212> PRT
<213> Artificial Sequence

<220>
<223> Made in a lab

<400> 624
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1           5           10          15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
 20          25          30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
 35          40          45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
 50          55          60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
 65          70          75          80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
 85          90          95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
 100         105         110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
 115         120         125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
 130         135         140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
 145         150         155         160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
 165         170         175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
 180         185         190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
 195         200         205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser
 210         215         220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser
 225         230         235         240
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly
 245         250         255
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro
 260         265         270
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro
 275         280         285
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr
 290         295         300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn
 305         310         315         320
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg
 325         330         335
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val
 340         345         350
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser
 355         360         365

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Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys
 370                               375                               380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp
385                               390                               395                               400
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe
                               405                               410                               415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu
                               420                               425                               430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe
                               435                               440                               445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
                               450                               455                               460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465                               470                               475                               480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
                               485                               490                               495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
                               500                               505                               510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
                               515                               520                               525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asp Tyr Pro
                               530                               535                               540
Met Gln Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                               550                               555                               560
Trp Ile Gly Pro Ser Gly Gly Trp Thr Val Tyr Ala Asp Ser Val Lys
                               565                               570                               575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
                               580                               585                               590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
                               595                               600                               605
Arg Asp Asp Gly Ile Ala Gly Phe Leu Trp Gly Gln Gly Thr Leu Val
                               610                               615                               620
Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly
625                               630                               635                               640
Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Leu Ser Leu Ser
                               645                               650                               655
Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Gln Ala Ser Arg Asp
                               660                               665                               670
Ile Ser Ile Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Val Pro
                               675                               680                               685
Lys Leu Leu Ile Phe Gly Ala Ser Thr Leu Arg Ser Gly Val Pro Ser
                               690                               695                               700
Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Thr
705                               710                               715                               720
Ser Leu Gln Pro Glu Asp Val Ala Thr Tyr Tyr Cys Gln Lys Tyr Asn
                               725                               730                               735
Ser Ala Pro Tyr Thr Phe Gly Pro Gly Thr Lys Val Glu Ile Arg
                               740                               745                               750

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<210> 625

<211> 759

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

&lt;400&gt; 625

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser	1	5	10	15
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	20	25	30	
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	35	40	45	
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	50	55	60	
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	65	70	75	80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	85	90	95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	100	105	110	
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	115	120	125	
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	130	135	140	
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	145	150	155	160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	165	170	175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	180	185	190	
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	195	200	205	
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	210	215	220	
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	225	230	235	240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	245	250	255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270	
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285	
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300	
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315	320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350	
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365	
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380	
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395	400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430	
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445	

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Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
 450                               455           460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465                               470           475           480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
                               485           490           495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
                               500           505           510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
 515                               520           525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr Leu
 530                               535           540
Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                               550           555           560
Gly Ile Trp Pro Ser Gly Gly His Thr Leu Tyr Ala Asp Ser Val Lys
                               565           570           575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
                               580           585           590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
 595                               600           605
Arg Glu Pro Leu Leu Trp Phe Gly Glu Leu Ser Tyr Asn Trp Phe Asp
 610                               615           620
Pro Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly
625                               630           635           640
Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met
                               645           650           655
Thr Gln Ser Pro Ser Ser Val Ser Ala Ser Val Gly Asp Arg Val Thr
                               660           665           670
Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser Trp Leu Ala Trp Tyr
 675                               680           685
Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser
 690                               695           700
Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly
705                               710           715           720
Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala
 725                               730           735
Thr Tyr Tyr Cys Gln Gln Ala Asn Ser Phe Pro Leu Thr Phe Gly Gly
 740                               745           750
Gly Thr Lys Val Glu Ile Lys
 755

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<210> 626

<211> 757

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 626

```

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1                               5           10           15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
 20                               25           30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
 35                               40           45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys

```

50	55	60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp		
65	70	75
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys		
	85	90
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg		95
	100	105
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu		110
	115	120
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg		125
	130	135
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val		140
145	150	155
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr		160
	165	170
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly		175
	180	185
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser		190
	195	200
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser		205
	210	215
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser		220
225	230	235
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly		240
	245	250
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro		255
	260	265
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro		270
	275	280
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr		285
	290	295
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn		300
305	310	315
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg		320
	325	330
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val		335
	340	345
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser		350
	355	360
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys		365
	370	375
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp		380
385	390	395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe		400
	405	410
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu		415
	420	425
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe		430
	435	440
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly		445
	450	455
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr		460
465	470	475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser		480
	485	490
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu		495
	500	505
		510



```

Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
    515                      520                      525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr Glu
    530                      535                      540
Asn Met Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
545                      550                      555                      560
Ser Gly Ile Tyr Pro Ser Gly Gly Leu Thr Tyr Tyr Ala Asp Ser Val
    565                      570                      575
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
    580                      585                      590
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
    595                      600                      605
Ala Arg Ser Arg Arg Tyr Tyr Asp Ser Ser Asp Ala Phe Asp Ile Trp
    610                      615                      620
Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly
625                      630                      635                      640
Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln
    645                      650                      655
Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
    660                      665                      670
Cys Arg Ala Ser Gln Ser Ile Asn Thr Tyr Leu Ala Trp Tyr Gln Gln
    675                      680                      685
Lys Pro Gly His Pro Pro Arg Leu Leu Ile Tyr Asp Ala Ser Asn Arg
    690                      695                      700
Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
705                      710                      715                      720
Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Val Val Tyr
    725                      730                      735
Tyr Cys Gln Gln Tyr Gly Arg Ser Arg Tyr Thr Phe Gly Gln Gly Thr
    740                      745                      750
Lys Leu Glu Ile Lys
    755

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<210> 627

<211> 757

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 627

```

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

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[illegible]

Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu  
                   580                                  585                                  590  
 Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Cys Tyr Cys Ala  
                   595                                  600                                  605  
 Arg Asp Pro Gly Asp Phe Trp Ser Gly Tyr Tyr Gly Met Asp Val Trp  
                   610                                  615                                  620  
 Gly Gln Gly Thr Thr Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly  
                   625                                  630                                  635                                  640  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln  
                   645                                  650                                  655  
 Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr  
                   660                                  665                                  670  
 Cys Arg Ala Ser Gln Gly Ile Ser Asn Tyr Leu Ala Trp Tyr Gln Gln  
                   675                                  680                                  685  
 Lys Pro Gly Lys Val Pro Lys Leu Leu Ile Asn Ala Ala Ser Thr Leu  
                   690                                  695                                  700  
 Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Glu  
                   705                                  710                                  715                                  720  
 Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr  
                   725                                  730                                  735  
 Tyr Cys Gln Gln Ser Asp Asn Ile Pro Tyr Thr Phe Gly Leu Gly Thr  
                   740                                  745                                  750  
 Lys Leu Glu Ile Lys  
                   755

<210> 628

<211> 758

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 628

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

				180				185				190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
			195				200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
			210				215					220			
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225						230					235				240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245						250				255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
			275				280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
			290			295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
				340				345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
			355				360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
			370			375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
			435				440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
			450			455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
			515				520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Phe	Tyr	Gly
			530			535					540				
Met	Ala	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Tyr	Pro	Ser	Gly	Gly	Tyr	Thr	Asp	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580				585						590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
			595				600					605			
Arg	Ser	Ala	Gly	Gly	Trp	Ile	Gly	Gly	Gly	Ala	Phe	Asp	Ile	Trp	Gly
			610			615					620				
Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly
625					630					635					640

Gly Gly Ser Gly Gly Gly Gly Ser Gln Ser Glu Leu Thr Gln Pro Pro  
                                 645                                650                                655  
 Ser Ala Ser Gly Thr Pro Gly Gln Arg Val Thr Ile Ser Cys Ser Gly  
                                 660                                665                                670  
 Ser Ser Ser Asn Ile Gly Ser Asn Thr Val Asn Trp Tyr Gln Gln Leu  
                                 675                                680                                685  
 Pro Gly Thr Ala Pro Lys Leu Leu Ile Tyr Ser Asn Asn Gln Arg Pro  
                                 690                                695                                700  
 Ser Gly Val Pro Asp Arg Phe Ser Gly Ser Lys Ser Gly Thr Ser Ala  
 705                                710                                715                                720  
 Ser Leu Ala Ile Ser Gly Leu Gln Ser Glu Asp Glu Ala Asp Tyr Tyr  
                                 725                                730                                735  
 Cys Ala Ala Trp Asp Asp Ser Leu Asn Gly Tyr Val Phe Gly Ile Gly  
                                 740                                745                                750  
 Thr Lys Val Thr Val Leu  
                                 755

<210> 629

<211> 760

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 629

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1                                5                                10                                15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
                                 20                                25                                30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
                                 35                                40                                45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
                                 50                                55                                60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65                                70                                75                                80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
                                 85                                90                                95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
                                 100                                105                                110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
                                 115                                120                                125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
                                 130                                135                                140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145                                150                                155                                160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
                                 165                                170                                175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
                                 180                                185                                190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
                                 195                                200                                205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
                                 210                                215                                220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225                                230                                235                                240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly

				245				250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
			260					265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325						330					335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
			340					345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405						410					415		
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
	435						440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485						490					495		
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
			500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515						520						525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Lys	Tyr	Pro	
	530					535					540					
Met	Met	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550						555					560	
Tyr	Ile	Tyr	Pro	Ser	Gly	Gly	Lys	Thr	Ala	Tyr	Ala	Asp	Ser	Val	Lys	
			565						570					575		
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
			580					585					590			
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	
	595						600					605				
Thr	Tyr	Gly	Ser	Gly	Ser	Tyr	Tyr	Leu	Tyr	Tyr	Tyr	Tyr	Tyr	Tyr	Met	
	610					615					620					
Asp	Val	Trp	Gly	Lys	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	
625				630						635					640	
Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	
			645						650					655		
Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val	Gly	Asp	Ser	Val	
		660						665					670			
Thr	Ile	Ala	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Ser	Thr	Tyr	Leu	Asn	Trp	
	675						680					685				
Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Glu	Ala	
	690					695					700					

Thr	Asn	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser
705					710					715					720
Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp	Val
				725				730						735	
Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ala	Asn	Asn	Phe	Pro	Phe	Thr	Phe	Gly
			740					745					750		
Pro	Gly	Thr	Lys	Val	Asp	Ile	Lys								
		755					760								

&lt;210&gt; 630

&lt;211&gt; 761

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 630

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
	195						200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn

305					310					315				320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
			405						410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470				475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
			485						490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
	515						520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Trp	Tyr	Ser
	530					535					540				
Met	Trp	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550				555						560
Tyr	Ile	Val	Pro	Ser	Gly	Gly	Glu	Thr	Ile	Tyr	Ala	Asp	Ser	Val	Lys
			565						570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
		580						585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
	595						600					605			
Lys	Asn	Leu	Gly	Glu	Gly	Phe	Trp	Ser	Asp	Tyr	Tyr	Pro	Pro	Leu	Asp
	610					615					620				
Tyr	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly
625					630				635						640
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Val	Leu	Thr
			645						650					655	
Gln	Pro	Pro	Ser	Ala	Ser	Gly	Thr	Pro	Gly	Gln	Arg	Val	Thr	Ile	Ser
		660						665					670		
Cys	Ser	Gly	Ser	Ser	Ser	Asn	Ile	Gly	Gly	Asn	Ala	Val	Asn	Trp	Tyr
		675					680					685			
Gln	Gln	Leu	Pro	Gly	Thr	Ala	Pro	Lys	Leu	Val	Leu	Tyr	Ser	Ser	Asn
	690					695					700				
His	Arg	Pro	Ser	Gly	Val	Pro	Asp	Arg	Phe	Ser	Gly	Ser	Arg	Ser	Gly
705					710					715					720
Thr	Ser	Ala	Ser	Leu	Ala	Ile	Ser	Gly	Leu	Gln	Ser	Gln	Asp	Glu	Ala
			725						730					735	
Asp	Tyr	Tyr	Cys	Ala	Ala	Trp	Asp	Asp	Ser	Leu	Asn	Gly	Pro	Leu	Phe
			740					745					750		
Gly	Gly	Gly	Thr	Lys	Leu	Thr	Val	Leu							
		755					760								



<210> 631  
 <211> 752  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 631  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225 230 235 240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
 245 250 255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
 260 265 270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
 275 280 285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr  
 290 295 300  
 Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn  
 305 310 315 320  
 Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg  
 325 330 335  
 Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val  
 340 345 350  
 Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser  
 355 360 365  
 Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys

370		375		380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				400
	405		410	415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420		425	430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435		440	445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450		455	460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				480
	485		490	495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500		505	510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515		520	525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr Gly				
	530		535	540
Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Ser Ile Tyr Pro Gly Met Thr Gln Tyr Ala Asp Ser Val Lys Gly Arg				
	565		570	575
Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln Met				
	580		585	590
Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg Asp				
	595		600	605
Arg Gly Ser Gly Ile Asp Ala Phe Asp Ile Trp Gly Gln Gly Thr Met				
	610		615	620
Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly				
625		630		635
Gly Gly Gly Ser Gln Ser Glu Leu Thr Gln Pro Pro Ser Val Ser Val				
	645		650	655
Ser Pro Gly Gln Thr Ala Thr Val Ser Cys Ser Gly Glu Lys Leu Gly				
	660		665	670
Asp Arg Tyr Ala Trp Trp Tyr Gln Gln Lys Ala Gly Gln Ser Pro Val				
	675		680	685
Leu Val Ile Tyr Gln Asp Thr Lys Arg Pro Ser Gly Val Pro Asp Arg				
	690		695	700
Phe Ser Gly Ser Asn Ser Gly Thr Ser Ala Ser Leu Val Ile Ser Gly				
705		710		715
Pro Gln Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp				
	725		730	735
Ser Leu Lys Gly Val Val Phe Gly Gly Ser Lys Leu Thr Val Leu				
	740		745	750

&lt;210&gt; 632

&lt;211&gt; 761

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 632

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser	1	5	10	15
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	20	25	30	
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	35	40	45	
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	50	55	60	
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	65	70	75	80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	85	90	95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	100	105	110	
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	115	120	125	
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	130	135	140	
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	145	150	155	160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	165	170	175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	180	185	190	
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	195	200	205	
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	210	215	220	
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	225	230	235	240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	245	250	255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270	
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285	
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300	
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315	320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350	
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365	
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380	
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395	400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430	
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445	
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly				

450	455	460													
Asn Val Phe Ser Cys Ser	Val Met His Glu Ala Leu His Asn His Tyr														
465	470	475													480
Thr Gln Lys Ser Leu Ser	Leu Ser Pro Gly Gln Arg His Asn Asn Ser														
	485	490													495
Ser Leu Asn Thr Arg Thr	Gln Lys Ala Arg His Ser Pro Asn Ser Glu														
	500	505													510
Val Gln Leu Leu Glu Ser	Gly Gly Gly Leu Val Gln Pro Gly Gly Ser														
	515	520													525
Leu Arg Leu Ser Cys Ala	Ala Ser Gly Phe Thr Phe Ser Trp Tyr Asp														
	530	535													540
Met Leu Trp Val Arg Gln	Ala Pro Gly Lys Gly Leu Glu Trp Val Ser														560
	545	550													555
Val Ile Ser Pro Ser Gly	Gly Arg Thr Phe Tyr Ala Asp Ser Val Lys														
	565	570													575
Gly Arg Phe Thr Ile Ser	Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu														
	580	585													590
Gln Met Asn Ser Leu Arg	Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala														
	595	600													605
Val Thr Arg Ser Met Tyr	Ser Ser Ser Trp Tyr Gly Ala Pro Pro Thr														
	610	615													620
His Trp Gly Gln Gly Thr	Leu Val Thr Val Ser Ser Gly Gly Gly Gly														
	625	630													635
Ser Gly Gly Gly Gly Ser	Gly Gly Gly Gly Gly Ser Gln Ser Glu Leu Thr														
	645	650													655
Gln Pro Pro Ser Ala Ser	Gly Thr Pro Gly Gln Arg Val Thr Ile Ser														
	660	665													670
Cys Ser Gly Arg Ser Ser	Asn Ile Gly Ser Asn Ser Val Asn Trp Tyr														
	675	680													685
Gln Gln Leu Pro Gly Thr	Ala Pro Lys Leu Leu Ile Tyr Ser Asn Asn														
	690	695													700
His Arg Pro Ser Gly Val	Pro Asp Arg Phe Ser Gly Ser Lys Ser Gly														
	705	710													715
Thr Ser Ala Ser Leu Ala	Ile Ser Gly Leu Gln Ser Glu Asp Glu Ala														
	725	730													735
Asp Tyr Tyr Cys Ala Ala	Trp Asp Asp Ser Leu Tyr Gln Gly Val Phe														
	740	745													750
Gly Gly Gly Thr Lys Leu	Thr Val Leu														
	755	760													

&lt;210&gt; 633

&lt;211&gt; 754

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 633

Met Asp Phe Gln Val Gln	Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
1	5 10 15
Val Ile Met Ser Arg Gly	Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
	20 25 30
Ala Pro Glu Pro Gly Ser	Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
	35 40 45
Thr Ala Gln Met Cys Cys	Ser Lys Cys Ser Pro Gly Gln His Ala Lys
	50 55 60

Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser

		515					520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Trp	Tyr	Lys		
	530					535					540						
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		
Gly	Ile	Ser	Ser	Ser	Gly	Gly	Leu	Thr	Lys	Tyr	Ala	Asp	Ser	Val	Lys		
				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Thr		
		595					600					605					
Thr	Glu	Arg	Arg	Gly	Asp	Gly	Gly	Ala	Phe	Asp	Ile	Trp	Gly	Gln	Gly		
	610					615					620						
Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly		
625					630					635					640		
Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser		
				645					650					655			
Ser	Leu	Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala		
			660				665						670				
Ser	Gln	Ser	Ile	Ser	Ser	Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly		
		675				680					685						
Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly		
	690					695					700						
Val	Pro	Ser	Lys	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu		
705					710					715					720		
Thr	Ile	Ser	Ser	Leu	Leu	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln		
				725					730					735			
Gln	Tyr	Glu	Tyr	Phe	Pro	Pro	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu		
			740					745					750				
Ile	Lys																

<210> 634  
 <211> 757  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400>	634																
Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr		
			20					25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln		
		35					40					45					
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys		
	50					55					60						
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp		
65					70					75					80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys		
				85					90					95			
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg		
			100				105					110					
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu		
		115					120					125					

Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
130						135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310					315						320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
		340						345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
			405						410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465				470					475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
			485						490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
		515					520				525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Glu	Tyr	Thr
	530					535					540				
Met	Tyr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545				550					555						560
Tyr	Ile	Ser	Pro	Ser	Gly	Gly	Thr	Thr	Asn	Tyr	Ala	Gly	Ser	Val	Lys
			565						570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu

			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Lys	Gly	Ala	Trp	Gly	Asp	Ile	Tyr	Tyr	Tyr	Gly	Met	Asp	Val	Trp	Gly		
	610					615					620						
Gln	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly		
625					630					635					640		
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser		
			645					650				655					
Pro	Asp	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys		
		660					665					670					
Arg	Ala	Ser	Gln	Ser	Val	Ser	Ser	Tyr	Ser	Ala	Trp	Tyr	Gln	Gln			
		675					680				685						
Lys	Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu	Ile	Tyr	Gly	Ala	Ser	Ser	Arg		
	690					695					700						
Ala	Thr	Gly	Ile	Pro	Asp	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp		
705				710					715						720		
Phe	Thr	Leu	Thr	Ile	Ser	Arg	Leu	Glu	Pro	Glu	Asp	Phe	Ala	Val	Tyr		
				725					730					735			
Tyr	Cys	Gln	Gln	Tyr	Gly	Ser	Ser	Ser	Leu	Thr	Phe	Gly	Gly	Gly	Thr		
			740					745				750					
Glu	Val	Glu	Ile	Lys													
		755															

&lt;210&gt; 635

&lt;211&gt; 749

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 635

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



Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
				340				345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
		515					520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	Trp
	530					535					540				
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Tyr	Pro	Ser	Gly	Gly	Ser	Thr	Glu	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600					605			
Lys	Gly	Ile	Trp	Phe	Asp	Pro	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val
	610					615					620				
Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly
625					630					635					640
Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ala	Thr	Leu	Ser	Ala	Ser

				645					650					655			
Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ser	Ser	Gln	Arg	Ile	Ser		
			660					665					670				
Gly	Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Gln	Pro	Gly	Lys	Ala	Pro	Lys	Leu		
		675					680					685					
Leu	Ile	Tyr	Ala	Ala	Ser	Thr	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Ile		
	690					695					700						
Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Ser	Leu	Ile	Ile	Ser	Ser	Leu		
705					710					715					720		
Gln	Pro	Glu	Asp	Val	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Pro		
			725						730						735		
Pro	Phe	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys					
			740					745									

&lt;210&gt; 636

&lt;211&gt; 755

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 636

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr		
			20					25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln		
		35					40					45					
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys		
	50					55					60						
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp		
65					70					75					80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys		
				85					90					95			
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg		
			100					105					110				
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu		
		115					120						125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg		
	130					135					140						
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val		
145					150					155					160		
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr		
			165						170					175			
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
		180						185					190				
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
		195					200					205					
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
	210					215					220						
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225					230					235					240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
			245					250						255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro		
			260					265					270				

Cys 305	Pro 290	Ala 275	Pro 280	Glu 285	Ala 290	Ala 295	Gly 280	Ala 295	Pro 300	Ser 305	Val 285	Phe 290	Leu 295	Phe 300	Pro 305
Trp 310	Tyr 315	Val 320	Asp 325	Gly 330	Val 335	Glu 340	Val 345	His 350	Asn 355	Ala 360	Lys 365	Thr 370	Lys 375	Pro 380	Arg 385
Glu 390	Glu 395	Gln 400	Tyr 405	Asn 410	Ser 415	Thr 420	Tyr 425	Arg 430	Val 435	Val 440	Ser 445	Val 450	Leu 455	Thr 460	Val 465
Leu 470	His 475	Gln 480	Asp 485	Trp 490	Leu 495	Asn 500	Gly 505	Lys 510	Ala 515	Tyr 520	Ala 525	Cys 530	Ala 535	Val 540	Ser 545
Asn 550	Lys 555	Ala 560	Leu 565	Pro 570	Ala 575	Pro 580	Ile 585	Glu 590	Lys 595	Thr 600	Ile 605	Ser 610	Lys 615	Ala 620	Lys 625
Gly 630	Gln 635	Pro 640	Arg 645	Glu 650	Pro 655	Gln 660	Val 665	Tyr 670	Thr 675	Leu 680	Pro 685	Pro 690	Ser 695	Arg 700	Asp 705
Glu 710	Leu 715	Thr 720	Lys 725	Asn 730	Gln 735	Val 740	Ser 745	Leu 750	Thr 755	Cys 760	Leu 765	Val 770	Lys 775	Gly 780	Phe 785
Tyr 790	Pro 795	Ser 800	Asp 805	Ile 810	Ala 815	Val 820	Glu 825	Trp 830	Glu 835	Ser 840	Asn 845	Gly 850	Gln 855	Pro 860	Glu 865
Asn 870	Asn 875	Tyr 880	Lys 885	Thr 890	Thr 895	Pro 900	Pro 905	Val 910	Leu 915	Asp 920	Ser 925	Asp 930	Gly 935	Ser 940	Phe 945
Phe 950	Leu 955	Tyr 960	Ser 965	Lys 970	Leu 975	Thr 980	Val 985	Asp 990	Lys 995	Ser 1000	Arg 1005	Trp 1010	Gln 1015	Gln 1020	Gly 1025
Asn 1030	Val 1035	Phe 1040	Ser 1045	Cys 1050	Ser 1055	Val 1060	Met 1065	His 1070	Glu 1075	Ala 1080	Leu 1085	His 1090	Asn 1095	His 1100	Tyr 1105
Thr 1110	Gln 1115	Lys 1120	Ser 1125	Leu 1130	Ser 1135	Leu 1140	Ser 1145	Pro 1150	Gly 1155	Gln 1160	Arg 1165	His 1170	Asn 1175	Asn 1180	Ser 1185
Ser 1190	Leu 1195	Asn 1200	Thr 1205	Arg 1210	Thr 1215	Gln 1220	Lys 1225	Ala 1230	Arg 1235	His 1240	Ser 1245	Pro 1250	Asn 1255	Ser 1260	Glu 1265
Val 1270	Gln 1275	Leu 1280	Leu 1285	Glu 1290	Ser 1295	Gly 1300	Gly 1305	Gly 1310	Leu 1315	Val 1320	Gln 1325	Pro 1330	Gly 1335	Gly 1340	Pro 1345
Leu 1350	Arg 1355	Leu 1360	Ser 1365	Cys 1370	Ala 1375	Ala 1380	Ser 1385	Gly 1390	Phe 1395	Thr 1400	Phe 1405	Ser 1410	Arg 1415	Tyr 1420	Gly 1425
Met 1430	Met 1435	Trp 1440	Val 1445	Arg 1450	Gln 1455	Ala 1460	Pro 1465	Gly 1470	Lys 1475	Gly 1480	Leu 1485	Glu 1490	Trp 1495	Val 1500	Ser 1505
Tyr 1510	Ile 1515	Ser 1520	Ser 1525	Ser 1530	Gly 1535	Gly 1540	Phe 1545	Thr 1550	Arg 1555	Tyr 1560	Ala 1565	Asp 1570	Ser 1575	Val 1580	Lys 1585
Gly 1590	Arg 1595	Phe 1600	Thr 1605	Ile 1610	Ser 1615	Arg 1620	Asp 1625	Asn 1630	Ser 1635	Lys 1640	Asn 1645	Thr 1650	Leu 1655	Tyr 1660	Leu 1665
Gln 1670	Met 1675	Asn 1680	Ser 1685	Leu 1690	Arg 1695	Ala 1700	Glu 1705	Asp 1710	Thr 1715	Ala 1720	Val 1725	Tyr 1730	Cys 1735	Ala 1740	
Arg 1745	Val 1750	Gly 1755	Gly 1760	Tyr 1765	Ser 1770	Tyr 1775	Gly 1780	Pro 1785	His 1790	Phe 1795	Asp 1800	Phe 1805	Trp 1810	Gly 1815	Gln 1820
Gly 1825	Thr 1830	Leu 1835	Val 1840	Thr 1845	Val 1850	Ser 1855	Ser 1860	Gly 1865	Gly 1870	Gly 1875	Gly 1880	Ser 1885	Gly 1890	Gly 1895	Gly 1900
Gly 1905	Ser 1910	Gly 1915	Gly 1920	Gly 1925	Gly 1930	Ser 1935	Gln 1940	Asp 1945	Ile 1950	Gln 1955	Met 1960	Thr 1965	Gln 1970	Ser 1975	Pro 1980
Ser 1985	Thr 1990	Leu 1995	Ser 2000	Ala 2005	Ser 2010	Val 2015	Gly 2020	Asp 2025	Arg 2030	Val 2035	Thr 2040	Ile 2045	Thr 2050	Cys 2055	Arg 2060
Ala 2065	Ser 2070	Gln													

Gln	Gln	Tyr	Lys	725	Ser	Tyr	Pro	Phe	Thr	730	Phe	Gly	Gly	Gly	Thr	735	Lys	Val
			740						745							750		
Glu	Ile	Lys																
		755																

<210> 637  
 <211> 757  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 637

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser			
1				5					10					15				
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr			
			20					25					30					
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln			
		35					40					45						
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys			
	50					55					60							
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp			
65				70					75					80				
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys			
			85					90					95					
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg			
		100					105					110						
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu			
		115				120					125							
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg			
	130				135						140							
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val			
145				150						155				160				
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr			
			165					170					175					
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly			
		180				185						190						
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser			
	195				200							205						
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser			
	210				215						220							
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser			
225				230					235					240				
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly			
			245					250					255					
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro			
		260					265					270						
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro			
		275				280					285							
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr			
	290				295						300							
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn			
305				310					315					320				
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg			
			325					330						335				

Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val  
 340 345 350  
 Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser  
 355 360 365  
 Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys  
 370 375 380  
 Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp  
 385 390 395 400  
 Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe  
 405 410 415  
 Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu  
 420 425 430  
 Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe  
 435 440 445  
 Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly  
 450 455 460  
 Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr  
 465 470 475 480  
 Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser  
 485 490 495  
 Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu  
 500 505 510  
 Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser  
 515 520 525  
 Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr His  
 530 535 540  
 Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser  
 545 550 555 560  
 Trp Ile Ser Pro Ser Gly Gly Phe Thr Lys Tyr Ala Asp Ser Val Lys  
 565 570 575  
 Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu  
 580 585 590  
 Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala  
 595 600 605  
 Arg Tyr Asp Ser Arg Ala Ala Ala Gly Thr Asn Ala Phe Asp Ile Trp  
 610 615 620  
 Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly  
 625 630 635 640  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln  
 645 650 655  
 Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Ser Val Thr Ile Thr  
 660 665 670  
 Cys Gln Ala Ser Gln Asp Ile Arg Ser Tyr Leu Asn Trp Phe Gln Gln  
 675 680 685  
 Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Asp Thr Val Asn Leu  
 690 695 700  
 Gln Thr Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp  
 705 710 715 720  
 Tyr Ser Phe Thr Ile Asn Asn Leu Gln Pro Glu Asp Phe Ala Thr Tyr  
 725 730 735  
 Cys Cys Gln Gln Tyr Glu Tyr Leu Pro Ile Thr Phe Gly Gln Gly Thr  
 740 745 750  
 Arg Leu Glu Ile Lys  
 755

&lt;210&gt; 638

&lt;211&gt; 757

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 638

```

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1      5      10      15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
 20      25      30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
 35      40      45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
 50      55      60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
 65      70      75      80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
 85      90      95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
 100     105     110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
 115     120     125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
 130     135     140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
 145     150     155     160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
 165     170     175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
 180     185     190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
 195     200     205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser
 210     215     220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser
 225     230     235     240
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly
 245     250     255
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro
 260     265     270
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro
 275     280     285
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr
 290     295     300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn
 305     310     315     320
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg
 325     330     335
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val
 340     345     350
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser
 355     360     365
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys
 370     375     380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp
 385     390     395     400

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Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe  
 405 410 415  
 Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu  
 420 425 430  
 Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe  
 435 440 445  
 Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly  
 450 455 460  
 Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr  
 465 470 475 480  
 Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser  
 485 490 495  
 Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu  
 500 505 510  
 Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser  
 515 520 525  
 Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr Lys  
 530 535 540  
 Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser  
 545 550 555 560  
 Gly Ile Ser Pro Ser Gly Gly Tyr Thr Tyr Tyr Ala Asp Ser Val Lys  
 565 570 575  
 Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu  
 580 585 590  
 Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala  
 595 600 605  
 Arg Gly Gly Tyr Gly Trp Ser Tyr Tyr Tyr Tyr Gly Met Asp Val Trp  
 610 615 620  
 Gly Gln Gly Thr Thr Val Thr Val Ser Ser Gly Gly Gly Ser Gly  
 625 630 635 640  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln  
 645 650 655  
 Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr  
 660 665 670  
 Cys Arg Ala Ser Gln Ser Ile Ser Asn Tyr Leu Asn Trp Tyr Gln Gln  
 675 680 685  
 Lys Ser Gly Lys Ala Pro Asn Leu Leu Met Tyr Ala Ala Ser Ser Leu  
 690 695 700  
 His Ser Gly Val Ser Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Glu  
 705 710 715 720  
 Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Tyr Ala Thr Tyr  
 725 730 735  
 Tyr Cys Gln Gln Ser Tyr Ser Thr Ser Gly Thr Phe Gly Gln Gly Thr  
 740 745 750  
 Lys Val Glu Ile Lys  
 755

<210> 639

<211> 752

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 639

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser

1				5				10				15				
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	
			20					25					30			
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	
		35					40					45				
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	
	50					55					60					
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	
65				70					75					80		
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	
			85						90					95		
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	
			100					105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	
		115				120						125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	
	130					135					140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	
145				150						155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	
			165					170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
			180					185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
	195						200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225				230						235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250						255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
	275					280						285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
		340						345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
	355						360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410						415		
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
		420					425						430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
	435					440						445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					



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Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465                               470           475           480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
                               485           490           495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
                               500           505           510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
                               515           520           525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr Trp
                               530           535           540
Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                               550           555           560
Val Ile Ser Ser Ser Gly Gly His Thr Phe Tyr Ala Asp Ser Val Lys
                               565           570           575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
                               580           585           590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
                               595           600           605
Arg Asp Tyr Glu Gly Gly Ser Asn Asp Tyr Trp Gly Gln Gly Thr Leu
610                               615           620
Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly
625                               630           635           640
Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu
                               645           650           655
Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln
660                               665           670
Ser Ile Ser Ser Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala
675                               680           685
Pro Lys Leu Leu Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro
690                               695           700
Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
705                               710           715           720
Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ser
725                               730           735
Tyr Ser Thr Pro Val Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
740                               745           750

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<210> 640

<211> 755

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 640

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

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				85				90					95			
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	
			100					105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	
		115					120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	
	130					135					140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	
145					150					155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	
			165					170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
		180						185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
		195					200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250						255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
		340						345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410						415		
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
		420						425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490						495		
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500						505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515					520						525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Asn	Tyr	Ala	
	530					535					540					

```

Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545          550          555          560
Ser Ile Tyr Ser Ser Gly Gly Tyr Thr Ala Tyr Ala Asp Ser Val Lys
          565          570          575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
          580          585          590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Met Tyr Tyr Cys Ala
          595          600          605
Arg Val Pro His Val Phe Arg Gly Glu Leu Asp Tyr Trp Gly Gln Gly
          610          615          620
Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly
625          630          635          640
Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser
          645          650          655
Ser Leu Ser Ala Ser Val Gly Asp Arg Val Ala Ile Thr Cys Arg Ala
          660          665          670
Ser Gln Ser Ile Asp Thr Tyr Leu Asn Trp Tyr Gln His Lys Pro Gly
          675          680          685
Lys Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly
          690          695          700
Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu
705          710          715          720
Thr Ile Ser Gly Leu Gln Pro Glu Asp Phe Ala Ser Tyr Phe Cys Gln
          725          730          735
Gln Ser Tyr Ser Ser Pro Gly Ile Thr Phe Gly Gly Gly Thr Arg Val
          740          745          750
Glu Ile Lys
          755

```

<210> 641

<211> 756

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 641

```

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

```

145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465				470					475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
		515				520						525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Phe	Tyr	Thr
	530					535					540				
Met	Trp	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545				550					555						560
Ser	Ile	Tyr	Pro	Ser	Gly	Gly	Gln	Thr	Leu	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600					605			

```

Arg Pro Asp Ser Tyr Gly Tyr Leu Tyr Tyr Gly Met Asp Val Trp Gly
 610          615          620
Gln Gly Thr Thr Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly
625          630          635          640
Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser
          645          650          655
Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys
          660          665          670
Arg Ala Ser Gln Ser Ile Gly Thr Tyr Leu Asn Trp Tyr Gln Gln Lys
          675          680          685
Pro Gly Lys Ala Pro Asn Leu Leu Ile Tyr Gly Thr Ser Ser Leu Gln
          690          695          700
Arg Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
705          710          715          720
Thr Leu Thr Ile Ser Thr Leu Gln Pro Glu Asp Phe Val His Tyr Tyr
          725          730          735
Cys Gln Gln Ser Tyr Thr Ser Pro Pro Thr Phe Gly Gln Gly Thr Lys
          740          745          750
Val Glu Val Lys
          755

```

```

<210> 642
<211> 747
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in a lab

```

```

<400> 642
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1          5          10          15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
          20          25          30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
          35          40          45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
          50          55          60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
65          70          75          80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
          85          90          95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
          100          105          110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
          115          120          125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
          130          135          140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
145          150          155          160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
          165          170          175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
          180          185          190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
          195          200          205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser

```

210		215		220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser				
225		230		235
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly				240
	245		250	255
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro				
	260	265		270
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro				
	275	280		285
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr				
	290	295		300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn				
305		310		315
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg				
	325		330	335
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val				
	340		345	350
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser				
	355	360		365
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys				
	370	375		380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				
	405		410	415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420	425		430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435	440		445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450	455		460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				
	485		490	495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500	505		510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515	520		525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr Pro				
	530	535		540
Met Glu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Gly Ile Gly Pro Ser Gly Gly Gln Thr Thr Tyr Ala Asp Ser Val Lys				
	565	570		575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu				
	580	585		590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala				
	595	600		605
Arg Gly Ser Tyr Ser Phe Ile Trp Gly Gln Gly Thr Met Val Thr Val				
	610	615		620
Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly				
625		630		635
Ser Gln Tyr Glu Leu Thr Gln Ser Pro Ser Val Ser Val Ser Pro Gly				
	645	650		655
Gln Thr Ala Ser Ile Pro Cys Ser Gly Asp Gln Leu Gly Gly Lys Phe				
	660	665		670

Thr	Ser	Trp	Tyr	Gln	Gln	Lys	Ala	Gly	Gln	Ser	Pro	Leu	Leu	Val	Ile
		675					680					685			
Tyr	Gln	Asp	Thr	Lys	Arg	Pro	Ser	Gly	Ile	Pro	Glu	Arg	Phe	Ser	Gly
		690				695				700					
Ser	Asn	Ser	Gly	Asn	Thr	Ala	Thr	Leu	Thr	Ile	Thr	Gly	Thr	Gln	Ala
705					710					715					720
Met	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Leu	Val	Trp	Asp	Ser	Asn	Thr	Tyr
				725					730					735	
Val	Phe	Gly	Pro	Gly	Thr	Lys	Val	Thr	Val	Leu					
			740					745							

&lt;210&gt; 643

&lt;211&gt; 759

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 643

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
		130				135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200						205		
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
		210				215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr

290		295		300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn				
305		310		315
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg				
	325		330	
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val				
	340		345	
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser				
	355		360	
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys				
	370		375	
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				
	405		410	
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420		425	
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435		440	
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450		455	
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				
	485		490	
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500		505	
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515		520	
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr Lys				
	530		535	
Met Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Val Ile Gly Ser Ser Gly Gly Arg Thr Pro Tyr Ala Asp Ser Val Lys				
	565		570	
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu				
	580		585	
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala				
	595		600	
Arg Ala Pro Leu Ser Gly Trp Phe Gly Gln Ala His Asp Ala Phe Asp				
	610		615	
Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly				
625		630		635
Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Tyr Glu Leu Thr				
	645		650	
Gln Ser Pro Ser Val Ser Val Ala Pro Gly Gln Thr Ala Arg Ile Thr				
	660		665	
Cys Gly Gly Asn Asn Ile Gly Ser Lys Ser Val His Trp Tyr Gln Gln				
	675		680	
Lys Pro Gly Gln Ala Pro Val Leu Val Val Tyr Asp Asp Ser Asp Arg				
	690		695	
Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asn Thr				
705		710		715
Ala Thr Leu Thr Val Ser Arg Val Glu Ala Gly Asp Glu Ala Asp Tyr				
	725		730	
Tyr Cys Gln Val Trp Asp Ser Ser Gly Gly Leu Gln Val Phe Gly Thr				
	740		745	
				750



Gly Thr Lys Val Thr Val Leu  
755

<210> 644

<211> 756

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 644

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75				80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100				105						110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
		260						265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser

		355					360					365					
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys		
	370					375					380						
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp		
385					390					395					400		
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe		
				405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu		
			420					425					430				
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe		
	435					440						445					
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly		
	450					455					460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr		
465					470					475					480		
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser		
				485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
			500					505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
	515					520					525						
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Thr	Tyr	Glu		
	530				535					540							
Met	Asn	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550				555						560		
Gly	Ile	Val	Pro	Ser	Gly	Gly	Val	Thr	Tyr	Tyr	Ala	Asp	Ser	Val	Lys		
				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595				600					605						
Arg	Gly	Pro	Tyr	Ser	Tyr	Gly	His	Asp	Tyr	Gly	Met	Asp	Val	Trp	Gly		
	610					615				620							
Gln	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly		
625					630					635					640		
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser		
				645				650						655			
Pro	Ser	Thr	Leu	Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys		
			660					665					670				
Arg	Ala	Ser	Gln	Ser	Ile	Ser	Ser	Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys		
	675					680						685					
Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Lys	Ala	Ser	Asn	Leu	Glu		
	690					695				700							
Ser	Glu	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Glu	Phe		
705					710				715						720		
Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Asp	Asp	Phe	Ala	Thr	Tyr	Tyr		
				725				730						735			
Cys	Gln	Gln	Tyr	Asn	Ala	Tyr	Pro	Phe	Ile	Phe	Gly	Pro	Gly	Thr	Lys		
			740					745					750				
Val	Asp	Ile	Lys														
	755																

&lt;210&gt; 645

&lt;211&gt; 751

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 645

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65					70					75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
		180						185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu

			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455				460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500					505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
	515					520				525						
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Val	Tyr	Pro	
	530				535				540							
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550					555						560	
Ser	Ile	Tyr	Ser	Ser	Gly	Gly	Phe	Thr	Met	Tyr	Ala	Asp	Ser	Val	Lys	
			565				570						575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
		580				585						590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Val	
	595					600				605						
Arg	Glu	Gly	Val	Ala	Asp	Ala	Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Met	
	610					615				620						
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	
625				630					635					640		
Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Thr	Leu	
			645				650						655			
Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	
		660					665					670				
Ser	Ile	Ser	Ser	Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	
	675					680						685				
Pro	Lys	Leu	Leu	Ile	Tyr	Lys	Ala	Ser	Ser	Leu	Glu	Ser	Gly	Val	Pro	
	690					695				700						
Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Glu	Phe	Thr	Leu	Thr	Ile	
705			710						715					720		
Ser	Ser	Leu	Gln	Pro	Asp	Asp	Phe	Ala	Thr	Tyr	His	Cys	Gln	Gln	Tyr	
			725				730						735			
Lys	Ser	Tyr	Trp	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys		
		740				745						750				

&lt;210&gt; 646

&lt;211&gt; 756

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 646

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser	
1				5					10					15		
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	
			20				25					30				
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	
		35					40					45				

Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
50						55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65						70				75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
				340				345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu

				500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
		515					520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	Thr		
	530					535					540						
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		
Ser	Ile	Ser	Pro	Ser	Gly	Gly	Met	Thr	Phe	Tyr	Ala	Asp	Ser	Val	Lys		
				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Arg	Thr	Tyr	Asp	Phe	Trp	Ser	Gly	Tyr	Phe	Asp	Tyr	Trp	Gly	Gln	Gly		
	610					615					620						
Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly		
625					630				635						640		
Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Glu	Leu	Thr	Gln	Pro	Pro	Ser	Val		
				645				650						655			
Ser	Glu	Ala	Pro	Arg	Gln	Arg	Val	Thr	Val	Ser	Cys	Ser	Gly	Ser	Pro		
			660					665					670				
Ser	Asn	Ile	Gly	Ser	Asn	Ser	Val	Asn	Trp	Tyr	Gln	Gln	Leu	Pro	Gly		
		675					680					685					
Lys	Ala	Pro	Lys	Val	Val	Ile	Tyr	Tyr	Asp	Asp	Leu	Val	Pro	Ser	Gly		
	690					695				700							
Val	Ser	Asp	Arg	Phe	Ser	Gly	Ser	Lys	Ser	Gly	Thr	Ser	Ala	Ser	Leu		
705					710					715					720		
Ala	Ile	Ser	Gly	Leu	Gln	Ser	Glu	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Ala		
				725					730					735			
Ala	Trp	Asp	Asp	Arg	Leu	Asn	Gly	Trp	Val	Phe	Gly	Gly	Gly	Thr	Lys		
			740					745					750				
Leu	Thr	Val	Leu														
		755															

&lt;210&gt; 647

&lt;211&gt; 759

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 647

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

Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290				295						300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val</									

				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Arg	Ser	Asn	Gln	Gly	Asp	Phe	Trp	Ser	Gly	Tyr	Pro	Phe	Ala	Phe	Asp		
	610					615					620						
Ile	Trp	Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly		
625					630					635					640		
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met		
			645					650						655			
Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr		
			660					665					670				
Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Ile	Ser	Ser	Trp	Leu	Ala	Trp	Tyr		
		675					680					685					
Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser		
	690					695					700						
Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly		
705					710				715						720		
Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala		
			725					730					735				
Thr	Tyr	Tyr	Cys	Gln	Gln	Ala	Asn	Ser	Phe	Pro	Leu	Thr	Phe	Gly	Gly		
			740					745					750				
Gly	Thr	Lys	Val	Glu	Ile	Lys											
		755															

&lt;210&gt; 648

&lt;211&gt; 754

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 648

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



Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
			180					185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
		195					200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310						315					320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
			340					345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470						475					480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
			500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515						520					525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Asn	Tyr	Asn	
	530					535					540					
Met	Tyr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550						555					560	
Tyr	Ile	Ser	Pro	Ser	Gly	Gly	Met	Thr	Lys	Tyr	Ala	Asp	Ser	Val	Lys	
			565					570						575		
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
			580					585					590			
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	
	595						600					605				
Arg	Arg	Gly	Val	Leu	Gly	Tyr	Tyr	Gly	Met	Asp	Val	Trp	Gly	Gln	Gly	
	610					615					620					
Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	

625					630					635				640	
Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser
				645					650					655	
Ser	Leu	Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala
			660					665					670		
Ser	Gln	Ser	Ile	Ser	Ser	Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly
		675					680					685			
Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly
	690					695					700				
Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu
705					710					715					720
Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln
				725					730					735	
Gln	Ser	Tyr	Ser	Thr	Pro	Tyr	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Leu	Glu
			740					745					750		
Ile	Lys														

&lt;210&gt; 649

&lt;211&gt; 757

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 649

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55				60					
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65				70					75					80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85				90					95		
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100				105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
	115					120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135				140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145				150						155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165				170					175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180				185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
	195					200						205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215				220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225				230						235					240

Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
			405						410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
	435						440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
			485						490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
	515						520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Leu	Tyr	Ser
	530					535					540				
Met	Asn	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Ser	Ser	Ser	Gly	Gly	Ala	Thr	Leu	Tyr	Ala	Asp	Ser	Val	Lys
			565						570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
	595						600					605			
Arg	Asp	Leu	Ile	Ser	Ser	Gly	Tyr	Tyr	Pro	Asp	Ala	Phe	Asp	Ile	Trp
	610					615					620				
Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly
625					630					635					640
Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln
			645						650					655	
Ser	Pro	Gly	Thr	Leu	Ser	Leu	Ser	Pro	Gly	Glu	Arg	Ala	Thr	Leu	Ser
			660					665					670		
Cys	Arg	Ala	Ser	Gln	Ser	Val	Ser	Ser	Ser	Tyr	Leu	Ala	Trp	Tyr	Gln
	675						680					685			
Gln	Lys	Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu	Ile	Tyr	Gly	Ala	Ser	Asn

690		695		700
Arg Val Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr				
705		710		715
Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Ser Ala Val				
	725		730	735
Tyr Tyr Cys His Gln Tyr Gly Ser Gln Tyr Thr Phe Gly Pro Gly Thr				
	740		745	750
Lys Leu Glu Ile Lys				
755				

&lt;210&gt; 650

&lt;211&gt; 762

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 650

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser	
1	5
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr	
	20
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln	
	35
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys	
	50
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp	
65	70
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys	
	85
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg	
	100
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu	
	115
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg	
	130
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val	
145	150
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr	
	165
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly	
	180
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser	
	195
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser	
	210
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser	
225	230
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly	
	245
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro	
	260
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro	
	275
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr	
	290

Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
		500						505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
		515					520					525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Gln	Tyr	Met
	530					535					540				
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Ser	Ser	Ser	Gly	Gly	Trp	Thr	Ser	Tyr	Ala	Asp	Ser	Val	Lys
				565					570					575	
Gly	Ser	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
		595					600					605			
Arg	Pro	Leu	Ala	Leu	Gly	Tyr	Asp	Phe	Trp	Ser	Gly	Tyr	Gln	Ala	Ala
	610					615					620				
Gly	Phe	Asp	Tyr	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser	Gly
625					630					635					640
Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp
			645						650					655	
Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val	Gly	Asp
		660						665					670		
Arg	Val	Thr	Ile	Ser	Cys	Gln	Ala	Ser	Gln	His	Ile	Ser	Asn	Phe	Leu
		675					680					685			
Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu	Ile	Tyr
	690					695					700				
Asp	Thr	Ser	Asn	Leu	Arg	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser
705					710					715					720
Gly	Ser	Gly	Thr	Asp	Phe	Thr	Phe	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu
			725						730					735	
Asp	Ile	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Asp	Asn	Phe	Pro	Leu	Thr
			740					745					750		
Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys						

755

760

<210> 651  
 <211> 755  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 651

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser	1	5	10	15
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	20	25	30	
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	35	40	45	
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	50	55	60	
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	65	70	75	80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	85	90	95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	100	105	110	
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	115	120	125	
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	130	135	140	
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	145	150	155	160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	165	170	175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	180	185	190	
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	195	200	205	
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	210	215	220	
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	225	230	235	240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	245	250	255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	260	265	270	
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	275	280	285	
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	290	295	300	
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315	320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350	
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365	

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Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys
 370                               375                               380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp
 385                               390                               395                               400
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe
                               405                               410                               415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu
 420                               425                               430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe
 435                               440                               445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
 450                               455                               460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
 465                               470                               475                               480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
                               485                               490                               495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
 500                               505                               510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
 515                               520                               525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr Ser
 530                               535                               540
Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
 545                               550                               555                               560
Ser Ile Val Pro Ser Gly Gly Leu Thr Trp Tyr Ala Asp Ser Val Lys
 565                               570                               575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
 580                               585                               590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
 595                               600                               605
Arg Asp Val Tyr Gly Asp Tyr Tyr Phe Asp Asn Trp Gly Gln Gly Thr
 610                               615                               620
Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser
 625                               630                               635                               640
Gly Gly Gly Gly Ser Gln Ser Glu Leu Thr Gln Pro Pro Ser Val Ser
 645                               650                               655
Glu Ala Pro Gly Gln Arg Val Thr Ile Ser Cys Ser Gly Ser Thr Ser
 660                               665                               670
Asn Ile Gly Asn Asn Ala Val Ile Trp Tyr Gln Gln Leu Pro Gly Lys
 675                               680                               685
Ala Pro Lys Leu Leu Ile Tyr Tyr Asp Asp Leu Leu Pro Ser Gly Val
 690                               695                               700
Ser Asp Arg Phe Ser Gly Ser Lys Ser Gly Thr Ser Gly Ser Leu Ala
 705                               710                               715                               720
Ile Ser Gly Leu Gln Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ser
 725                               730                               735
Trp Asp Asp Asn Leu Asn Gly Pro Leu Phe Gly Gly Gly Thr Lys Leu
 740                               745                               750
Thr Val Leu
 755

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&lt;210&gt; 652

&lt;211&gt; 753

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Made in a lab

<400> 652

Met 1	Asp	Phe	Gln	Val 5	Gln	Ile	Phe	Ser	Phe 10	Leu	Leu	Ile	Ser	Ala 15	Ser
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala 25	Gln	Val	Ala	Phe	Thr 30	Pro	Tyr
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys 40	Arg	Leu	Arg	Glu	Tyr 45	Tyr	Asp	Gln
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys 55	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
Val 65	Phe	Cys	Thr	Lys	Thr	Ser	Asp 70	Thr	Val	Cys 75	Asp	Ser	Cys	Glu	Asp 80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val 105	Glu	Thr	Gln	Ala	Cys 110	Thr	Arg
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys 120	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu 135	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
Pro 145	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val 160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys 185	Asn	Val	Val	Ala	Ile 190	Pro	Gly
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys 200	Thr	Ser	Thr	Ser	Pro	Thr 205	Arg	Ser
Met	Ala	Pro	Gly	Ala	Val	His	Leu 215	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
Gln 225	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser 240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp 265	Lys	Thr	His	Thr	Cys 270	Pro	Pro
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly 280	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met 295	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
Cys 305	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn 320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn 330	Ala	Lys	Thr	Lys	Pro	Arg
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly 360	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile 375	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
Gly 385	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp 400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp 425	Glu	Ser	Asn	Gly	Gln	Pro	Glu



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Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe
    435                      440          445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
    450                      455          460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465                      470          475          480
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
    485                      490          495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
    500                      505          510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
    515                      520          525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr Asn
    530                      535          540
Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                      550          555          560
Ser Ile Ser Pro Ser Gly Gly Arg Thr Gly Tyr Ala Asp Ser Val Lys
    565                      570          575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
    580                      585          590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
    595                      600          605
Arg Val Ser Gly Phe Tyr Ala Ala Phe Asp Tyr Trp Gly Gln Gly Thr
    610                      615          620
Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser
625                      630          635          640
Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Thr
    645                      650          655
Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser
    660                      665          670
Gln Ser Ile Ser Ser Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys
    675                      680          685
Ala Pro Lys Leu Leu Ile Tyr Gln Ala Ser Thr Leu Glu Ser Gly Val
    690                      695          700
Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
705                      710          715          720
Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln
    725                      730          735
Ala Asn Ser Phe Pro Ile Thr Phe Gly Gly Gly Thr Lys Val Glu Ile
    740                      745          750
Asn

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<210> 653

<211> 754

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 653

```

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
  1              5              10              15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
    20              25              30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln

```

		35					40				45				
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50						55				60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65							70				75				80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310					315						320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470				475						480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	

Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu  
 500 505 510  
 Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser  
 515 520 525  
 Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Trp Tyr Trp  
 530 535 540  
 Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser  
 545 550 555 560  
 Ser Ile Pro Pro Ser Gly Gly His Thr Ser Tyr Ala Asp Ser Val Lys  
 565 570 575  
 Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu  
 580 585 590  
 Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala  
 595 600 605  
 Lys Ser Ser Gly Trp Tyr Glu Asp Tyr Phe Asp Tyr Trp Gly Gln Gly  
 610 615 620  
 Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly  
 625 630 635 640  
 Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser  
 645 650 655  
 Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Gln Ala  
 660 665 670  
 Ser Gln Asp Ile Thr Asn Tyr Leu Asn Trp Tyr Leu His Lys Pro Gly  
 675 680 685  
 Lys Ala Pro Lys Leu Leu Ile Tyr Asp Ala Ser Asn Leu Gln Thr Gly  
 690 695 700  
 Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Phe  
 705 710 715 720  
 Thr Ile Thr Ser Leu Gln Pro Glu Asp Phe Gly Thr Tyr Tyr Cys Gln  
 725 730 735  
 Gln Tyr Asp Thr Leu His Pro Ser Phe Gly Pro Gly Thr Thr Val Asp  
 740 745 750  
 Ile Lys

<210> 654

<211> 759

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 654

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg

				100					105					110			
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu		
		115						120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg		
	130						135				140						
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val		
145					150					155					160		
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr		
			165					170						175			
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
	180							185					190				
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
	195						200					205					
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
	210					215					220						
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225					230					235					240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
			245					250						255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro		
	260							265					270				
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro		
	275						280					285					
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr		
	290				295						300						
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn		
305				310					315						320		
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg		
			325					330						335			
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val		
			340					345					350				
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser		
	355						360					365					
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys		
	370					375					380						
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp		
385				390						395					400		
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe		
			405					410						415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu		
			420				425						430				
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe		
	435						440					445					
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly		
	450					455					460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr		
465				470					475						480		
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser		
			485					490						495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
			500					505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
	515						520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Pro	Tyr	Glu		
	530					535					540						
Met	Ala	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545					550					555					560		

Val	Ile	Gly	Pro	Ser	Gly	Gly	Tyr	Thr	Glu	Tyr	Ala	Asp	Ser	Val	Lys	
				565					570					575		
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
			580					585					590			
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	
		595					600					605				
Arg	Gly	Tyr	Asp	Phe	Trp	Ser	Gly	Tyr	Tyr	Asp	Ala	Phe	Asp	Ile	Trp	
	610					615				620						
Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	
625					630					635					640	
Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Ala	Leu	Thr	Gln	Pro	
				645					650					655		
Pro	Ser	Ala	Ser	Gly	Ser	Pro	Gly	Gln	Ser	Val	Thr	Ile	Ser	Cys	Thr	
			660					665					670			
Gly	Thr	Ser	Ser	Asp	Val	Gly	Val	Tyr	Asp	Ser	Val	Ser	Trp	Tyr	Gln	
		675					680					685				
Gln	His	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Met	Ile	Tyr	Asp	Val	Ser	Asp	
	690					695					700					
Arg	Pro	Ser	Gly	Val	Ser	Asn	Arg	Phe	Ser	Gly	Ser	Lys	Ser	Gly	Tyr	
705					710				715						720	
Thr	Ala	Ser	Leu	Thr	Ile	Ser	Ala	Leu	Gln	Ala	Glu	Asp	Glu	Ala	Asp	
				725				730					735			
Tyr	Tyr	Cys	Gly	Ser	Tyr	Arg	Ala	Ser	Ser	Ser	Tyr	Val	Phe	Gly	Thr	
			740					745					750			
Gly	Thr	Lys	Val	Thr	Val	Leu										
		755														

&lt;210&gt; 655

&lt;211&gt; 753

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 655

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

				165				170					175				
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly		
			180					185					190				
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser		
		195					200					205					
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser		
	210					215					220						
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser		
225					230					235					240		
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly		
				245					250						255		
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro		
			260					265					270				
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro		
		275					280					285					
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr		
	290					295					300						
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn		
305				310						315					320		
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg		
			325						330					335			
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val		
			340					345					350				
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser		
		355					360					365					
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys		
	370					375					380						
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp		
385				390						395					400		
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe		
			405						410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu		
			420					425					430				
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe		
		435					440					445					
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly		
	450					455					460						
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr		
465				470					475						480		
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser		
			485						490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu		
		500						505					510				
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
	515						520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr	Phe		
	530					535					540						
Met	Gly	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545				550						555					560		
Ser	Ile	Trp	Pro	Ser	Gly	Gly	Asn	Thr	Thr	Tyr	Ala	Asp	Ser	Val	Lys		
			565						570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
		580						585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
	595					600						605					
Arg	His	Val	Gly	Trp	Gly	Arg	Tyr	Phe	Asp	Tyr	Trp	Gly	Gln	Gly	Thr		
	610					615					620						

Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser  
 625 630 635 640  
 Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser  
 645 650 655  
 Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser  
 660 665 670  
 Gln Asn Ile Tyr His Tyr Leu His Trp Tyr Gln Gln Lys Pro Gly Lys  
 675 680 685  
 Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser Thr Leu Glu Ser Gly Val  
 690 695 700  
 Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Leu Thr Leu Thr  
 705 710 715 720  
 Ile Asn Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln  
 725 730 735  
 Thr Tyr Asp Thr Pro Leu Thr Phe Gly Gly Gly Ser Lys Val Glu Val  
 740 745 750  
 Lys

<210> 656  
 <211> 757  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 656  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser

225					230					235				240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr
				245					250					255
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro
			260					265					270	
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe
		275					280					285		Pro
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val
	290					295				300				Thr
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe
305					310					315				Asn
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro
			325						330					Arg
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr
			340					345					350	Val
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val
		355					360					365		Ser
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala
	370					375					380			Lys
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg
385				390						395				Asp
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly
			405						410					Phe
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro
			420					425					430	Glu
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser
	435						440					445		Phe
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln
	450					455					460			Gly
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His
465					470				475					Tyr
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn
			485						490					Ser
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser
		500						505					510	Glu
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
	515					520						525		Ser
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ala	Tyr
	530				535						540			Arg
Met	Ile	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
545					550				555					Ser
Tyr	Ile	Ser	Ser	Ser	Gly	Gly	Arg	Thr	Asp	Tyr	Ala	Asp	Ser	Val
			565					570					575	Lys
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
			580					585					590	Leu
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
	595					600						605		Thr
Thr	Gly	Gly	Leu	Arg	Tyr	Phe	Asp	Trp	Leu	Ala	Pro	Ser	Met	Asp
	610				615						620			Tyr
Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly
625					630				635					Ser
Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Ala	Leu	Thr
			645					650					655	Gln
Pro	Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln	Thr	Ala	Ser	Ile	Thr
		660					665					670		Cys
Ser	Gly	Asp	Lys	Leu	Gly	Glu	Lys	Phe	Ala	Ser	Trp	Tyr	Gln	Gln
		675				680						685		Arg



Pro Gly Gln Ser Pro Ile Leu Val Ile Tyr Gln Asp Ser Lys Arg Pro  
 690 695 700  
 Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asn Thr Ala  
 705 710 715 720  
 Thr Leu Thr Ile Ser Gly Thr Gln Thr Met Asp Glu Ala Asp Tyr Tyr  
 725 730 735  
 Cys Gln Ala Trp Gly Gly Ser Thr Ala Tyr Val Phe Gly Ser Gly Thr  
 740 745 750  
 Lys Val Thr Val Leu  
 755

<210> 657

<211> 752

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 657

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225 230 235 240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
 245 250 255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
 260 265 270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
 275 280 285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr

290		295		300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn				
305		310		315
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg				
	325		330	
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val				
	340		345	
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser				
	355		360	
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys				
	370		375	
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				
	405		410	
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420		425	
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435		440	
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450		455	
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				
	485		490	
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500		505	
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515		520	
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ala Tyr Ser				
	530		535	
Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Tyr Ile Tyr Pro Ser Gly Gly Ile Thr Thr Tyr Ala Asp Ser Val Lys				
	565		570	
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu				
	580		585	
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala				
	595		600	
Arg Glu Gly Gln Val Phe Asp Ile Trp Gly Gln Gly Thr Thr Val Thr				
	610		615	
Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly				
625		630		635
Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ala Thr Leu Ser Leu				
	645		650	
Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val				
	660		665	
Thr Thr Tyr Leu Ala Trp Tyr Gln Gln Arg Pro Gly Gln Ala Pro Arg				
	675		680	
Leu Leu Ile Tyr Asp Ala Ser Asn Arg Ala Thr Gly Ile Pro Ala Arg				
	690		695	
Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser				
705		710		715
Val Glu Pro Glu Asp Tyr Ala Val Tyr Tyr Cys Gln Gln Arg Ser Asn				
	725		730	
Trp Pro Pro Ser Ile Thr Phe Gly Gln Gly Thr Arg Leu Glu Ser Lys				
	740		745	
				750

<210> 658  
 <211> 754  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 658  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225 230 235 240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
 245 250 255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
 260 265 270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
 275 280 285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr  
 290 295 300  
 Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn  
 305 310 315 320  
 Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg  
 325 330 335  
 Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val  
 340 345 350  
 Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser  
 355 360 365  
 Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys

370		375		380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp				
385		390		395
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe				400
	405		410	415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu				
	420		425	430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe				
	435		440	445
Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly				
	450		455	460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr				
465		470		475
Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser				480
	485		490	495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu				
	500		505	510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser				
	515		520	525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr Ala				
	530		535	540
Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser				
545		550		555
Arg Ile Val Pro Ser Gly Gly Met Thr Ser Tyr Ala Asp Ser Val Lys				
	565		570	575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu				
	580		585	590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Thr				
	595		600	605
Thr Asp Asp Phe Trp Ser Gly Met Asp Val Trp Gly Gln Gly Thr Thr				
	610		615	620
Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly				
625		630		635
Gly Gly Gly Ser Gln Ser Ala Leu Thr Gln Pro Ala Ser Val Ser Gly				
	645		650	655
Ser Pro Gly Gln Ser Ile Thr Ile Ser Cys Thr Gly Thr Ser Ser Asp				
	660		665	670
Val Gly Ser Tyr Asn Leu Val Ser Trp Tyr Gln Gln His Pro Gly Lys				
	675		680	685
Ala Pro Lys Val Met Ile Tyr Glu Gly Ser Lys Arg Pro Ser Gly Val				
	690		695	700
Ser Asn Arg Phe Ser Gly Ser Lys Ser Gly Asn Thr Ala Ser Leu Thr				
705		710		715
Ile Ser Gly Leu Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Cys Ser				
	725		730	735
Tyr Ala Gly Arg Ser Thr Tyr Val Phe Gly Thr Gly Thr Lys Val Thr				
	740		745	750
Val Leu				

&lt;210&gt; 659

&lt;211&gt; 754

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 659

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Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1          5          10          15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
 20          25          30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
 35          40          45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
 50          55          60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
 65          70          75          80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
 85          90          95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
 100          105          110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
 115          120          125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
 130          135          140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
 145          150          155          160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
 165          170          175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
 180          185          190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
 195          200          205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser
 210          215          220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser
 225          230          235          240
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly
 245          250          255
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro
 260          265          270
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro
 275          280          285
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr
 290          295          300
Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn
 305          310          315          320
Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg
 325          330          335
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val
 340          345          350
Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser
 355          360          365
Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys
 370          375          380
Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp
 385          390          395          400
Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe
 405          410          415
Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu
 420          425          430
Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe

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[illegible]

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<210> 660
<211> 761
<212> PRT
<213> Artificial Sequence
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<220>  
<223> Made in a lab

<400> 660

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			

Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
50						55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65						70				75					80
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145					150					155					160
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
		195					200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225					230					235					240
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
				340				345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
				485					490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu

				500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser		
		515					520					525					
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ala	Tyr	Thr		
	530					535					540						
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser		
545				550					555						560		
Ser	Ile	Tyr	Pro	Ser	Gly	Gly	Thr	Thr	Pro	Tyr	Ala	Asp	Ser	Val	Lys		
			565						570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
		580						585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Thr	Tyr	Tyr	Cys	Ala		
	595						600					605					
Arg	Val	Val	Asn	Ile	Asp	Phe	Trp	Ser	Gly	Tyr	Asn	Met	Arg	Ser	Ala		
	610					615					620						
Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly		
625				630					635						640		
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile		
			645					650						655			
Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala	Ser	Val	Gly	Asp	Thr		
			660					665					670				
Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Val	Ser	Asp	Trp	Leu	Ala		
	675						680					685					
Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Asn	Leu	Leu	Ile	Tyr	Gly		
	690					695				700							
Ala	Ser	Ser	Leu	Gln	Thr	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Gly	Gly		
705				710					715						720		
Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp		
			725						730					735			
Phe	Ala	Thr	Tyr	Cys	Gln	Gln	Ala	His	Ser	Phe	Pro	Phe	Thr	Phe			
			740				745					750					
Gly	Gly	Gly	Thr	Arg	Val	Glu	Ile	Lys									
	755						760										

&lt;210&gt; 661

&lt;211&gt; 757

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 661

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



Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	
		115					120					125				
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	
	130					135					140					
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	
145					150					155					160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	
			165					170						175		
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
		180						185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
		195					200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310					315						320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330					335			
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
		340						345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390					395						400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435						440				445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470					475						480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
		500						505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
		515					520					525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Asp	Tyr	Leu	
	530					535					540					
Met	Gly	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550					555						560	
Val	Ile	Ser	Ser	Ser	Gly	Gly	Pro	Thr	Ala	Tyr	Ala	Asp	Ser	Val	Lys	

				565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu		
			580					585					590				
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala		
		595					600					605					
Arg	Val	Gly	Leu	Asp	Tyr	Gly	Ile	Leu	Gly	Ala	Phe	Asp	Ile	Trp	Gly		
	610					615					620						
Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly		
625					630					635					640		
Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser		
			645					650						655			
Pro	Ala	Thr	Leu	Ser	Val	Ser	Pro	Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys		
			660					665					670				
Arg	Ala	Ser	Gln	Asn	Ile	Asn	Thr	His	Leu	Ala	Trp	Tyr	Gln	Gln	Lys		
		675					680					685					
Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu	Ile	Phe	Gly	Ala	Ser	Thr	Arg	Ala		
	690					695				700							
Thr	Gly	Ile	Pro	Ala	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Glu	Phe		
705					710					715					720		
Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Ser	Glu	Asp	Phe	Thr	Val	Tyr	Tyr		
				725				730						735			
Cys	Gln	Gln	Tyr	Gly	Ser	Ser	Leu	Thr	Trp	Thr	Phe	Gly	Gln	Gly	Thr		
			740					745					750				
Lys	Val	Glu	Ile	Lys													
		755															

&lt;210&gt; 662

&lt;211&gt; 755

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 662

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

Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	
			180					185					190			
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	
		195					200					205				
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	
	210					215					220					
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	
225					230					235					240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	
			245					250					255			
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	
		260						265					270			
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	
		275					280					285				
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	
	290					295					300					
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	
305				310						315					320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	
			325					330						335		
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	
			340					345					350			
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	
		355					360					365				
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	
	370					375					380					
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	
385				390						395					400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	
			405					410					415			
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	
			420					425					430			
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	
		435					440					445				
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	
	450					455					460					
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	
465				470						475					480	
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	
			485					490					495			
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	
			500					505					510			
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	
	515						520					525				
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Glu	Tyr	Gly	
	530					535					540					
Met	Ser	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	
545				550						555					560	
Ser	Ile	Arg	Ser	Ser	Gly	Gly	Trp	Thr	Lys	Tyr	Ala	Asp	Ser	Val	Lys	
			565					570					575			
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	
			580					585					590			
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	
	595						600					605				
Lys	Gly	Leu	Gly	Ala	Thr	Ser	Gly	Glu	Phe	Asp	Tyr	Trp	Gly	Gln	Gly	
	610					615					620					
Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	

```

625          630          635          640
Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Gly
645          650          655
Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala
660          665          670
Ser Gln Ser Val Ser Gly Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Leu
675          680          685
Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Ser Ser Arg Ala Thr
690          695          700
Gly Thr Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Glu Phe Thr
705          710          715          720
Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys
725          730          735
Leu Gln His Asn Ser Tyr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val
740          745          750
Glu Ile Lys
755

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<210> 663
<211> 755
<212> PRT
<213> Artificial Sequence

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<220>
<223> Made in a lab

```

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<400> 663
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
1      5      10
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
20     25     30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
35     40     45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
50     55     60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
65     70     75     80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
85     90     95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
100    105    110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
115    120    125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
130    135    140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
145    150    155    160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
165    170    175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
180    185    190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
195    200    205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser
210    215    220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser
225    230    235    240

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Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305					310					315					320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
			325						330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385					390					395					400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
			405						410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
	435						440					445			
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly
	450					455					460				
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr
465					470					475					480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser
			485						490					495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu
			500					505					510		
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser
	515					520						525			
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Lys	Tyr	Asp
	530					535					540				
Met	Trp	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser
545					550					555					560
Ser	Ile	Trp	Pro	Ser	Gly	Gly	Trp	Thr	Ser	Tyr	Ala	Asp	Ser	Val	Lys
			565						570					575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala
	595						600					605			
Lys	Ser	Arg	Gly	Ser	Pro	Trp	Tyr	Gly	Asp	Phe	Asp	His	Trp	Gly	Gln
	610					615					620				
Gly	Thr	Leu	Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly
625					630					635					640
Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gln	Ser	Glu	Leu	Thr	Gln	Pro	Pro	Ser
			645						650					655	
Val	Ser	Val	Ala	Pro	Gly	Glu	Thr	Ala	Thr	Val	Thr	Cys	Gly	Gly	Ile
			660					665				670			
Asn	Ile	Gly	Ser	Lys	Ser	Val	His	Trp	Tyr	Gln	Gln	Lys	Pro	Asp	Gln
	675						680					685			
Ala	Pro	Val	Val	Val	Met	Tyr	Phe	Asp	Ser	Asp	Arg	Pro	Ser	Gly	Ile

690		695		700
Pro Glu Arg Phe Ser Gly Ser Asn Ser Gly Asp Thr Ala Thr Leu Thr				
705		710		715
Ile Ser Arg Val Glu Ala Gly Asp Glu Ala Asp Tyr Tyr Cys Gln Val				
	725		730	735
Trp Asp Ser Arg Ser Asp Gln Tyr Val Phe Gly Phe Gly Thr Lys Val				
	740		745	750
Thr Val Leu				
755				

&lt;210&gt; 664

&lt;211&gt; 752

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 664

Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser				
1		5		10
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr				
	20		25	30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln				
	35		40	45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys				
	50		55	60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp				
65		70		75
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys				
	85		90	95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg				
	100		105	110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu				
	115		120	125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg				
	130		135	140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val				
145		150		155
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr				
	165		170	175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly				
	180		185	190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser				
	195		200	205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser				
	210		215	220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser				
225		230		235
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly				
	245		250	255
Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro				
	260		265	270
Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro				
	275		280	285
Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr				
	290		295	300

Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	305	310	315	320
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	325	330	335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	340	345	350	
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	355	360	365	
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	370	375	380	
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	385	390	395	400
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	405	410	415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	420	425	430	
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	435	440	445	
Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	450	455	460	
Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	465	470	475	480
Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	485	490	495	
Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	500	505	510	
Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	515	520	525	
Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Met	Tyr	Thr	530	535	540	
Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val	Ser	545	550	555	560
Trp	Ile	Ser	Pro	Ser	Gly	Gly	Trp	Thr	Lys	Tyr	Ala	Asp	Ser	Val	Lys	565	570	575	
Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu	580	585	590	
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys	Ala	595	600	605	
Val	Ser	Pro	Trp	Gly	Gly	Pro	Phe	Asp	Tyr	Trp	Gly	Gln	Gly	Thr	Leu	610	615	620	
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	625	630	635	640
Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Thr	Leu	645	650	655	
Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	660	665	670	
Ser	Ile	Ser	Ser	Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	675	680	685	
Pro	Lys	Leu	Leu	Ile	Tyr	Lys	Ala	Ser	Thr	Leu	Glu	Ser	Gly	Val	Pro	690	695	700	
Ser	Arg	Phe	Ser	Ala	Gly	Gly	Ser	Gly	Thr	Glu	Phe	Thr	Leu	Thr	Ile	705	710	715	720
Ser	Ser	Leu	Gln	Pro	Asp	Asp	Phe	Gly	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	725	730	735	
Asn	Ser	Tyr	Ser	Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Ile	740	745	750	

<210> 665  
 <211> 757  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 665  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr  
 20 25 30  
 Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln  
 35 40 45  
 Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys  
 50 55 60  
 Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp  
 65 70 75 80  
 Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys  
 85 90 95  
 Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg  
 100 105 110  
 Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu  
 115 120 125  
 Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg  
 130 135 140  
 Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val  
 145 150 155 160  
 Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr  
 165 170 175  
 Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly  
 180 185 190  
 Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser  
 195 200 205  
 Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser  
 210 215 220  
 Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser  
 225 230 235 240  
 Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly  
 245 250 255  
 Asp Thr Gly Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro  
 260 265 270  
 Cys Pro Ala Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro  
 275 280 285  
 Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr  
 290 295 300  
 Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn  
 305 310 315 320  
 Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg  
 325 330 335  
 Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val  
 340 345 350  
 Leu His Gln Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser  
 355 360 365  
 Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys  
 370 375 380



Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp  
 385 390 395 400  
 Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe  
 405 410 415  
 Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu  
 420 425 430  
 Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe  
 435 440 445  
 Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly  
 450 455 460  
 Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr  
 465 470 475 480  
 Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser  
 485 490 495  
 Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu  
 500 505 510  
 Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser  
 515 520 525  
 Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gln Tyr Phe  
 530 535 540  
 Met Met Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser  
 545 550 555 560  
 Ser Ile Ser Pro Ser Gly Gly Tyr Thr Gln Tyr Ala Asp Ser Val Lys  
 565 570 575  
 Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu  
 580 585 590  
 Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala  
 595 600 605  
 Lys Thr Gly Asp Tyr Thr Arg Tyr Tyr Ser Tyr Gly Met Asp Val Trp  
 610 615 620  
 Gly Gln Gly Thr Leu Val Thr Val Ser Ser Gly Gly Gly Gly Ser Gly  
 625 630 635 640  
 Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln  
 645 650 655  
 Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr  
 660 665 670  
 Cys Arg Ala Ser Gln Ser Ile Ser Ser Tyr Leu Asn Trp Tyr Gln Gln  
 675 680 685  
 Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser Ser Leu  
 690 695 700  
 Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp  
 705 710 715 720  
 Phe Thr Leu Ile Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr  
 725 730 735  
 Tyr Cys Gln Gln Ser Tyr Ser Thr Pro Leu Thr Phe Gly Gly Gly Thr  
 740 745 750  
 Lys Val Glu Ile Lys  
 755

<210> 666

<211> 758

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

&lt;400&gt; 666

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr
			20					25					30		
Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln
		35					40					45			
Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys
	50					55					60				
Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp
65				70						75				80	
Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys
				85					90					95	
Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg
			100					105					110		
Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu
		115					120					125			
Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg
	130					135					140				
Pro	Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val
145				150						155				160	
Cys	Lys	Pro	Cys	Ala	Pro	Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr
				165					170					175	
Asp	Ile	Cys	Arg	Pro	His	Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly
			180					185					190		
Asn	Ala	Ser	Met	Asp	Ala	Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser
	195						200					205			
Met	Ala	Pro	Gly	Ala	Val	His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser
	210					215					220				
Gln	His	Thr	Gln	Pro	Thr	Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser
225				230						235				240	
Phe	Leu	Leu	Pro	Met	Gly	Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly
				245					250					255	
Asp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro
			260					265					270		
Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro
		275					280					285			
Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr
	290					295					300				
Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn
305				310						315				320	
Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg
				325					330					335	
Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val
			340					345					350		
Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser
		355					360					365			
Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys
	370					375					380				
Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp
385				390						395				400	
Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe
				405					410					415	
Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu
			420					425					430		
Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe
		435					440					445			

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Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly
  450                      455                      460
Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr
465                      470                      475                      480
Thr Gln Lys Ser      Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser
                      485                      490                      495
Ser Leu Asn Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Glu
      500                      505                      510
Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
      515                      520                      525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gly Tyr Ser
      530                      535                      540
Met Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                      550                      555                      560
Gly Ile Tyr Ser      Ser Gly Gly Trp Thr Trp Tyr Ala Asp Ser Val Lys
                      565                      570                      575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
      580                      585                      590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Thr
      595                      600                      605
Arg Asp Gln Ser Phe Ser Tyr Asp Ser Ser Leu Asp Ala Phe Asp Ile
      610                      615                      620
Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly Ser
625                      630                      635                      640
Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr
      645                      650                      655
Gln Ser Pro Ser Ser Leu Ser Ala Ser Val Gly Asp Arg Val Thr Ile
      660                      665                      670
Thr Cys Arg Ala Ser Gln Ser Ile Asn Asn Tyr Leu Asn Trp Tyr Gln
      675                      680                      685
Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Thr Ala Ser Thr
      690                      695                      700
Leu His Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr
705                      710                      715                      720
Asp Phe Thr Leu Thr Ile Ser Asn Leu Gln Pro Glu Asp Phe Ala Thr
      725                      730                      735
Tyr Tyr Cys Gln Gln Ser Tyr Ala Thr Leu Arg Thr Phe Gly Gln Gly
      740                      745                      750
Thr Lys Val Glu Ile Lys
      755

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&lt;210&gt; 667

&lt;211&gt; 760

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 667

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Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
  1                      5                      10                      15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
      20                      25                      30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
      35                      40                      45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys

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50		55		60
Val Phe Cys Thr Lys Thr Ser Asp Thr	Val Cys Asp Ser Cys Glu Asp			
65	70	75	80	
Ser Thr Tyr Thr Gln Leu Trp Asn Trp	Val Pro Glu Cys Leu Ser Cys			
	85	90	95	
Gly Ser Arg Cys Ser Ser Asp Gln Val	Glu Thr Gln Ala Cys Thr Arg			
	100	105	110	
Glu Gln Asn Arg Ile Cys Thr Cys Arg	Pro Gly Trp Tyr Cys Ala Leu			
	115	120	125	
Ser Lys Gln Glu Gly Cys Arg Leu Cys	Ala Pro Leu Arg Lys Cys Arg			
	130	135	140	
Pro Gly Phe Gly Val Ala Arg Pro Gly	Thr Glu Thr Ser Asp Val Val			
145	150	155	160	
Cys Lys Pro Cys Ala Pro Gly Thr Phe	Ser Asn Thr Thr Ser Ser Thr			
	165	170	175	
Asp Ile Cys Arg Pro His Gln Ile Cys	Asn Val Val Ala Ile Pro Gly			
	180	185	190	
Asn Ala Ser Met Asp Ala Val Cys Thr	Ser Thr Ser Pro Thr Arg Ser			
	195	200	205	
Met Ala Pro Gly Ala Val His Leu Pro	Gln Pro Val Ser Thr Arg Ser			
	210	215	220	
Gln His Thr Gln Pro Thr Pro Glu Pro	Ser Thr Ala Pro Ser Thr Ser			
225	230	235	240	
Phe Leu Leu Pro Met Gly Pro Ser Pro	Pro Ala Glu Gly Ser Thr Gly			
	245	250	255	
Asp Thr Gly Glu Pro Lys Ser Ser Asp	Lys Thr His Thr Cys Pro Pro			
	260	265	270	
Cys Pro Ala Pro Glu Ala Ala Gly Ala	Pro Ser Val Phe Leu Phe Pro			
	275	280	285	
Pro Lys Pro Lys Asp Thr Leu Met Ile	Ser Arg Thr Pro Glu Val Thr			
	290	295	300	
Cys Val Val Val Asp Val Ser His Glu	Asp Pro Glu Val Lys Phe Asn			
305	310	315	320	
Trp Tyr Val Asp Gly Val Glu Val His	Asn Ala Lys Thr Lys Pro Arg			
	325	330	335	
Glu Glu Gln Tyr Asn Ser Thr Tyr Arg	Val Val Ser Val Leu Thr Val			
	340	345	350	
Leu His Gln Asp Trp Leu Asn Gly Lys	Ala Tyr Ala Cys Ala Val Ser			
	355	360	365	
Asn Lys Ala Leu Pro Ala Pro Ile Glu	Lys Thr Ile Ser Lys Ala Lys			
	370	375	380	
Gly Gln Pro Arg Glu Pro Gln Val Tyr	Thr Leu Pro Pro Ser Arg Asp			
385	390	395	400	
Glu Leu Thr Lys Asn Gln Val Ser Leu	Thr Cys Leu Val Lys Gly Phe			
	405	410	415	
Tyr Pro Ser Asp Ile Ala Val Glu Trp	Glu Ser Asn Gly Gln Pro Glu			
	420	425	430	
Asn Asn Tyr Lys Thr Thr Pro Pro Val	Leu Asp Ser Asp Gly Ser Phe			
	435	440	445	
Phe Leu Tyr Ser Lys Leu Thr Val Asp	Lys Ser Arg Trp Gln Gln Gly			
	450	455	460	
Asn Val Phe Ser Cys Ser Val Met His	Glu Ala Leu His Asn His Tyr			
465	470	475	480	
Thr Gln Lys Ser Leu Ser Leu Ser Pro	Gly Gln Arg His Asn Asn Ser			
	485	490	495	
Ser Leu Asn Thr Arg Thr Gln Lys Ala	Arg His Ser Pro Asn Ser Glu			
	500	505	510	

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Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
    515                520                525
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr Gly
    530                535                540
Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
545                550                555                560
Ser Ile Val Pro Ser Gly Gly Leu Thr Arg Tyr Ala Asp Ser Val Lys
    565                570                575
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
    580                585                590
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Thr Tyr Tyr Cys Ala
    595                600                605
Arg Leu Ala Tyr Tyr Asp Phe Trp Ser Gly Arg Asp Ala Phe Asp Ile
    610                615                620
Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser Gly Gly Gly Gly Ser
625                630                635                640
Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Ser Val Leu Thr Gln
    645                650                655
Pro Ala Ser Val Ser Gly Ser Pro Gly Gln Ser Ile Thr Ile Ser Cys
    660                665                670
Ser Gly Ser Ser Ser Asp Val Gly His Tyr Asp Tyr Val Ser Trp Tyr
    675                680                685
Gln Gln His Pro Gly Lys Ala Pro Lys Leu Met Ile Tyr Asp Val Ser
    690                695                700
Asn Arg Pro Ser Gly Val Ser Asn Arg Phe Ser Gly Ser Lys Ser Gly
705                710                715                720
Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu Gln Ala Asp Asp Glu Ala
    725                730                735
Glu Tyr Tyr Cys Ser Ser Tyr Thr Ser Ser Gly Thr Arg Val Phe Gly
    740                745                750
Thr Gly Thr Lys Val Thr Val Leu
    755                760

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<210> 668
<211> 752
<212> PRT
<213> Artificial Sequence

```

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<220>
<223> Made in a lab

```

```

<400> 668
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1          5          10          15
Val Ile Met Ser Arg Gly Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
    20          25          30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
    35          40          45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
    50          55          60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
65          70          75          80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
    85          90          95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
    100         105         110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu

```

[illegible]

Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr	Leu
			580					585					590		
Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Thr	Tyr	Tyr	Cys	Ala
		595					600					605			
Thr	Arg	Thr	Trp	Ala	Asp	Ala	Phe	Asp	Val	Trp	Gly	Gln	Gly	Thr	Thr
	610				615						620				
Val	Thr	Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly
625					630					635					640
Gly	Gly	Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Thr	Leu
			645					650						655	
Ser	Ala	Ser	Val	Gly	Asp	Arg	Val	Thr	Phe	Thr	Cys	Arg	Ala	Ser	Gln
			660					665					670		
Ser	Val	Asn	Asn	Trp	Val	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala
		675					680					685			
Pro	Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro
	690					695					700				
Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile
705					710					715					720
Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser
			725					730					735		
Tyr	Ser	Met	Pro	Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys
			740				745						750		

&lt;210&gt; 669

&lt;211&gt; 752

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 669

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

		195				200				205					
Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly
	210					215					220				
Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala
225					230					235					240
Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Tyr	Ser	Thr	Pro	Arg	Thr	Phe	Gly	Gln
			245						250					255	
Gly	Thr	Lys	Val	Glu	Ile	Lys	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys
		260						265				270			
Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro
	275						280					285			
Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser
	290					295					300				
Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp
305					310					315					320
Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn
			325						330					335	
Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val
		340						345				350			
Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala
	355					360					365				
Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys
	370					375				380					
Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr
385					390					395					400
Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr
			405						410					415	
Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu
		420						425				430			
Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu
	435					440					445				
Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys
	450					455					460				
Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu
465					470					475					480
Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly
			485					490						495	
Gln	Arg	His	Asn	Asn	Ser	Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg
		500						505				510			
His	Ser	Pro	Asn	Ser	Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	Ala
	515					520					525				
Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	Thr
	530					535					540				
Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	Val
545					550					555					560
Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	Ser
			565						570					575	
Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	Gly
		580						585				590			
Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	Glu
	595					600						605			
Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	Ser
	610					615					620				
Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	Pro
625					630					635					640
Gly	Phe	Gly	Val	Ala	Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	Cys
			645						650					655	



```

Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr Asp
      660                      665                      670
Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly Asn
      675                      680                      685
Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser Met
      690                      695                      700
Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser Gln
705                      710                      715                      720
His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser Phe
      725                      730                      735
Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly Asp
      740                      745                      750

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<210> 670
<211> 759
<212> PRT
<213> Artificial Sequence

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<220>
<223> Made in a lab

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```

<400> 670
Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1          5          10          15
Val Ile Met Ser Arg Gly Glu Val Gln Leu Leu Glu Ser Gly Gly Gly
 20          25          30
Leu Val Gln Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly
 35          40          45
Phe Thr Phe Ser Trp Tyr Leu Met His Trp Val Arg Gln Ala Pro Gly
 50          55          60
Lys Gly Leu Glu Trp Val Ser Gly Ile Trp Pro Ser Gly Gly His Thr
 65          70          75          80
Leu Tyr Ala Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn
      85          90          95
Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp
      100          105          110
Thr Ala Val Tyr Tyr Cys Ala Arg Glu Pro Leu Leu Trp Phe Gly Glu
      115          120          125
Leu Ser Tyr Asn Trp Phe Asp Pro Trp Gly Gln Gly Thr Leu Val Thr
      130          135          140
Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly
145          150          155          160
Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Val Ser Ala
      165          170          175
Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile
      180          185          190
Ser Ser Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
      195          200          205
Leu Leu Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg
      210          215          220
Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser
225          230          235          240
Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Ser
      245          250          255
Phe Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys Thr Gly
      260          265          270
Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala

```

		275					280					285			
Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro
	290					295					300				
Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val
305					310					315					320
Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val
				325					330					335	
Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln
			340					345					350		
Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln
		355					360					365			
Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala
	370					375					380				
Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro
385					390					395					400
Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr
				405					410					415	
Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser
			420					425					430		
Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr
		435					440					445			
Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr
	450					455					460				
Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe
465					470					475					480
Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys
				485					490					495	
Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	Ser	Leu	Asn
			500					505					510		
Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Leu	Pro	Ala	Gln
		515					520					525			
Val	Ala	Phe	Thr	Pro	Tyr	Ala	Pro	Glu	Pro	Gly	Ser	Thr	Cys	Arg	Leu
	530					535					540				
Arg	Glu	Tyr	Tyr	Asp	Gln	Thr	Ala	Gln	Met	Cys	Cys	Ser	Lys	Cys	Ser
545					550					555					560
Pro	Gly	Gln	His	Ala	Lys	Val	Phe	Cys	Thr	Lys	Thr	Ser	Asp	Thr	Val
				565					570					575	
Cys	Asp	Ser	Cys	Glu	Asp	Ser	Thr	Tyr	Thr	Gln	Leu	Trp	Asn	Trp	Val
			580					585					590		
Pro	Glu	Cys	Leu	Ser	Cys	Gly	Ser	Arg	Cys	Ser	Ser	Asp	Gln	Val	Glu
		595					600					605			
Thr	Gln	Ala	Cys	Thr	Arg	Glu	Gln	Asn	Arg	Ile	Cys	Thr	Cys	Arg	Pro
	610					615					620				
Gly	Trp	Tyr	Cys	Ala	Leu	Ser	Lys	Gln	Glu	Gly	Cys	Arg	Leu	Cys	Ala
625					630					635					

Thr Ala Pro Ser Thr Ser Phe Leu Leu Pro Met Gly Pro Ser Pro Pro  
 740 745 750  
 Ala Glu Gly Ser Thr Gly Asp  
 755

<210> 671  
 <211> 2270  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 671  
 aagcttgccg ccatggattt tcaagtgcag attttcagct tcctgctaata cagtgtttca 60  
 gtcataatgt cgcgaggatt gcccgcccag gtggcattta caccctacgc cccggagccc 120  
 gggagcacat gccggctcag agaatactat gaccagacag ctgagatgtg ctgcagcaaa 180  
 tgctcgccgg gccaacatgc aaaagtcttc tgtaccaaga cctcggacac cgtgtgtgac 240  
 tcctgtgagg acagcacata caccagctc tgggaactggg ttcccgagtg cttgagctgt 300  
 ggctcccgtc gtagctctga ccaggtggaa actcaagcct gcaactcggga acagaaccgc 360  
 atctgcacct gcaggcccgg ctggtactgc gcgtgagca agcaggagggt gtgccggctg 420  
 tgcgcgccgc tgcgcaagtg ccgcccgggc ttcggcgtgg ccagaccagg aactgaaaca 480  
 tcagacgtgg tgtgcaagcc ctgtgccccg gggacgttct ccaacacgac ttcattccacg 540  
 gatatttgca ggccccacca gatctgtaac gtggtggcca tccctgggaa tgcaagcatg 600  
 gatgcagtct gcacgtccac gtccccacc cggagtatgg ccccgagggc agtacactta 660  
 cccagccag tgtccacacg atcccaacac acgcagccaa ctccagaacc cagcactgct 720  
 ccaagcacct ccttctctgt cccaatgggc cccagccccc cagctgaagg gagcactggc 780  
 gacaccggtg agcccaaate ttctgacaaa actcacacat gccaccgtg cccagcacct 840  
 gaagccgctg gtagcccgct agtcttcttc ttcccccaa aaccaagga caccctcatg 900  
 atctcccga cccctgaggt cacatgcgtg gtggtggacg tgagccacga agaccctgag 960  
 gtcaagtcca actggtacgt ggacggcgtg gagggtgcata atgccaagac aaagccgctg 1020  
 gaggagcagt acaacagcac gtaccgtgtg gtcagcgtcc tcaccgtcct gcaccaggac 1080  
 tggctgaatg gcaaggcgta cgcgtgcgct gtctccaaca aagccctccc agccccatc 1140  
 gagaaaacca tctccaaagc caaaggcgag ccccgagaac cacagggtgta caccctgccc 1200  
 ccatcccggg atgagctgac caagaaccag gtcagcctga cctgcctggt caaaggcttc 1260  
 tatccaagcg acatcgccgt ggagtgggag agcaatgggc agccggagaa caactacaag 1320  
 accacgcctc ccgtgctgga ctccgacggc tccttcttcc tctacagcaa gctcaccgtg 1380  
 gacaagagca ggtggcagca ggggaacgtc ttctcatgct ccgtgatgca tgaggctctg 1440  
 cacaaccact acacgcagaa gagcctctcc ctgtctccgg gtcagaggca caacaattct 1500  
 tccctgaata caagaactca gaaagcacgt cattctccga attctgaagt tcaattgtta 1560  
 gagtctggtg gcggtcttgt tcagcctggt ggttctttac gtctttcttg cgtgcttcc 1620  
 ggattcactt tctctaagta ctctatgcat tgggttcgcc aagctcctgg taaaggtttg 1680  
 gagtgggttt ctctatctg gccttctggt ggctggacta cttatgctga ctccgttaaa 1740  
 ggtcgcttca ctatctctag agacaactct aagaatactc tctacttgca gatgaacagc 1800  
 ttaagggtctg aggacacggc cgtgtattac tgtgcgagag gtggttatta ctacgccctt 1860  
 gactactggg gccagggaac cctggtcacc gtctcaagcg gtggcggcgg ttccgggggt 1920  
 ggcggaagtg gaggtggagg gtagtcagag gaattgactc agccaccctc agtgtccgtg 1980  
 tccccaggac agacagccac catcacctgc tctggagaaa aattggggga tatatatgct 2040  
 tcctggtatc agcagaagcc aggccagtc ccggtcttgg tcatctatca agataccaag 2100  
 cggccctcag ggatccctgt gcgattctct ggctccaact ctgggaacac agccactctg 2160  
 accatcagcg ggaccaggc tatggatgag gctgactatt actgtcaggc gtgggacagc 2220  
 agcacggtat tcggcgagg gaccaagctg accgtcctat gagcgggcgc 2270

<210> 672  
 <211> 2297  
 <212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 672

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gggagcacat	gcccgtcag	agaatactat	gaccagacag	ctcagatgtg	ctgcagcaaa	180
tgctcgccgg	gccaacatgc	aaaagtcttc	tgtaccaaga	cctcggacac	cgtgtgtgac	240
tcctgtgagg	acagcacata	caccagctc	tggaactggg	ttcccagagt	cttgagctgt	300
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tgcgcgccgc	tgcgcaagtg	ccgcccgggc	ttcggcgtgg	ccagaccagg	aactgaaaca	480
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gatattttgca	ggccccacca	gatctgtaac	gtggtggcca	tccttgggaa	tgcaagcatg	600
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gacaccgggtg	agcccaaate	ttctgacaaa	actcacacat	gccaccgtg	cccagcacct	840
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atctcccggg	cccctgaggt	cacatgcgtg	gtggtggacg	tgagccacga	agaccctgag	960
gtcaagttca	actggtacgt	ggacggcgtg	gaggtgcata	atgccaaagac	aaagccgcgg	1020
gaggagcagt	acaacagcac	gtaccgtgtg	gtcagcgtcc	tcaccgtcct	gcaccaggac	1080
tggttgaatg	gcaaggcgta	cgctgctggc	gtctccaaca	aagccctccc	agcccccatc	1140
gagaaaaacca	tctccaaagc	caaagggcag	ccccgagaac	cacaggtgta	cacctgtccc	1200
ccatcccggg	atgagctgac	caagaaccag	gtcagcctga	cctgcctggt	caaaggcttc	1260
tatccaagcg	acatcgccgt	ggagtgggag	agcaatgggc	agccggagaa	caactacaag	1320
accacgcctc	ccgtgctgga	ctccgacggc	tccttcttcc	tctacagcaa	gtcaccgtg	1380
gacaagagca	gggtggcagca	ggggaacgtc	ttctcatgct	ccgtgatgca	tgaggctctg	1440
cacaaccact	acacgcagaa	gagcctctcc	ctgtctccgg	gtcagaggca	caacaattct	1500
tccttgaata	caagaactca	gaaagcacgt	cattctccga	attctgaagt	tcaattgtta	1560
gagtctgggtg	gcggtcttgt	tcagcctggt	ggttcttttac	gtctttcttg	cgctgcttcc	1620
ggattcactt	tctcttttta	ccagatgcat	tgggttcgcc	aagctcctgg	taaaggtttg	1680
gagtgggttt	ctggtatcta	tccttctggt	ggctatacta	agtatgctga	ctccgttaaa	1740
ggctcgcttca	ctatctctag	agacaactct	aagaatactc	tctacttgca	gatgaacagc	1800
ttaagggctg	aggacactgc	agtctactat	tgtgcgagag	atctagggta	cggcagcagc	1860
tggtactact	ttgactactg	gggccaggga	accctggtca	ccgtctcaag	cgggtggcggc	1920
ggttcggggg	gtggcggaag	tggaggtgga	gggagtcaga	gcgctttgac	tcagcctgcc	1980
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<210> 673

<211> 2276

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 673

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&lt;210&gt; 674

&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 674

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&lt;210&gt; 675

&lt;211&gt; 2306

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 675

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&lt;210&gt; 676

&lt;211&gt; 2300

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 676

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&lt;210&gt; 677

&lt;211&gt; 2285

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 677

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gccgc						2285

&lt;210&gt; 678

&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 678

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&lt;210&gt; 679

&lt;211&gt; 2376

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 679

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&lt;210&gt; 680

&lt;211&gt; 2306

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 680

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&lt;210&gt; 681

&lt;211&gt; 2288

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 681

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gcggccgc						2288

&lt;210&gt; 682

&lt;211&gt; 2389

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 682

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&lt;210&gt; 683

&lt;211&gt; 2290

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 683

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&lt;210&gt; 684

&lt;211&gt; 2300

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 684

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&lt;210&gt; 685

&lt;211&gt; 2376

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 685

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&lt;210&gt; 686

<211> 2285  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 686

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 <211> 2361  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 687



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&lt;210&gt; 688

&lt;211&gt; 2276

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 688

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&lt;210&gt; 689

&lt;211&gt; 2300

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 689

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gagatcaaat	gagcggccgc					2300

&lt;210&gt; 690

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 690

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&lt;210&gt; 691

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 691

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aaatgagcgg	ccgc					2294

&lt;210&gt; 692

&lt;211&gt; 2297

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 692

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 <211> 2303  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 693

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 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 694

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&lt;210&gt; 695

&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 695

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&lt;210&gt; 696

&lt;211&gt; 2306

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 696

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&lt;210&gt; 697

&lt;211&gt; 2371

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 697

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&lt;210&gt; 698

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 698

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&lt;210&gt; 699

&lt;211&gt; 2359

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 699

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2359

&lt;210&gt; 700

&lt;211&gt; 2288

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 700

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&lt;210&gt; 701

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Made in a lab

<400> 701

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<210> 702

<211> 2301

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 702

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&lt;210&gt; 703

&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 703

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&lt;210&gt; 704

&lt;211&gt; 2288

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 704

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&lt;210&gt; 705

&lt;211&gt; 2291

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 705

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&lt;210&gt; 706

&lt;211&gt; 2264

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 706

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 <211> 2300  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 707  
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 <211> 2378  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

&lt;400&gt; 708

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&lt;210&gt; 709

&lt;211&gt; 2369

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 709

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&lt;211&gt; 2291

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 710

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&lt;210&gt; 711

&lt;211&gt; 2299

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 711

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&lt;210&gt; 712

&lt;211&gt; 2362

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 712

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&lt;210&gt; 713

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 713

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 <211> 2308  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

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 <211> 2288  
 <212> DNA



<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 715

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<210> 716

<211> 2282

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 716

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gc 2282

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<210> 717

<211> 2285

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 717

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gccgc						2285

&lt;210&gt; 718

&lt;211&gt; 2300

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 718

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&lt;211&gt; 2282

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 719

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gc						2282

&lt;210&gt; 720

&lt;211&gt; 2294

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 720

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&lt;210&gt; 721

&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 721

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&lt;210&gt; 722

&lt;211&gt; 2285

&lt;212&gt; DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 722

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gccgc						2285

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<211> 2285

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 723

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<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 724

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&lt;211&gt; 2383

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 725

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&lt;211&gt; 2412

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 726

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&lt;211&gt; 2288

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 727

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&lt;211&gt; 2279

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 728

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 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 729

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 <211> 2297  
 <212> DNA  
 <213> Artificial Sequence

<220>

<223> Made in a lab

<400> 730

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<211> 2303

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 731

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&lt;210&gt; 732

&lt;211&gt; 2326

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 732

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&lt;211&gt; 2280

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 733

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gaagattttg	caacttacta	ctgtcaacag	agttacagta	cccctcggac	gttcggccaa	780
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accgtcctgc	accaggactg	gctgaatggc	aaggcgtacg	cgtgcgcggg	ctccaacaaa	1140
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tacagcaagc	tcaccgtgga	caagagcagg	tggcagcagg	ggaacgtctt	ctcatgctcc	1440
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cagaggcaca	acaattcttc	cctgaatata	agaactcaga	aagcacgtca	ttctccgaat	1560
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caccagatct	gtaacgtggg	ggccatccct	gggaatgcaa	gcatggatgc	agtctgcacg	2100
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acacgatccc	aacacacgca	gccaaactca	gaaccagca	ctgctccaag	cacctccttc	2220
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&lt;210&gt; 734

&lt;211&gt; 2300

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 734

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cattgggttc	gccaaagctc	tggtaaaagt	tggagtgagg	tttctgggat	ctggccttct	240
ggtggccata	ctcttttatgc	tgactccgtt	aaaggctcgt	tcactatctc	tagagacaac	300
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aagaaccagg	tcagcctgac	ctgcctgggtc	aaaggcttct	atccaagcga	catcgccgtg	1320
gagtggggaga	gcaatgggca	gccggagaac	aactacaaga	ccacgcctcc	cgtgctggac	1380
tccgacggct	ccttcttctc	ctacagcaag	ctcaccgtgg	acaagagcag	gtggcagcag	1440
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<210> 735

<211> 693

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 735

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Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser
 1             5             10             15
Val Ile Met Ser Arg Gly Glu Val Gln Leu Leu Glu Ser Gly Gly Gly
      20             25             30
Leu Val Gln Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly
      35             40             45
Phe Thr Phe Ser Trp Tyr Leu Met His Trp Val Arg Gln Ala Pro Gly
      50             55             60
Lys Gly Leu Glu Trp Val Ser Gly Ile Trp Pro Ser Gly Gly His Thr
65             70             75             80
Leu Tyr Ala Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn
      85             90             95
Ser Lys Asn Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp
      100            105            110
Thr Ala Val Tyr Tyr Cys Ala Arg Glu Pro Leu Leu Trp Phe Gly Glu
      115            120            125
Leu Ser Tyr Asn Trp Phe Asp Pro Trp Gly Gln Gly Thr Leu Val Thr
      130            135            140
Val Ser Ser Gly Gly Gly Gly Ser Gly Gly Gly Gly Ser Gly Gly Gly
145            150            155            160
Gly Ser Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Val Ser Ala
      165            170            175
Ser Val Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile
      180            185            190
Ser Ser Trp Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys
      195            200            205
Leu Leu Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg
      210            215            220
Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser
225            230            235            240
Leu Gln Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Ser
      245            250            255
Phe Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys Thr Gly
      260            265            270
Glu Pro Lys Ser Ser Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala
      275            280            285
Pro Glu Ala Ala Gly Ala Pro Ser Val Phe Leu Phe Pro Pro Lys Pro
      290            295            300

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Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val
305              310              315              320
Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val
              325              330              335
Asp Gly Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln
              340              345              350
Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln
              355              360              365
Asp Trp Leu Asn Gly Lys Ala Tyr Ala Cys Ala Val Ser Asn Lys Ala
              370              375              380
Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro
385              390              395              400
Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu Leu Thr
              405              410              415
Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser
              420              425              430
Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr
              435              440              445
Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr
              450              455              460
Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe
465              470              475              480
Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys
              485              490              495
Ser Leu Ser Leu Ser Pro Gly Gln Arg His Asn Asn Ser Ser Leu Asn
              500              505              510
Thr Arg Thr Gln Lys Ala Arg His Ser Pro Asn Ser Thr Ile Pro Pro
              515              520              525
His Val Gln Lys Ser Asp Val Glu Met Glu Ala Gln Lys Asp Glu Ile
              530              535              540
Ile Cys Pro Ser Cys Asn Arg Thr Ala His Pro Leu Arg His Ile Asn
545              550              555              560
Asn Asp Met Ile Val Thr Asp Asn Asn Gly Ala Val Lys Phe Pro Gln
              565              570              575
Leu Cys Lys Phe Cys Asp Val Arg Phe Ser Thr Cys Asp Asn Gln Lys
              580              585              590
Ser Cys Met Ser Asn Cys Ser Ile Thr Ser Ile Cys Glu Lys Pro Gln
              595              600              605
Glu Val Cys Val Ala Val Trp Arg Lys Asn Asp Glu Asn Ile Thr Leu
              610              615              620
Glu Thr Val Cys His Asp Pro Lys Leu Pro Tyr His Asp Phe Ile Leu
625              630              635              640
Glu Asp Ala Ala Ser Pro Lys Cys Ile Met Lys Glu Lys Lys Lys Pro
              645              650              655
Gly Glu Thr Phe Phe Met Cys Ser Cys Ser Ser Asp Glu Cys Asn Asp
              660              665              670
Asn Ile Ile Phe Ser Glu Glu Tyr Asn Thr Ser Asn Pro Asp Leu Leu
              675              680              685
Leu Val Ile Phe Gln
              690

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&lt;210&gt; 736

&lt;211&gt; 668

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

<223> Made in a lab

<400> 736

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser
1				5					10					15	
Val	Ile	Met	Ser	Arg	Gly	Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly
			20					25					30		
Leu	Val	Gln	Pro	Gly	Gly	Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly
		35					40					45			
Phe	Thr	Phe	Ser	Trp	Tyr	Leu	Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly
	50					55					60				
Lys	Gly	Leu	Glu	Trp	Val	Ser	Gly	Ile	Trp	Pro	Ser	Gly	Gly	His	Thr
65					70					75					80
Leu	Tyr	Ala	Asp	Ser	Val	Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn
				85					90					95	
Ser	Lys	Asn	Thr	Leu	Tyr	Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp
			100					105					110		
Thr	Ala	Val	Tyr	Tyr	Cys	Ala	Arg	Glu	Pro	Leu	Leu	Trp	Phe	Gly	Glu
		115					120					125			
Leu	Ser	Tyr	Asn	Trp	Phe	Asp	Pro	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr
		130				135					140				
Val	Ser	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly
145				150					155						160
Gly	Ser	Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala
			165					170					175		
Ser	Val	Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Ile
		180						185					190		
Ser	Ser	Trp	Leu	Ala	Trp	Tyr	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	
		195				200					205				
Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg
		210				215					220				
Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser
225				230					235						240
Leu	Gln	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ala	Asn	Ser
			245					250						255	
Phe	Pro	Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys	Thr	Gly
		260					265						270		
Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala
		275				280						285			
Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro
		290				295					300				
Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val
305				310						315					320
Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val
			325					330					335		
Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln
		340					345						350		
Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln
		355				360						365			
Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala
		370				375					380				
Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro
385				390						395					400
Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr
			405				410						415		
Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser

			420					425					430				
Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr		
		435						440					445				
Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr		
		450						455					460				
Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe		
465					470					475					480		
Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys		
				485						490					495		
Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	Ser	Leu	Asn		
			500					505						510			
Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Thr	Ile	Pro	Pro		
		515					520						525				
His	Val	Gln	Lys	Ser	Val	Asn	Asn	Asp	Met	Ile	Val	Thr	Asp	Asn	Asn		
		530				535					540						
Gly	Ala	Val	Lys	Phe	Pro	Gln	Leu	Cys	Lys	Phe	Cys	Asp	Val	Arg	Phe		
545					550					555					560		
Ser	Thr	Cys	Asp	Asn	Gln	Lys	Ser	Cys	Met	Ser	Asn	Cys	Ser	Ile	Thr		
				565					570					575			
Ser	Ile	Cys	Glu	Lys	Pro	Gln	Glu	Val	Cys	Val	Ala	Val	Trp	Arg	Lys		
			580					585					590				
Asn	Asp	Glu	Asn	Ile	Thr	Leu	Glu	Thr	Val	Cys	His	Asp	Pro	Lys	Leu		
		595					600						605				
Pro	Tyr	His	Asp	Phe	Ile	Leu	Glu	Asp	Ala	Ala	Ser	Pro	Lys	Cys	Ile		
	610					615					620						
Met	Lys	Glu	Lys	Lys	Lys	Pro	Gly	Glu	Thr	Phe	Phe	Met	Cys	Ser	Cys		
625					630					635					640		
Ser	Ser	Asp	Glu	Cys	Asn	Asp	Asn	Ile	Ile	Phe	Ser	Glu	Glu	Tyr	Asn		
				645					650					655			
Thr	Ser	Asn	Pro	Asp	Leu	Leu	Leu	Val	Ile	Phe	Gln						
			660					665									

&lt;210&gt; 737

&lt;211&gt; 693

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 737

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

		115					120					125				
Pro	Lys	Leu	Pro	Tyr	His	Asp	Phe	Ile	Leu	Glu	Asp	Ala	Ala	Ser	Pro	
	130					135					140					
Lys	Cys	Ile	Met	Lys	Glu	Lys	Lys	Lys	Pro	Gly	Glu	Thr	Phe	Phe	Met	
145					150					155					160	
Cys	Ser	Cys	Ser	Ser	Asp	Glu	Cys	Asn	Asp	Asn	Ile	Ile	Phe	Ser	Glu	
				165					170					175		
Glu	Tyr	Asn	Thr	Ser	Asn	Pro	Asp	Leu	Leu	Leu	Val	Ile	Phe	Gln	Thr	
			180					185					190			
Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	
		195					200					205				
Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	
		210				215					220					
Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	
225					230					235					240	
Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	
				245					250					255		
Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	
			260					265					270			
Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	
		275					280					285				
Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	
		290				295					300					
Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	
305					310					315					320	
Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	
				325					330					335		
Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	
			340					345					350			
Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	
		355					360					365				
Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	
		370				375					380					
Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	
385					390					395					400	
Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	
				405					410					415		
Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln	Arg	His	Asn	Asn	Ser	Ser	Leu	
			420					425				430				
Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His	Ser	Pro	Asn	Ser	Glu	Val	Gln	
		435					440					445				
Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly	Ser	Leu	Arg	
		450				455					460					
Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Trp	Tyr	Leu	Met	His	
465					470					475						

Gly Gly Gly Ser Gly Gly Gly Gly Ser Gln Asp Ile Gln Met Thr Gln  
                   580                  585                  590  
 Ser Pro Ser Ser Val Ser Ala Ser Val Gly Asp Arg Val Thr Ile Thr  
                   595                  600                  605  
 Cys Arg Ala Ser Gln Gly Ile Ser Ser Trp Leu Ala Trp Tyr Gln Gln  
           610                  615                  620  
 Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile Tyr Ala Ala Ser Ser Leu  
 625                  630                  635                  640  
 Gln Ser Gly Val Pro Ser Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp  
                   645                  650                  655  
 Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro Glu Asp Phe Ala Thr Tyr  
                   660                  665                  670  
 Tyr Cys Gln Gln Ala Asn Ser Phe Pro Leu Thr Phe Gly Gly Gly Thr  
           675                  680                  685  
 Lys Val Glu Ile Lys  
           690

<210> 738

<211> 668

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 738

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

Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val
				245					250					255	
Ser	Val	Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr
			260					265					270		
Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu	Lys	Thr
		275					280					285			
Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr	Thr	Leu
	290					295					300				
Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr	Lys	Asn	Gln	Val	Ser	Leu	Thr	Cys
305					310					315					320
Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp	Glu	Ser
				325					330					335	
Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val	Leu	Asp
			340					345					350		
Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp	Lys	Ser
		355					360					365			
Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His	Glu	Ala
	370					375					380				
Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro	Gly	Gln
385					390					395					400
Arg	His	Asn	Asn	Ser	Ser	Leu	Asn	Thr	Arg	Thr	Gln	Lys	Ala	Arg	His
				405					410					415	
Ser	Pro	Asn	Ser	Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val
			420					425					430		
Gln	Pro	Gly	Gly	Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr
		435					440					445			
Phe	Ser	Trp	Tyr	Leu	Met	His	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly
	450					455				460					
Leu	Glu	Trp	Val	Ser	Gly	Ile	Trp	Pro	Ser	Gly	Gly	His	Thr	Leu	Tyr
465					470					475					480
Ala	Asp	Ser	Val	Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys
				485					490					495	
Asn	Thr	Leu	Tyr	Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala
			500					505					510		
Val	Tyr	Tyr	Cys	Ala	Arg	Glu	Pro	Leu	Leu	Trp	Phe	Gly	Glu	Leu	Ser
		515					520					525			
Tyr	Asn	Trp	Phe	Asp	Pro	Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser
	530					535					540				
Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser	Gly	Gly	Gly	Gly	Ser
545					550					555					560
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Val	Ser	Ala	Ser	Val
				565					570					575	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Ile	Ser	Ser
			580					585					590		
Trp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		595					600					605			
Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser
	610					615					620				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln
625					630					635					640
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ala	Asn	Ser	Phe	Pro
				645					650					655	
Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			660					665							

&lt;210&gt; 739

&lt;211&gt; 2102



<212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 739  
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 ggtggttctt tacgtctttc ctgcgtgct tccggattca ctttctcttg gtaccttatg 180  
 cattgggttc gccaaagctcc tggtaaaggt ttggagtggg tttctggtat ctggccttct 240  
 ggtggccata ctctttatgc tgactccgtt aaaggctcgt tcactatctc tagagacaac 300  
 tctaagaata ctctctactt gcagatgaac agcttaaggg ctgaggacac ggccgtgtat 360  
 tactgtgcga gagagccgtt actatggttc ggggagttat cgtacaactg gttcgacccc 420  
 tggggccagg gcaccctggt caccgtctca agcgggtggc gcggttcggg ggggtggcga 480  
 agtggaggtg gagggagtca agacatccag atgaccagct ctccatcttc cgtgtctgca 540  
 tctgtaggag acagagtcac catcacttgt cgggcgagtc aggggtattag cagctgggta 600  
 gcctggtatc agcagaaacc agggaaagcc cctaagctcc tgatctatgc tgcattcagt 660  
 ttgcaaagtg ggggtcccatc aagggttcagc ggcagtggtat ctgggacaga tttcactctc 720  
 accatcagca gcctgcagcc tgaagatttt gcaacttact attgtcaaca ggctaacagt 780  
 ttccctctca ctttcggcgg agggaccaag gtggagatca aaaccggtga gcccaaactc 840  
 tctgacaaaa ctcacacatg cccaccgtgc ccagcacctg aagccgcggg tgcaccgtca 900  
 gtcttctctt tcccccaaaa acccaaggac accctcatga tctcccgga ccctgagggtc 960  
 acatgcgtgg tgggtggacgt gagccacgaa gaccctgagg tcaagttcaa ctggtacgtg 1020  
 gacggcgtgg aggtgcataa tgccaagaca aagccgcggg aggagcagta caacagcacg 1080  
 taccgtgtgg tcagcgtcct caccgtcctg caccaggact ggctgaatgg caaggcgtac 1140  
 gcgtgcgcgg tctccaacaa agccctccca gccccatcg agaaaaccat ctccaaagcc 1200  
 aaagggcagc cccgagaacc acaggtgtac accctgcccc catcccggga tgagctgacc 1260  
 aagaaccagg tcagcctgac ctgcctggtc aaaggcttct atccaagcga catcgccgtg 1320  
 gagtgggaga gcaatgggca gccggagaa aactacaaga ccacgcctcc cgtgctggac 1380  
 tccgacggct ccttcttctt ctacagcaag ctaccgtgg acaagagcag gtggcagcag 1440  
 gggaaactgt tctcatgctc cgtgatgcat gaggtctctg acaaccacta cacgcagaag 1500  
 agcctctccc tgtctccggg tcagaggcac aacaattctt cctgaatac aagaactcag 1560  
 aaagcacgtc atttctccga ttctacgac ccaccgcagc ttcagaagtc ggatgtggaa 1620  
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 agacatatta ataacgacat gatagtcact gacaacaacg gtgcagtcaa gtttccacaa 1740  
 ctgtgtaaat tttgtgatgt gagattttcc acctgtgaca accagaaatc ctgcatgagc 1800  
 aactgcagca tcacctccat ctgtgagaag ccacaggaag tctgtgtggc tgtatggaga 1860  
 aagaatgacg agaacataac actagagaca gtttgccatg accccaagct cccctaccat 1920  
 gactttatct tgggaagatgc tgcttctcca aagtgcatta tgaaggaaaa aaaaaagcct 1980  
 ggtgagactt tcttcatgtg ttctgtagc tctgatgagt gcaatgacaa catcatcttc 2040  
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 gc 2102

<210> 740  
 <211> 2027  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 740  
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 gtcataatgt cgcgaggaga agttcaattg ttagagtctg gtggcggtct tgttcagcct 120

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ggtgggttctt tacgtctttc ctgcgctgct tccggattca ctttctcttg gtaccttatg 180
cattggggttc gccaaagctcc tggtaaagggt ttggagtgagg tttctggtat ctggccttct 240
ggtggccata ctctttatgc tgactccgtt aaaggctcgt tcaactatctc tagagacaac 300
tctaagaata ctctctactt gcagatgaac agcttaaggg ctgaggacac ggccgtgtat 360
tactgtgcga gagagccggt actatgggtc ggggagttat cgtacaactg gttcgacccc 420
tgggggccagg gcaccctggg caccgtctca agcgggtggc gcgggttcggg ggggtggcgga 480
agtggagggtg gagggagtc aagacatccag atgacccagt ctccatcttc cgtgtctgca 540
tctgtaggag acagagtcac catcacttgt cgggcgagtc aggggtattag cagctgggta 600
gcctgggtatc agcagaaacc agggaaagcc cctaagctcc tgatctatgc tgcattccagt 660
ttgcaaagtg ggggtcccatc aagggttcagc ggcagtggtat ctgggacaga tttcactctc 720
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tctgacaaaa cttcacatg cccaccgtgc ccagacctga aagccgcggg tgcaccgtca 900
gtcttcctct tcccccaaa acccaaggac accctcatga tctcccggac ccctgagggtc 960
acatgcgtgg tgggtggacgt gagccacgaa gaccctgagg tcaagttcaa ctggtacgtg 1020
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gagtgggaga gcaatgggca gccggagaac aactacaaga ccacgcctcc cgtgctggac 1380
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aaagcacgtc atttctccga ttctacgac ccaccgcacg ttcagaagtc ggttaataac 1620
gacatgatag tcaactgaca caacgggtgca gtcaagtttc cacaactgtg taaattttgt 1680
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ataacactag agacagtttg ccatgacccc aagctcccc accatgactt tattctggaa 1860
gatgctgctt ctccaaagtg cattatgaag gaaaaaaaaa agcctgggtg gactttcttc 1920
atgtgttctt gtagctctga tgagtgaat gacaacatca tcttctcaga agaataaac 1980
accagcaatc ctgacttggt gtagtcata tttcaatgag cggccgc 2027

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<210> 741

<211> 2102

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 741

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gccagaaaag atgaaatcat ctgcccagc tgtaatagga ctgcccattc actgagacat 180
attaataacg acatgatagt cactgacaac aacgggtgcag tcaagtttcc acaactgtgt 240
aaatttttgt atgtgagatt ttccacctgt gacaaccaga aatcctgcat gagcaactgc 300
agcatcacct ccatctgtga gaagccacag gaagtctgtg tggctgtatg gagaaagaat 360
gacgagaaca taacactaga gacagtttgc catgaccca agctccccta ccatgacttt 420
attctggaag atgtgtcttc tccaaagtgc attatgaagg aaaaaaaaaa gcctgggtgag 480
actttcttca tgtgttctct tagctctgat gagtgcattg acaacatcat cttctcagaa 540
gaatataaca ccagcaatcc tgacttggtg ctagtcatat ttcaaaccgg tgagccaaa 600
tcttctgaca aaactcacac atgcccaccg tgcccagcac ctgaagccgc ggggtgcaccg 660
tcagtcttcc tcttcccccc aaaacccaag gacaccctca tgatctcccg gaccctgag 720
gtcacatgcg tgggtgggga cgtgagccac gaagaccctg aggtcaagtt caactgggtac 780
gtggacggcg tggagggtgca taatgccaa acaaagccgc gggaggagca gtacaacagc 840

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gccaaagggc	agccccgaga	accacaggtg	tacaccctgc	ccccatccc	ggatgagctg	1020
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gc						2102

&lt;210&gt; 742

&lt;211&gt; 2027

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 742

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atagtcactg	acaacaacgg	tgcagtcaag	tttccacaac	tgtgtaaatt	ttgtgatgtg	180
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cgagaaccac	aggtgtacac	cctgccccca	tcccgggatg	agctgaccaa	gaaccaggctc	960
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gcgagagagc cgttactatg gttcggggag ttatcgtaga actgggtcga cccctggggc 1620
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tatcagcaga aaccagggaa agcccctaag ctcctgatct atgctgcatc cagtttgcaa 1860
agtgggggtc catcaagggt cagcggcagt ggatctggga cagatttcac tctcaccatc 1920
agcagcctgc agcctgaaga ttttgcaact tactattgtc aacaggctaa cagtttccct 1980
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<210> 743

<211> 191

<212> PRT

<213> Human

<400> 743

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

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<210> 744

<211> 166

<212> PRT

<213> Human

<400> 744

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

```

65					70					75				80	
Gln	Glu	Val	Cys	Val	Ala	Val	Trp	Arg	Lys	Asn	Asp	Glu	Asn	Ile	Thr
				85					90					95	
Leu	Glu	Thr	Val	Cys	His	Asp	Pro	Lys	Leu	Pro	Tyr	His	Asp	Phe	Ile
			100					105					110		
Leu	Glu	Asp	Ala	Ala	Ser	Pro	Lys	Cys	Ile	Met	Lys	Glu	Lys	Lys	Lys
		115					120					125			
Pro	Gly	Glu	Thr	Phe	Phe	Met	Cys	Ser	Cys	Ser	Ser	Asp	Glu	Cys	Asn
	130					135					140				
Asp	Asn	Ile	Ile	Phe	Ser	Glu	Glu	Tyr	Asn	Thr	Ser	Asn	Pro	Asp	Leu
145					150					155					160
Leu	Leu	Val	Ile	Phe	Gln										
				165											

<210> 745  
 <211> 233  
 <212> PRT  
 <213> Human

<400> 745															
Met	Leu	Pro	Cys	Leu	Val	Val	Leu	Leu	Ala	Ala	Leu	Leu	Ser	Leu	Arg
1				5					10					15	
Leu	Gly	Ser	Asp	Ala	His	Gly	Thr	Glu	Leu	Pro	Ser	Pro	Pro	Ser	Val
			20					25					30		
Trp	Phe	Glu	Ala	Glu	Phe	Phe	His	His	Ile	Leu	His	Trp	Thr	Pro	Ile
	35						40					45			
Pro	Asn	Gln	Ser	Glu	Ser	Thr	Cys	Tyr	Glu	Val	Ala	Leu	Leu	Arg	Tyr
	50					55					60				
Gly	Ile	Glu	Ser	Trp	Asn	Ser	Ile	Ser	Asn	Cys	Ser	Gln	Thr	Leu	Ser
65					70					75					80
Tyr	Asp	Leu	Thr	Ala	Val	Thr	Leu	Asp	Leu	Tyr	His	Ser	Asn	Gly	Tyr
				85					90					95	
Arg	Ala	Arg	Val	Arg	Ala	Val	Asp	Gly	Ser	Arg	His	Ser	Asn	Trp	Thr
			100					105					110		
Val	Thr	Asn	Thr	Arg	Phe	Ser	Val	Asp	Glu	Val	Thr	Leu	Thr	Val	Gly
	115						120					125			
Ser	Val	Asn	Leu	Glu	Ile	His	Asn	Gly	Phe	Ile	Leu	Gly	Lys	Ile	Gln
	130					135					140				
Leu	Pro	Arg	Pro	Lys	Met	Ala	Pro	Ala	Asn	Asp	Thr	Tyr	Glu	Ser	Ile
145					150					155					160
Phe	Ser	His	Phe	Arg	Glu	Tyr	Glu	Ile	Ala	Ile	Arg	Lys	Val	Pro	Gly
				165					170					175	
Asn	Phe	Thr	Phe	Thr	His	Lys	Lys	Val	Lys	His	Glu	Asn	Phe	Ser	Leu
		180						185					190		
Leu	Thr	Ser	Gly	Glu	Val	Gly	Glu	Phe	Cys	Val	Gln	Val	Lys	Pro	Ser
	195						200					205			
Val	Ala	Ser	Arg	Ser	Asn	Lys	Gly	Met	Trp	Ser	Lys	Glu	Glu	Cys	Ile
	210					215					220				
Ser	Leu	Thr	Arg	Gln	Tyr	Phe	Thr	Val							
225					230										

<210> 746  
 <211> 126  
 <212> PRT  
 <213> Human

<400> 746

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

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<210> 747
<211> 1356
<212> PRT
<213> Human

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<400> 747
Met Gln Ser Lys Val Leu Leu Ala Val Ala Leu Trp Leu Cys Val Glu
 1      5      10      15
Thr Arg Ala Ala Ser Val Gly Leu Pro Ser Val Ser Leu Asp Leu Pro
      20      25      30
Arg Leu Ser Ile Gln Lys Asp Ile Leu Thr Ile Lys Ala Asn Thr Thr
      35      40      45
Leu Gln Ile Thr Cys Arg Gly Gln Arg Asp Leu Asp Trp Leu Trp Pro
      50      55      60
Asn Asn Gln Ser Gly Ser Glu Gln Arg Val Glu Val Thr Glu Cys Ser
65      70      75      80
Asp Gly Leu Phe Cys Lys Thr Leu Thr Ile Pro Lys Val Ile Gly Asn
      85      90      95
Asp Thr Gly Ala Tyr Lys Cys Phe Tyr Arg Glu Thr Asp Leu Ala Ser
      100      105      110
Val Ile Tyr Val Tyr Val Gln Asp Tyr Arg Ser Pro Phe Ile Ala Ser
      115      120      125
Val Ser Asp Gln His Gly Val Val Tyr Ile Thr Glu Asn Lys Asn Lys
      130      135      140
Thr Val Val Ile Pro Cys Leu Gly Ser Ile Ser Asn Leu Asn Val Ser
145      150      155      160
Leu Cys Ala Arg Tyr Pro Glu Lys Arg Phe Val Pro Asp Gly Asn Arg
      165      170      175
Ile Ser Trp Asp Ser Lys Lys Gly Phe Thr Ile Pro Ser Tyr Met Ile
      180      185      190
Ser Tyr Ala Gly Met Val Phe Cys Glu Ala Lys Ile Asn Asp Glu Ser
      195      200      205
Tyr Gln Ser Ile Met Tyr Ile Val Val Val Val Gly Tyr Arg Ile Tyr
      210      215      220
Asp Val Val Leu Ser Pro Ser His Gly Ile Glu Leu Ser Val Gly Glu
225      230      235      240
Lys Leu Val Leu Asn Cys Thr Ala Arg Thr Glu Leu Asn Val Gly Ile
      245      250      255
Asp Phe Asn Trp Glu Tyr Pro Ser Ser Lys His Gln His Lys Lys Leu
      260      265      270

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Val	Asn	Arg	Asp	Leu	Lys	Thr	Gln	Ser	Gly	Ser	Glu	Met	Lys	Lys	Phe
		275					280					285			
Leu	Ser	Thr	Leu	Thr	Ile	Asp	Gly	Val	Thr	Arg	Ser	Asp	Gln	Gly	Leu
	290					295					300				
Tyr	Thr	Cys	Ala	Ala	Ser	Ser	Gly	Leu	Met	Thr	Lys	Lys	Asn	Ser	Thr
305					310					315					320
Phe	Val	Arg	Val	His	Glu	Lys	Pro	Phe	Val	Ala	Phe	Gly	Ser	Gly	Met
				325					330					335	
Glu	Ser	Leu	Val	Glu	Ala	Thr	Val	Gly	Glu	Arg	Val	Arg	Ile	Pro	Ala
			340					345					350		
Lys	Tyr	Leu	Gly	Tyr	Pro	Pro	Pro	Glu	Ile	Lys	Trp	Tyr	Lys	Asn	Gly
	355						360					365			
Ile	Pro	Leu	Glu	Ser	Asn	His	Thr	Ile	Lys	Ala	Gly	His	Val	Leu	Thr
	370					375					380				
Ile	Met	Glu	Val	Ser	Glu	Arg	Asp	Thr	Gly	Asn	Tyr	Thr	Val	Ile	Leu
385					390					395					400
Thr	Asn	Pro	Ile	Ser	Lys	Glu	Lys	Gln	Ser	His	Val	Val	Ser	Leu	Val
				405					410					415	
Val	Tyr	Val	Pro	Pro	Gln	Ile	Gly	Glu	Lys	Ser	Leu	Ile	Ser	Pro	Val
			420					425					430		
Asp	Ser	Tyr	Gln	Tyr	Gly	Thr	Thr	Gln	Thr	Leu	Thr	Cys	Thr	Val	Tyr
		435					440					445			
Ala	Ile	Pro	Pro	Pro	His	His	Ile	His	Trp	Tyr	Trp	Gln	Leu	Glu	Glu
	450					455					460				
Glu	Cys	Ala	Asn	Glu	Pro	Ser	Gln	Ala	Val	Ser	Val	Thr	Asn	Pro	Tyr
465					470					475					480
Pro	Cys	Glu	Glu	Trp	Arg	Ser	Val	Glu	Asp	Phe	Gln	Gly	Gly	Asn	Lys
				485					490					495	
Ile	Glu	Val	Asn	Lys	Asn	Gln	Phe	Ala	Leu	Ile	Glu	Gly	Lys	Asn	Lys
			500					505					510		
Thr	Val	Ser	Thr	Leu	Val	Ile	Gln	Ala	Ala	Asn	Val	Ser	Ala	Leu	Tyr
		515					520					525			
Lys	Cys	Glu	Ala	Val	Asn	Lys	Val	Gly	Arg	Gly	Glu	Arg	Val	Ile	Ser
	530					535					540				
Phe	His	Val	Thr	Arg	Gly	Pro	Glu	Ile	Thr	Leu	Gln	Pro	Asp	Met	Gln
545					550					555					560
Pro	Thr	Glu	Gln	Glu	Ser	Val	Ser	Leu	Trp	Cys	Thr	Ala	Asp	Arg	Ser
				565					570					575	
Thr	Phe	Glu	Asn	Leu	Thr	Trp	Tyr	Lys	Leu	Gly	Pro	Gln	Pro	Leu	Pro
			580					585					590		
Ile	His	Val	Gly	Glu	Leu	Pro	Thr	Pro	Val	Cys	Lys	Asn	Leu	Asp	Thr
		595					600					605			
Leu	Trp	Lys	Leu	Asn	Ala	Thr	Met	Phe	Ser	Asn	Ser	Thr	Asn	Asp	Ile
	610					615					620				
Leu	Ile	Met	Glu	Leu	Lys	Asn	Ala	Ser	Leu	Gln	Asp	Gln	Gly	Asp	Tyr
625					630					635					640
Val	Cys	Leu	Ala	Gln	Asp	Arg	Lys	Thr	Lys	Lys	Arg	His	Cys	Val	Val
				645					650					655	
Arg	Gln	Leu	Thr	Val	Leu	Glu	Arg	Val	Ala	Pro	Thr	Ile	Thr	Gly	Asn
			660					665					670		
Leu	Glu	Asn	Gln	Thr	Thr	Ser	Ile	Gly	Glu	Ser	Ile	Glu	Val	Ser	Cys
		675					680					685			
Thr	Ala	Ser	Gly	Asn	Pro	Pro	Pro	Gln	Ile	Met	Trp	Phe	Lys	Asp	Asn
	690					695					700				
Glu	Thr	Leu	Val	Glu	Asp	Ser	Gly	Ile	Val	Leu	Lys	Asp	Gly	Asn	Arg
705					710					715					720
Asn	Leu	Thr	Ile	Arg	Arg	Val	Arg	Lys	Glu	Asp	Glu	Gly	Leu	Tyr	Thr

				725					730					735			
Cys	Gln	Ala	Cys	Ser	Val	Leu	Gly	Cys	Ala	Lys	Val	Glu	Ala	Phe	Phe		
			740					745					750				
Ile	Ile	Glu	Gly	Ala	Gln	Glu	Lys	Thr	Asn	Leu	Glu	Ile	Ile	Ile	Leu		
		755					760					765					
Val	Gly	Thr	Ala	Val	Ile	Ala	Met	Phe	Phe	Trp	Leu	Leu	Leu	Val	Ile		
	770					775					780						
Ile	Leu	Arg	Thr	Val	Lys	Arg	Ala	Asn	Gly	Gly	Glu	Leu	Lys	Thr	Gly		
785					790					795					800		
Tyr	Leu	Ser	Ile	Val	Met	Asp	Pro	Asp	Glu	Leu	Pro	Leu	Asp	Glu	His		
				805					810					815			
Cys	Glu	Arg	Leu	Pro	Tyr	Asp	Ala	Ser	Lys	Trp	Glu	Phe	Pro	Arg	Asp		
			820					825					830				
Arg	Leu	Lys	Leu	Gly	Lys	Pro	Leu	Gly	Arg	Gly	Ala	Phe	Gly	Gln	Val		
		835					840					845					
Ile	Glu	Ala	Asp	Ala	Phe	Gly	Ile	Asp	Lys	Thr	Ala	Thr	Cys	Arg	Thr		
	850					855					860						
Val	Ala	Val	Lys	Met	Leu	Lys	Glu	Gly	Ala	Thr	His	Ser	Glu	His	Arg		
865					870					875					880		
Ala	Leu	Met	Ser	Glu	Leu	Lys	Ile	Leu	Ile	His	Ile	Gly	His	His	Leu		
				885				890						895			
Asn	Val	Val	Asn	Leu	Leu	Gly	Ala	Cys	Thr	Lys	Pro	Gly	Gly	Pro	Leu		
			900					905					910				
Met	Val	Ile	Val	Glu	Phe	Cys	Lys	Phe	Gly	Asn	Leu	Ser	Thr	Tyr	Leu		
	915						920					925					
Arg	Ser	Lys	Arg	Asn	Glu	Phe	Val	Pro	Tyr	Lys	Thr	Lys	Gly	Ala	Arg		
	930					935					940						
Phe	Arg	Gln	Gly	Lys	Asp	Tyr	Val	Gly	Ala	Ile	Pro	Val	Asp	Leu	Lys		
945					950					955					960		
Arg	Arg	Leu	Asp	Ser	Ile	Thr	Ser	Ser	Gln	Ser	Ser	Ala	Ser	Ser	Gly		
				965					970					975			
Phe	Val	Glu	Glu	Lys	Ser	Leu	Ser	Asp	Val	Glu	Glu	Glu	Glu	Ala	Pro		
			980					985					990				
Glu	Asp	Leu	Tyr	Lys	Asp	Phe	Leu	Thr	Leu	Glu	His	Leu	Ile	Cys	Tyr		
	995						1000					1005					
Ser	Phe	Gln	Val	Ala	Lys	Gly	Met	Glu	Phe	Leu	Ala	Ser	Arg	Lys	Cys		
	1010					1015					1020						
Ile	His	Arg	Asp	Leu	Ala	Ala	Arg	Asn	Ile	Leu	Leu	Ser	Glu	Lys	Asn		
1025					1030					1035					1040		
Val	Val	Lys	Ile	Cys	Asp	Phe	Gly	Leu	Ala	Arg	Asp	Ile	Tyr	Lys	Asp		
				1045					1050					1055			
Pro	Asp	Tyr	Val	Arg	Lys	Gly	Asp	Ala	Arg	Leu	Pro	Leu	Lys	Trp	Met		
			1060					1065					1070				
Ala	Pro	Glu	Thr	Ile	Phe	Asp	Arg	Val	Tyr	Thr	Ile	Gln	Ser	Asp	Val		
	1075						1080					1085					
Trp	Ser	Phe	Gly	Val	Leu	Leu	Trp	Glu	Ile	Phe	Ser	Leu	Gly	Ala	Ser		
	1090					1095					1100						
Pro	Tyr	Pro	Gly	Val	Lys	Ile	Asp	Glu	Glu	Phe	Cys	Arg	Arg	Leu	Lys		
1105					1110					1115					1120		
Glu	Gly	Thr	Arg	Met	Arg	Ala	Pro	Asp	Tyr	Thr	Thr	Pro	Glu	Met	Tyr		
				1125					1130					1135			
Gln	Thr	Met	Leu	Asp	Cys	Trp	His	Gly	Glu	Pro	Ser	Gln	Arg	Pro	Thr		
			1140					1145					1150				
Phe	Ser	Glu	Leu	Val	Glu	His	Leu	Gly	Asn	Leu	Leu	Gln	Ala	Asn	Ala		
	1155						1160					1165					
Gln	Gln	Asp	Gly	Lys	Asp	Tyr	Ile	Val	Leu	Pro	Ile	Ser	Glu	Thr	Leu		
	1170					1175					1180						



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Ser Met Glu Glu Asp Ser Gly Leu Ser Leu Pro Thr Ser Pro Val Ser
1185                1190                1195                1200
Cys Met Glu Glu Glu Glu Val Cys Asp Pro Lys Phe His Tyr Asp Asn
                1205                1210                1215
Thr Ala Gly Ile Ser Gln Tyr Leu Gln Asn Ser Lys Arg Lys Ser Arg
                1220                1225                1230
Pro Val Ser Val Lys Thr Phe Glu Asp Ile Pro Leu Glu Glu Pro Glu
                1235                1240                1245
Val Lys Val Ile Pro Asp Asp Asn Gln Thr Asp Ser Gly Met Val Leu
                1250                1255                1260
Ala Ser Glu Glu Leu Lys Thr Leu Glu Asp Arg Thr Lys Leu Ser Pro
1265                1270                1275                1280
Ser Phe Gly Gly Met Val Pro Ser Lys Ser Arg Glu Ser Val Ala Ser
                1285                1290                1295
Glu Gly Ser Asn Gln Thr Ser Gly Tyr Gln Ser Gly Tyr His Ser Asp
                1300                1305                1310
Asp Thr Asp Thr Thr Val Tyr Ser Ser Glu Glu Ala Glu Leu Leu Lys
                1315                1320                1325
Leu Ile Glu Ile Gly Val Gln Thr Gly Ser Thr Ala Gln Ile Leu Gln
                1330                1335                1340
Pro Asp Ser Gly Thr Thr Leu Ser Ser Pro Pro Val
1345                1350                1355

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<210> 748
<211> 257
<212> PRT
<213> Human

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<400> 748
Met Ala Pro Val Ala Val Trp Ala Ala Leu Ala Val Gly Leu Glu Leu
 1                5                10                15
Trp Ala Ala Ala His Ala Leu Pro Ala Gln Val Ala Phe Thr Pro Tyr
                20                25                30
Ala Pro Glu Pro Gly Ser Thr Cys Arg Leu Arg Glu Tyr Tyr Asp Gln
                35                40                45
Thr Ala Gln Met Cys Cys Ser Lys Cys Ser Pro Gly Gln His Ala Lys
                50                55                60
Val Phe Cys Thr Lys Thr Ser Asp Thr Val Cys Asp Ser Cys Glu Asp
65                70                75                80
Ser Thr Tyr Thr Gln Leu Trp Asn Trp Val Pro Glu Cys Leu Ser Cys
                85                90                95
Gly Ser Arg Cys Ser Ser Asp Gln Val Glu Thr Gln Ala Cys Thr Arg
                100                105                110
Glu Gln Asn Arg Ile Cys Thr Cys Arg Pro Gly Trp Tyr Cys Ala Leu
                115                120                125
Ser Lys Gln Glu Gly Cys Arg Leu Cys Ala Pro Leu Arg Lys Cys Arg
                130                135                140
Pro Gly Phe Gly Val Ala Arg Pro Gly Thr Glu Thr Ser Asp Val Val
145                150                155                160
Cys Lys Pro Cys Ala Pro Gly Thr Phe Ser Asn Thr Thr Ser Ser Thr
                165                170                175
Asp Ile Cys Arg Pro His Gln Ile Cys Asn Val Val Ala Ile Pro Gly
                180                185                190
Asn Ala Ser Met Asp Ala Val Cys Thr Ser Thr Ser Pro Thr Arg Ser
                195                200                205
Met Ala Pro Gly Ala Val His Leu Pro Gln Pro Val Ser Thr Arg Ser

```

210		215		220
Gln His Thr Gln Pro Thr Pro Glu Pro Ser Thr Ala Pro Ser Thr Ser				
225		230		235
Phe Leu Leu Pro Met Gly Pro Ser Pro Pro Ala Glu Gly Ser Thr Gly				
	245		250	255

Asp

<210> 749  
 <211> 211  
 <212> PRT  
 <213> Human

<400> 749

Met Gly Leu Ser Thr Val Pro Asp Leu Leu Leu Pro Leu Val Leu Leu				
1	5	10	15	
Glu Leu Leu Val Gly Ile Tyr Pro Ser Gly Val Ile Gly Leu Val Pro				
	20	25	30	
His Leu Gly Asp Arg Glu Lys Arg Asp Ser Val Cys Pro Gln Gly Lys				
	35	40	45	
Tyr Ile His Pro Gln Asn Asn Ser Ile Cys Cys Thr Lys Cys His Lys				
	50	55	60	
Gly Thr Tyr Leu Tyr Asn Asp Cys Pro Gly Pro Gly Gln Asp Thr Asp				
65	70	75	80	
Cys Arg Glu Cys Glu Ser Gly Ser Phe Thr Ala Ser Glu Asn His Leu				
	85	90	95	
Arg His Cys Leu Ser Cys Ser Lys Cys Arg Lys Glu Met Gly Gln Val				
	100	105	110	
Glu Ile Ser Ser Cys Thr Val Asp Arg Asp Thr Val Cys Gly Cys Arg				
	115	120	125	
Lys Asn Gln Tyr Arg His Tyr Trp Ser Glu Asn Leu Phe Gln Cys Phe				
	130	135	140	
Asn Cys Ser Leu Cys Leu Asn Gly Thr Val His Leu Ser Cys Gln Glu				
145	150	155	160	
Lys Gln Asn Thr Val Cys Thr Cys His Ala Gly Phe Phe Leu Arg Glu				
	165	170	175	
Asn Glu Cys Val Ser Cys Ser Asn Cys Lys Lys Ser Leu Glu Cys Thr				
	180	185	190	
Lys Leu Cys Leu Pro Gln Ile Glu Asn Val Lys Gly Thr Glu Asp Ser				
	195	200	205	
Gly Thr Thr				
210				

<210> 750  
 <211> 950  
 <212> PRT  
 <213> Human

<400> 750

Met Lys Ala Pro Ala Val Leu Ala Pro Gly Ile Leu Val Leu Leu Phe				
1	5	10	15	
Thr Leu Val Gln Arg Ser Asn Gly Glu Cys Lys Glu Ala Leu Ala Lys				
	20	25	30	
Ser Glu Met Asn Val Asn Met Lys Tyr Gln Leu Pro Asn Phe Thr Ala				
	35	40	45	
Glu Thr Pro Ile Gln Asn Val Ile Leu His Glu His His Ile Phe Leu				
	50	55	60	

Gly	Ala	Thr	Asn	Tyr	Ile	Tyr	Val	Leu	Asn	Glu	Glu	Asp	Leu	Gln	Lys
65					70					75					80
Val	Ala	Glu	Tyr	Lys	Thr	Gly	Pro	Val	Leu	Glu	His	Pro	Asp	Cys	Phe
				85					90					95	
Pro	Cys	Gln	Asp	Cys	Ser	Ser	Lys	Ala	Asn	Leu	Ser	Gly	Gly	Val	Trp
			100					105					110		
Lys	Asp	Asn	Ile	Asn	Met	Ala	Leu	Val	Val	Asp	Thr	Tyr	Tyr	Asp	Asp
		115					120					125			
Gln	Leu	Ile	Ser	Cys	Gly	Ser	Val	Asn	Arg	Gly	Thr	Cys	Gln	Arg	His
	130						135				140				
Val	Phe	Pro	His	Asn	His	Thr	Ala	Asp	Ile	Gln	Ser	Glu	Val	His	Cys
145					150					155					160
Ile	Phe	Ser	Pro	Gln	Ile	Glu	Glu	Pro	Ser	Gln	Cys	Pro	Asp	Cys	Val
				165					170					175	
Val	Ser	Ala	Leu	Gly	Ala	Lys	Val	Leu	Ser	Ser	Val	Lys	Asp	Arg	Phe
			180					185					190		
Ile	Asn	Phe	Phe	Val	Gly	Asn	Thr	Ile	Asn	Ser	Ser	Tyr	Phe	Pro	Asp
	195						200					205			
His	Pro	Leu	His	Ser	Ile	Ser	Val	Arg	Arg	Leu	Lys	Glu	Thr	Lys	Asp
	210						215				220				
Gly	Phe	Met	Phe	Leu	Thr	Asp	Gln	Ser	Tyr	Ile	Asp	Val	Leu	Pro	Glu
225					230					235					240
Phe	Arg	Asp	Ser	Tyr	Pro	Ile	Lys	Tyr	Val	His	Ala	Phe	Glu	Ser	Asn
				245					250					255	
Asn	Phe	Ile	Tyr	Phe	Leu	Thr	Val	Gln	Arg	Glu	Thr	Leu	Asp	Ala	Gln
			260					265					270		
Thr	Phe	His	Thr	Arg	Ile	Ile	Arg	Phe	Cys	Ser	Ile	Asn	Ser	Gly	Leu
	275						280					285			
His	Ser	Tyr	Met	Glu	Met	Pro	Leu	Glu	Cys	Ile	Leu	Thr	Glu	Lys	Arg
	290					295					300				
Lys	Lys	Arg	Ser	Thr	Lys	Lys	Glu	Val	Phe	Asn	Ile	Leu	Gln	Ala	Ala
305					310					315					320
Tyr	Val	Ser	Lys	Pro	Gly	Ala	Gln	Leu	Ala	Arg	Gln	Ile	Gly	Ala	Ser
				325					330					335	
Leu	Asn	Asp	Asp	Ile	Leu	Phe	Gly	Val	Phe	Ala	Gln	Ser	Lys	Pro	Asp
			340					345					350		
Ser	Ala	Glu	Pro	Met	Asp	Arg	Ser	Ala	Met	Cys	Ala	Phe	Pro	Ile	Lys
	355						360					365			
Tyr	Val	Asn	Asp	Phe	Phe	Asn	Lys	Ile	Val	Asn	Lys	Asn	Asn	Val	Arg
	370					375					380				
Cys	Leu	Gln	His	Phe	Tyr	Gly	Pro	Asn	His	Glu	His	Cys	Phe	Asn	Arg
385					390					395					400
Thr	Leu	Leu	Arg	Asn	Ser	Ser	Gly	Cys	Glu	Ala	Arg	Arg	Asp	Glu	Tyr
				405					410					415	
Arg	Thr	Glu	Phe	Thr	Thr	Ala	Leu	Gln	Arg	Val	Asp	Leu	Phe	Met	Gly
			420					425					430		
Gln	Phe	Ser	Glu	Val	Leu	Leu	Thr	Ser	Ile	Ser	Thr	Phe	Ile	Lys	Gly
	435						440					445			
Asp	Leu	Thr	Ile	Ala	Asn	Leu	Gly	Thr	Ser	Glu	Gly	Arg	Phe	Met	Gln
	450					455					460				
Val	Val	Val	Ser	Arg	Ser	Gly	Pro	Ser	Thr	Pro	His	Val	Asn	Phe	Leu
465					470					475					480
Leu	Asp	Ser	His	Pro	Val	Ser	Pro	Glu	Val	Ile	Val	Glu	His	Thr	Leu
				485					490					495	
Asn	Gln	Asn	Gly	Tyr	Thr	Leu	Val	Ile	Thr	Gly	Lys	Lys	Ile	Thr	Lys
			500					505					510		
Ile	Pro	Leu	Asn	Gly	Leu	Gly	Cys	Arg	His	Phe	Gln	Ser	Cys	Ser	Gln

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 <213> Human

<400> 751

Met	Lys	Ala	Pro	Ala	Val	Leu	Ala	Pro	Gly	Ile	Leu	Val	Leu	Leu	Phe	1	5	10	15
Thr	Leu	Val	Gln	Arg	Ser	Asn	Gly	Glu	Cys	Lys	Glu	Ala	Leu	Ala	Lys	20	25	30	
Ser	Glu	Met	Asn	Val	Asn	Met	Lys	Tyr	Gln	Leu	Pro	Asn	Phe	Thr	Ala	35	40	45	
Glu	Thr	Pro	Ile	Gln	Asn	Val	Ile	Leu	His	Glu	His	His	Ile	Phe	Leu	50	55	60	
Gly	Ala	Thr	Asn	Tyr	Ile	Tyr	Val	Leu	Asn	Glu	Glu	Asp	Leu	Gln	Lys	65	70	75	80
Val	Ala	Glu	Tyr	Lys	Thr	Gly	Pro	Val	Leu	Glu	His	Pro	Asp	Cys	Phe	85	90	95	
Pro	Cys	Gln	Asp	Cys	Ser	Ser	Lys	Ala	Asn	Leu	Ser	Gly	Gly	Val	Trp	100	105	110	
Lys	Asp	Asn	Ile	Asn	Met	Ala	Leu	Val	Val	Asp	Thr	Tyr	Tyr	Asp	Asp	115	120	125	
Gln	Leu	Ile	Ser	Cys	Gly	Ser	Val	Asn	Arg	Gly	Thr	Cys	Gln	Arg	His	130	135	140	
Val	Phe	Pro	His	Asn	His	Thr	Ala	Asp	Ile	Gln	Ser	Glu	Val	His	Cys	145	150	155	160
Ile	Phe	Ser	Pro	Gln	Ile	Glu	Glu	Pro	Ser	Gln	Cys	Pro	Asp	Cys	Val	165	170	175	
Val	Ser	Ala	Leu	Gly	Ala	Lys	Val	Leu	Ser	Ser	Val	Lys	Asp	Arg	Phe	180	185	190	
Ile	Asn	Phe	Phe	Val	Gly	Asn	Thr	Ile	Asn	Ser	Ser	Tyr	Phe	Pro	Asp	195	200	205	
His	Pro	Leu	His	Ser	Ile	Ser	Val	Arg	Arg	Leu	Lys	Glu	Thr	Lys	Asp	210	215	220	
Gly	Phe	Met	Phe	Leu	Thr	Asp	Gln	Ser	Tyr	Ile	Asp	Val	Leu	Pro	Glu	225	230	235	240
Phe	Arg	Asp	Ser	Tyr	Pro	Ile	Lys	Tyr	Val	His	Ala	Phe	Glu	Ser	Asn	245	250	255	
Asn	Phe	Ile	Tyr	Phe	Leu	Thr	Val	Gln	Arg	Glu	Thr	Leu	Asp	Ala	Gln	260	265	270	
Thr	Phe	His	Thr	Arg	Ile	Ile	Arg	Phe	Cys	Ser	Ile	Asn	Ser	Gly	Leu	275	280	285	
His	Ser	Tyr	Met	Glu	Met	Pro	Leu	Glu	Cys	Ile	Leu	Thr	Glu	Lys	Arg	290	295	300	
Lys	Lys	Arg	Ser	Thr	Lys	Lys	Glu	Val	Phe	Asn	Ile	Leu	Gln	Ala	Ala	305	310	315	320
Tyr	Val	Ser	Lys	Pro	Gly	Ala	Gln	Leu	Ala	Arg	Gln	Ile	Gly	Ala	Ser	325	330	335	
Leu	Asn	Asp	Asp	Ile	Leu	Phe	Gly	Val	Phe	Ala	Gln	Ser	Lys	Pro	Asp	340	345	350	
Ser	Ala	Glu	Pro	Met	Asp	Arg	Ser	Ala	Met	Cys	Ala	Phe	Pro	Ile	Lys	355	360	365	
Tyr	Val	Asn	Asp	Phe	Phe	Asn	Lys	Ile	Val	Asn	Lys	Asn	Asn	Val	Arg	370	375	380	
Cys	Leu	Gln	His	Phe	Tyr	Gly	Pro	Asn	His	Glu	His	Cys	Phe	Asn	Arg	385	390	395	400
Thr	Leu	Leu	Arg	Asn	Ser	Ser	Gly	Cys	Glu	Ala	Arg	Arg	Asp	Glu	Tyr	405	410	415	

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Arg Thr Glu Phe Thr Thr Ala Leu Gln Arg Val Asp Leu Phe Met Gly
      420      425      430
Gln Phe Ser Glu Val Leu Leu Thr Ser Ile Ser Thr Phe Ile Lys Gly
      435      440      445
Asp Leu Thr Ile Ala Asn Leu Gly Thr Ser Glu Gly Arg Phe Met Gln
      450      455      460
Val Val Val Ser Arg Ser Gly Pro Ser Thr Pro His Val Asn Phe Leu
465      470      475      480
Leu Asp Ser His Pro Val Ser Pro Glu Val Ile Val Glu His Thr Leu
      485      490      495
Asn Gln Asn Gly
      500

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<211> 567
<212> PRT
<213> Human

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<400> 752
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Thr Leu Val Gln Arg Ser Asn Gly Glu Cys Lys Glu Ala Leu Ala Lys
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Ser Glu Met Asn Val Asn Met Lys Tyr Gln Leu Pro Asn Phe Thr Ala
      35      40      45
Glu Thr Pro Ile Gln Asn Val Ile Leu His Glu His His Ile Phe Leu
      50      55      60
Gly Ala Thr Asn Tyr Ile Tyr Val Leu Asn Glu Glu Asp Leu Gln Lys
65      70      75      80
Val Ala Glu Tyr Lys Thr Gly Pro Val Leu Glu His Pro Asp Cys Phe
      85      90      95
Pro Cys Gln Asp Cys Ser Ser Lys Ala Asn Leu Ser Gly Gly Val Trp
      100      105      110
Lys Asp Asn Ile Asn Met Ala Leu Val Val Asp Thr Tyr Tyr Asp Asp
      115      120      125
Gln Leu Ile Ser Cys Gly Ser Val Asn Arg Gly Thr Cys Gln Arg His
      130      135      140
Val Phe Pro His Asn His Thr Ala Asp Ile Gln Ser Glu Val His Cys
145      150      155      160
Ile Phe Ser Pro Gln Ile Glu Glu Pro Ser Gln Cys Pro Asp Cys Val
      165      170      175
Val Ser Ala Leu Gly Ala Lys Val Leu Ser Ser Val Lys Asp Arg Phe
      180      185      190
Ile Asn Phe Phe Val Gly Asn Thr Ile Asn Ser Ser Tyr Phe Pro Asp
      195      200      205
His Pro Leu His Ser Ile Ser Val Arg Arg Leu Lys Glu Thr Lys Asp
      210      215      220
Gly Phe Met Phe Leu Thr Asp Gln Ser Tyr Ile Asp Val Leu Pro Glu
225      230      235      240
Phe Arg Asp Ser Tyr Pro Ile Lys Tyr Val His Ala Phe Glu Ser Asn
      245      250      255
Asn Phe Ile Tyr Phe Leu Thr Val Gln Arg Glu Thr Leu Asp Ala Gln
      260      265      270
Thr Phe His Thr Arg Ile Ile Arg Phe Cys Ser Ile Asn Ser Gly Leu
      275      280      285
His Ser Tyr Met Glu Met Pro Leu Glu Cys Ile Leu Thr Glu Lys Arg
      290      295      300

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Lys Lys Arg Ser Thr Lys Lys Glu Val Phe Asn Ile Leu Gln Ala Ala  
 305 310 315 320  
 Tyr Val Ser Lys Pro Gly Ala Gln Leu Ala Arg Gln Ile Gly Ala Ser  
 325 330 335  
 Leu Asn Asp Asp Ile Leu Phe Gly Val Phe Ala Gln Ser Lys Pro Asp  
 340 345 350  
 Ser Ala Glu Pro Met Asp Arg Ser Ala Met Cys Ala Phe Pro Ile Lys  
 355 360 365  
 Tyr Val Asn Asp Phe Phe Asn Lys Ile Val Asn Lys Asn Asn Val Arg  
 370 375 380  
 Cys Leu Gln His Phe Tyr Gly Pro Asn His Glu His Cys Phe Asn Arg  
 385 390 395 400  
 Thr Leu Leu Arg Asn Ser Ser Gly Cys Glu Ala Arg Arg Asp Glu Tyr  
 405 410 415  
 Arg Thr Glu Phe Thr Thr Ala Leu Gln Arg Val Asp Leu Phe Met Gly  
 420 425 430  
 Gln Phe Ser Glu Val Leu Leu Thr Ser Ile Ser Thr Phe Ile Lys Gly  
 435 440 445  
 Asp Leu Thr Ile Ala Asn Leu Gly Thr Ser Glu Gly Arg Phe Met Gln  
 450 455 460  
 Val Val Val Ser Arg Ser Gly Pro Ser Thr Pro His Val Asn Phe Leu  
 465 470 475 480  
 Leu Asp Ser His Pro Val Ser Pro Glu Val Ile Val Glu His Thr Leu  
 485 490 495  
 Asn Gln Asn Gly Tyr Thr Leu Val Ile Thr Gly Lys Lys Ile Thr Lys  
 500 505 510  
 Ile Pro Leu Asn Gly Leu Gly Cys Arg His Phe Gln Ser Cys Ser Gln  
 515 520 525  
 Cys Leu Ser Ala Pro Pro Phe Val Gln Cys Gly Trp Cys His Asp Lys  
 530 535 540  
 Cys Val Arg Ser Glu Glu Cys Leu Ser Gly Thr Trp Thr Gln Gln Ile  
 545 550 555 560  
 Cys Leu Pro Ala Ile Tyr Lys  
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 <212> PRT  
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 Leu Phe Leu Ser Ala Ala Leu Ser Leu Trp Pro Thr Ser Gly Glu Ile  
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 Cys Gly Pro Gly Ile Asp Ile Arg Asn Asp Tyr Gln Gln Leu Lys Arg  
 35 40 45  
 Leu Glu Asn Cys Thr Val Ile Glu Gly Tyr Leu His Ile Leu Leu Ile  
 50 55 60  
 Ser Lys Ala Glu Asp Tyr Arg Ser Tyr Arg Phe Pro Lys Leu Thr Val  
 65 70 75 80  
 Ile Thr Glu Tyr Leu Leu Leu Phe Arg Val Ala Gly Leu Glu Ser Leu  
 85 90 95  
 Gly Asp Leu Phe Pro Asn Leu Thr Val Ile Arg Gly Trp Lys Leu Phe  
 100 105 110  
 Tyr Asn Tyr Ala Leu Val Ile Phe Glu Met Thr Asn Leu Lys Asp Ile  
 115 120 125

Gly	Leu	Tyr	Asn	Leu	Arg	Asn	Ile	Thr	Arg	Gly	Ala	Ile	Arg	Ile	Glu
130						135					140				
Lys	Asn	Ala	Asp	Leu	Cys	Tyr	Leu	Ser	Thr	Val	Asp	Trp	Ser	Leu	Ile
145					150					155					160
Leu	Asp	Ala	Val	Ser	Asn	Asn	Tyr	Ile	Val	Gly	Asn	Lys	Pro	Pro	Lys
				165					170					175	
Glu	Cys	Gly	Asp	Leu	Cys	Pro	Gly	Thr	Met	Glu	Glu	Lys	Pro	Met	Cys
			180					185					190		
Glu	Lys	Thr	Thr	Ile	Asn	Asn	Glu	Tyr	Asn	Tyr	Arg	Cys	Trp	Thr	Thr
	195						200					205			
Asn	Arg	Cys	Gln	Lys	Met	Cys	Pro	Ser	Thr	Cys	Gly	Lys	Arg	Ala	Cys
210						215					220				
Thr	Glu	Asn	Asn	Glu	Cys	Cys	His	Pro	Glu	Cys	Leu	Gly	Ser	Cys	Ser
225					230					235					240
Ala	Pro	Asp	Asn	Asp	Thr	Ala	Cys	Val	Ala	Cys	Arg	His	Tyr	Tyr	Tyr
				245					250					255	
Ala	Gly	Val	Cys	Val	Pro	Ala	Cys	Pro	Pro	Asn	Thr	Tyr	Arg	Phe	Glu
			260					265					270		
Gly	Trp	Arg	Cys	Val	Asp	Arg	Asp	Phe	Cys	Ala	Asn	Ile	Leu	Ser	Ala
	275						280					285			
Glu	Ser	Ser	Asp	Ser	Glu	Gly	Phe	Val	Ile	His	Asp	Gly	Glu	Cys	Met
290						295					300				
Gln	Glu	Cys	Pro	Ser	Gly	Phe	Ile	Arg	Asn	Gly	Ser	Gln	Ser	Met	Tyr
305					310					315					320
Cys	Ile	Pro	Cys	Glu	Gly	Pro	Cys	Pro	Lys	Val	Cys	Glu	Glu	Glu	Lys
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Lys	Thr	Lys	Thr	Ile	Asp	Ser	Val	Thr	Ser	Ala	Gln	Met	Leu	Gln	Gly
			340					345					350		
Cys	Thr	Ile	Phe	Lys	Gly	Asn	Leu	Ile	Asn	Ile	Arg	Arg	Gly	Asn	
	355					360					365				
Asn	Ile	Ala	Ser	Glu	Leu	Glu	Asn	Phe	Met	Gly	Leu	Ile	Glu	Val	Val
370						375					380				
Thr	Gly	Tyr	Val	Lys	Ile	Arg	His	Ser	His	Ala	Leu	Val	Ser	Leu	Ser
385					390					395					400
Phe	Leu	Lys	Asn	Leu	Arg	Leu	Ile	Leu	Gly	Glu	Glu	Gln	Leu	Glu	Gly
			405						410					415	
Asn	Tyr	Ser	Phe	Tyr	Val	Leu	Asp	Asn	Gln	Asn	Leu	Gln	Gln	Leu	Trp
			420					425					430		
Asp	Trp	Asp	His	Arg	Asn	Leu	Thr	Ile	Lys	Ala	Gly	Lys	Met	Tyr	Phe
		435				440					445				
Ala	Phe	Asn	Pro	Lys	Leu	Cys	Val	Ser	Glu	Ile	Tyr	Arg	Met	Glu	Glu
450						455					460				
Val	Thr	Gly	Thr	Lys	Gly	Arg	Gln	Ser	Lys	Gly	Asp	Ile	Asn	Thr	Arg
465					470					475					480
Asn	Asn	Gly	Glu	Arg	Ala	Ser	Cys	Glu	Ser	Asp	Val	Leu	His	Phe	Thr
				485					490					495	
Ser	Thr	Thr	Thr	Ser	Lys	Asn	Arg	Ile	Ile	Ile	Thr	Trp	His	Arg	Tyr
			500					505					510		
Arg	Pro	Pro	Asp	Tyr	Arg	Asp	Leu	Ile	Ser	Phe	Thr	Val	Tyr	Tyr	Lys
		515					520					525			
Glu	Ala	Pro	Phe	Lys	Asn	Val	Thr	Glu	Tyr	Asp	Gly	Gln	Asp	Ala	Cys
	530					535					540				
Gly	Ser	Asn	Ser	Trp	Asn	Met	Val	Asp	Val	Asp	Leu	Pro	Pro	Asn	Lys
545					550					555					560
Asp	Val	Glu	Pro	Gly	Ile	Leu	Leu	His	Gly	Leu	Lys	Pro	Trp	Thr	Gln
				565					570					575	
Tyr	Ala	Val	Tyr	Val	Lys	Ala	Val	Thr	Leu	Thr	Met	Val	Glu	Asn	Asp



			580					585					590				
His	Ile	Arg	Gly	Ala	Lys	Ser	Glu	Ile	Leu	Tyr	Ile	Arg	Thr	Asn	Ala		
		595					600					605					
Ser	Val	Pro	Ser	Ile	Pro	Leu	Asp	Val	Leu	Ser	Ala	Ser	Asn	Ser	Ser		
	610					615					620						
Ser	Gln	Leu	Ile	Val	Lys	Trp	Asn	Pro	Pro	Ser	Leu	Pro	Asn	Gly	Asn		
625					630					635					640		
Leu	Ser	Tyr	Tyr	Ile	Val	Arg	Trp	Gln	Arg	Gln	Pro	Gln	Asp	Gly	Tyr		
				645					650					655			
Leu	Tyr	Arg	His	Asn	Tyr	Cys	Ser	Lys	Asp	Lys	Ile	Pro	Ile	Arg	Lys		
			660					665					670				
Tyr	Ala	Asp	Gly	Thr	Ile	Asp	Ile	Glu	Glu	Val	Thr	Glu	Asn	Pro	Lys		
		675					680					685					
Thr	Glu	Val	Cys	Gly	Gly	Glu	Lys	Gly	Pro	Cys	Cys	Ala	Cys	Pro	Lys		
	690					695					700						
Thr	Glu	Ala	Glu	Lys	Gln	Ala	Glu	Lys	Glu	Glu	Ala	Glu	Tyr	Arg	Lys		
705					710					715					720		
Val	Phe	Glu	Asn	Phe	Leu	His	Asn	Ser	Ile	Phe	Val	Pro	Arg	Pro	Glu		
				725					730					735			
Arg	Lys	Arg	Arg	Asp	Val	Met	Gln	Val	Ala	Asn	Thr	Thr	Met	Ser	Ser		
				740				745					750				
Arg	Ser	Arg	Asn	Thr	Thr	Ala	Ala	Asp	Thr	Tyr	Asn	Ile	Thr	Asp	Pro		
		755					760					765					
Glu	Glu	Leu	Glu	Thr	Glu	Tyr	Pro	Phe	Phe	Glu	Ser	Arg	Val	Asp	Asn		
	770					775					780						
Lys	Glu	Arg	Thr	Val	Ile	Ser	Asn	Leu	Arg	Pro	Phe	Thr	Leu	Tyr	Arg		
785					790					795					800		
Ile	Asp	Ile	His	Ser	Cys	Asn	His	Glu	Ala	Glu	Lys	Leu	Gly	Cys	Ser		
				805				810						815			
Ala	Ser	Asn	Phe	Val	Phe	Ala	Arg	Thr	Met	Pro	Ala	Glu	Gly	Ala	Asp		
			820					825					830				
Asp	Ile	Pro	Gly	Pro	Val	Thr	Trp	Glu	Pro	Arg	Pro	Glu	Asn	Ser	Ile		
		835					840					845					
Phe	Leu	Lys	Trp	Pro	Glu	Pro	Glu	Asn	Pro	Asn	Gly	Leu	Ile	Leu	Met		
	850					855					860						
Tyr	Glu	Ile	Lys	Tyr	Gly	Ser	Gln	Val	Glu	Asp	Gln	Arg	Glu	Cys	Val		
865					870					875					880		
Ser	Arg	Gln	Glu	Tyr	Arg	Lys	Tyr	Gly	Gly	Ala	Lys	Leu	Asn	Arg	Leu		
				885				890					895				
Asn	Pro	Gly	Asn	Tyr	Thr	Ala	Arg	Ile	Gln	Ala	Thr	Ser	Leu	Ser	Gly		
			900					905					910				
Asn	Gly	Ser	Trp	Thr	Asp	Pro	Val	Phe	Phe	Tyr	Val	Gln	Ala	Lys	Thr		
		915					920					925					
Gly	Tyr	Glu	Asn	Phe	Ile	His	Leu	Ile	Ile	Ala	Leu	Pro	Val	Ala	Val		
	930					935				940		</					

Met Arg Glu Arg Ile Glu Phe Leu Asn Glu Ala Ser Val Met Lys Glu  
 1045 1050 1055  
 Phe Asn Cys His His Val Val Arg Leu Leu Gly Val Val Ser Gln Gly  
 1060 1065 1070  
 Gln Pro Thr Leu Val Ile Met Glu Leu Met Thr Arg Gly Asp Leu Lys  
 1075 1080 1085  
 Ser Tyr Leu Arg Ser Leu Arg Pro Glu Met Glu Asn Asn Pro Val Leu  
 1090 1095 1100  
 Ala Pro Pro Ser Leu Ser Lys Met Ile Gln Met Ala Gly Glu Ile Ala  
 1105 1110 1115 1120  
 Asp Gly Met Ala Tyr Leu Asn Ala Asn Lys Phe Val His Arg Asp Leu  
 1125 1130 1135  
 Ala Ala Arg Asn Cys Met Val Ala Glu Asp Phe Thr Val Lys Ile Gly  
 1140 1145 1150  
 Asp Phe Gly Met Thr Arg Asp Ile Tyr Glu Thr Asp Tyr Tyr Arg Lys  
 1155 1160 1165  
 Gly Gly Lys Gly Leu Leu Pro Val Arg Trp Met Ser Pro Glu Ser Leu  
 1170 1175 1180  
 Lys Asp Gly Val Phe Thr Thr Tyr Ser Asp Val Trp Ser Phe Gly Val  
 1185 1190 1195 1200  
 Val Leu Trp Glu Ile Ala Thr Leu Ala Glu Gln Pro Tyr Gln Gly Leu  
 1205 1210 1215  
 Ser Asn Glu Gln Val Leu Arg Phe Val Met Glu Gly Gly Leu Leu Asp  
 1220 1225 1230  
 Lys Pro Asp Asn Cys Pro Asp Met Leu Phe Glu Leu Met Arg Met Cys  
 1235 1240 1245  
 Trp Gln Tyr Asn Pro Lys Met Arg Pro Ser Phe Leu Glu Ile Ile Ser  
 1250 1255 1260  
 Ser Ile Lys Glu Glu Met Glu Pro Gly Phe Arg Glu Val Ser Phe Tyr  
 1265 1270 1275 1280  
 Tyr Ser Glu Glu Asn Lys Leu Pro Glu Pro Glu Glu Leu Asp Leu Glu  
 1285 1290 1295  
 Pro Glu Asn Met Glu Ser Val Pro Leu Asp Pro Ser Ala Ser Ser Ser  
 1300 1305 1310  
 Ser Leu Pro Leu Pro Asp Arg His Ser Gly His Lys Ala Glu Asn Gly  
 1315 1320 1325  
 Pro Gly Pro Gly Val Leu Val Leu Arg Ala Ser Phe Asp Glu Arg Gln  
 1330 1335 1340  
 Pro Tyr Ala His Met Asn Gly Gly Arg Lys Asn Glu Arg Ala Leu Pro  
 1345 1350 1355 1360  
 Leu Pro Gln Ser Ser Thr Cys  
 1365

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 <213> Human

<400> 754  
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 Cys Ser Ala Glu Lys Leu Ala Leu Cys Pro Pro Val Ser Ala Ser Cys  
 35 40 45  
 Ser Glu Val Thr Arg Ser Ala Gly Cys Gly Cys Cys Pro Met Cys Ala  
 50 55 60

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

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 <213> Human

<400> 755

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

[illegible]

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<212>	PRT
<213>	Human

<400> 756															
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Gly	Leu	Gly 35	Pro	Val	Val	Arg	Cys 40	Glu	Pro	Cys	Asp	Ala 45	Arg	Ala	Leu
Ala	Gln 50	Cys	Ala	Pro	Pro	Pro 55	Ala	Val	Cys	Ala	Glu 60	Leu	Val	Arg	Glu
Pro 65	Gly	Cys	Gly	Cys 70	Cys	Leu	Thr	Cys	Ala 75	Leu	Ser	Glu	Gly	Gln 80	Pro
Cys	Gly	Ile	Tyr	Thr 85	Glu	Arg	Cys	Gly	Ser 90	Gly	Leu	Arg	Cys 95	Gln	Pro
Ser	Pro	Asp	Glu 100	Ala	Arg	Pro	Leu	Gln 105	Ala	Leu	Leu	Asp	Gly 110	Arg	Gly
Leu	Cys	Val 115	Asn	Ala	Ser	Ala 120	Val	Ser	Arg	Leu	Arg 125	Ala	Tyr	Leu	Leu
Pro	Ala 130	Pro	Pro	Ala	Pro	Gly 135	Glu	Pro	Pro	Ala 140	Pro	Gly	Asn	Ala	Ser
Glu 145	Ser	Glu	Glu	Asp 150	Arg	Ser	Ala	Gly	Ser	Val 155	Glu	Ser	Pro	Ser	Val
Ser	Ser	Thr	His 165	Arg	Val	Ser	Asp	Pro	Lys 170	Phe	His	Pro	Leu 175	His	Ser
Lys	Ile	Ile	Ile 180	Ile	Lys	Lys	Gly	His 185	Ala	Lys	Asp	Ser	Gln 190	Arg	Tyr
Lys	Val	Asp 195	Tyr	Glu	Ser	Gln 200	Ser	Thr	Asp	Thr	Gln	Asn 205	Phe	Ser	Ser
Glu	Ser 210	Lys	Arg	Glu	Thr	Glu 215	Tyr	Gly	Pro	Cys	Arg 220	Arg	Glu	Met	Glu
Asp 225	Thr	Leu	Asn	His 230	Leu	Lys	Phe	Leu	Asn 235	Val	Leu	Ser	Pro	Arg	Gly
Val	His	Ile	Pro	Asn 245	Cys	Asp	Lys	Lys	Gly 250	Phe	Tyr	Lys	Lys 255	Lys	Gln
Cys	Arg	Pro	Ser	Lys	Gly	Arg	Lys	Arg	Gly	Phe	Cys	Trp	Cys	Val	Asp

			260					265					270						
Lys	Tyr	Gly	Gln	Pro	Leu	Pro	Gly	Tyr	Thr	Thr	Lys	Gly	Lys	Glu	Asp				
		275					280					285							
Val	His	Cys	Tyr	Ser	Met	Gln	Ser	Lys											
	290					295													

<210> 757  
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 <212> PRT  
 <213> Human

<400> 757

Met	Gln	Arg	Ala	Arg	Pro	Thr	Leu	Trp	Ala	Ala	Ala	Leu	Thr	Leu	Leu				
1				5					10					15					
Val	Leu	Leu	Arg	Gly	Pro	Pro	Val	Ala	Arg	Ala	Gly	Ala	Ser	Ser	Ala				
			20					25					30						
Gly	Leu	Gly	Pro	Val	Val	Arg	Cys	Glu	Pro	Cys	Asp	Ala	Arg	Ala	Leu				
		35					40					45							
Ala	Gln	Cys	Ala	Pro	Pro	Pro	Ala	Val	Cys	Ala	Glu	Leu	Val	Arg	Glu				
	50					55					60								
Pro	Gly	Cys	Gly	Cys	Cys	Leu	Thr	Cys	Ala	Leu	Ser	Glu	Gly	Gln	Pro				
65					70				75					80					
Cys	Gly	Ile	Tyr	Thr	Glu	Arg	Cys	Gly	Ser	Gly	Leu	Arg	Cys	Gln	Pro				
				85				90						95					
Ser	Pro	Asp	Glu	Ala	Arg	Pro	Leu	Gln	Ala	Leu	Leu	Asp	Gly	Arg	Gly				
			100					105					110						
Leu	Cys	Val	Asn	Ala	Ser	Ala	Val	Ser	Arg	Leu	Arg	Ala	Tyr	Leu	Leu				
	115						120					125							
Pro	Ala	Pro	Pro	Ala	Pro	Gly	Asn	Ala	Ser	Glu	Ser	Glu	Glu	Asp	Arg				
	130					135					140								
Ser	Ala	Gly	Ser	Val	Glu	Ser	Pro	Ser	Val	Ser	Ser	Thr	His	Arg	Val				
145					150					155					160				
Ser	Asp	Pro	Lys	Phe	His	Pro	Leu	His	Ser	Lys	Ile	Ile	Ile	Ile	Lys				
				165					170					175					
Lys	Gly	His	Ala	Lys	Asp	Ser	Gln	Arg	Tyr	Lys	Val	Asp	Tyr	Glu	Ser				
		180						185					190						
Gln	Ser	Thr	Asp	Thr	Gln	Asn	Phe	Ser	Ser	Glu	Ser	Lys	Arg	Glu	Thr				
		195					200					205							
Glu	Tyr	Gly	Pro	Cys	Arg	Arg	Glu	Met	Glu	Asp	Thr	Leu	Asn	His	Leu				
	210					215				220									
Lys	Phe	Leu	Asn	Val	Leu	Ser	Pro	Arg	Gly	Val	His	Ile	Pro	Asn	Cys				
225					230					235					240				
Asp	Lys	Lys	Gly	Phe	Tyr	Lys	Lys	Lys	Gln	Cys	Arg	Pro	Ser	Lys	Gly				
			245						250					255					
Arg	Lys	Arg	Gly	Phe	Cys	Trp	Cys	Val	Asp	Lys	Tyr	Gly	Gln	Pro	Leu				
		260						265					270						
Pro	Gly	Tyr	Thr	Thr	Lys	Gly	Lys	Glu	Asp	Val	His	Cys	Tyr	Ser	Met				
		275					280					285							
Gln	Ser	Lys																	
	290																		

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 <212> PRT  
 <213> Human

<400> 758

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

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<213> Human

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Pro Ala Gln Ser Leu Gly Ser Phe Val His Cys Glu Pro Cys Asp Glu
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Lys Ala Leu Ser Met Cys Pro Pro Ser Pro Leu Gly Cys Glu Leu Val
          35          40          45
Lys Glu Pro Gly Cys Gly Cys Cys Met Thr Cys Ala Leu Ala Glu Gly
          50          55          60
Gln Ser Cys Gly Val Tyr Thr Glu Arg Cys Ala Gln Gly Leu Arg Cys
65          70          75          80
Leu Pro Arg Gln Asp Glu Glu Lys Pro Leu His Ala Leu Leu His Gly
          85          90          95
Arg Gly Val Cys Leu Asn Glu Lys Ser Tyr Arg Glu Gln Val Lys Ile
          100          105          110
Glu Arg Asp Ser Arg Glu His Glu Glu Pro Thr Thr Ser Glu Met Ala
          115          120          125
Glu Glu Thr Tyr Ser Pro Lys Ile Phe Arg Pro Lys His Thr Arg Ile

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130		135		140
Ser Glu Leu Lys Ala Glu Ala Val Lys Lys Asp Arg Arg Lys Lys Leu				
145		150		155
Thr Gln Ser Lys Phe Val Gly Gly Ala Glu Asn Thr Ala His Pro Arg				
	165		170	
Ile Ile Ser Ala Pro Glu Met Arg Gln Glu Ser Glu Gln Gly Pro Cys				
	180		185	
Arg Arg His Met Glu Ala Ser Leu Gln Glu Leu Lys Ala Ser Pro Arg				
	195		200	
Met Val Pro Arg Ala Val Tyr Leu Pro Asn Cys Asp Arg Lys Gly Phe				
	210		215	
Tyr Lys Arg Lys Gln Cys Lys Pro Ser Arg Gly Arg Lys Arg Gly Ile				
225		230		235
Cys Trp Cys Val Asp Lys Tyr Gly Met Lys Leu Pro Gly Met Glu Tyr				
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Val Asp Gly Asp Phe Gln Cys His Thr Phe Asp Ser Ser Asn Val Glu				
	260		265	
				270

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

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<212> PRT  
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 Thr Ala Pro Cys Ser Arg Gly Ser Ser Trp Ser Ala Asp Leu Asp Lys  
 35 40 45  
 Cys Met Asp Cys Ala Ser Cys Arg Ala Arg Pro His Ser Asp Phe Cys  
 50 55 60  
 Leu Gly Cys Ala Ala Ala Pro Pro Ala Pro Phe Arg Leu Leu Trp  
 65 70 75

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 Gly Lys Asn Asn Arg Pro Ser  
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<210> 767

<211> 9

<212> PRT

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<223> Made in lab

<400> 767

Gln Ala Trp Asp Ser Asn Thr Val Val  
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Ser Gly Ser Ser Ser Asn Ile Gly Ser Asp Tyr Val His  
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<400> 769

Arg Asn Asn Lys Arg Pro Ser  
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Glu Asp Asn Lys Arg Pro Ser  
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Gln Thr Trp Ala Ser Gly Thr Val Leu  
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Ala Ala Trp Asp Asp Ser Leu Asn Gly Trp Val  
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Gly Gly Asn Asn Ile Gly Asp Lys Ser Val His  
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Glu Asp Lys Asn Arg Pro Ala  
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Gln Val Trp Asp Ser Ser Thr Asp His His Val  
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<211> 11

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Gly Gly Asn Asn Ile Gly Thr Thr Ser Val Gln  
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Gln Thr Trp Val Lys Gly Ala Gly Ile  
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Arg Ala Ser Gln Ser Ile Arg Asn Tyr Leu Asn  
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Ala Ala Ser Ser Leu Gln Ser  
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<223> Made in lab

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1 5

<210> 804

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Arg Ala Ser Gln Ser Val Ser Ser Tyr Leu Ala

1 5 10

<210> 805

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Asp Ala Ser Asn Arg Ala Thr

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1 5 10 15

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<210> 809

<211> 9

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Gly Ala Ser Ser Arg Ala Thr

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<210> 812

<211> 5

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Gln Gln Arg Ser Ile

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Val Ala Ser Arg Leu Gln Gly  
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 Asp Ala Tyr Asn Leu Lys Ala  
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Arg Ala Ser Gln Gly Ile Lys Asn Asp Leu Gly  
1 5 10

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<213> Artificial Sequence

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<223> Made in lab

<400> 823

Ala Ala Ser Ser Leu Gln Ser  
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<211> 9

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Gln Gln Ser Asn Ser Phe Pro Leu Thr  
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Gln Ala Ser His Asp Ile Asn Asn Tyr Leu Asn  
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Asp Ala Ser Asn Leu Gln Ser  
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<210> 827

<211> 9

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<213> Artificial Sequence

<220>

<223> Made in lab

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Gln Gln Tyr Asp Thr Leu Pro Val Thr  
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Ala Gly Ser Ser Ser Asn Ile Gly Ser Asn Ser Val Tyr  
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<220>

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<400> 829

Ser Asn Asn Lys Arg Pro Ser  
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<210> 830

<211> 11

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<223> Made in lab

<400> 830

Ala Ala Trp Asp Asp Ser Leu Arg Ser Val Val  
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<210> 831

<211> 13

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<213> Artificial Sequence

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<223> Made in lab

<400> 831

Ser Gly Ser Ser Ser Thr Ile Gly Ser Asn Phe Val Asn  
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<220>  
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 Ala Thr Ser Ser Leu Gln Ser  
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Leu Gln Ala Asn Thr Leu Pro Leu Thr  
 1 5

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<220>  
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 Arg Ala Ser Leu Gly Val Ser Asn Tyr Leu Ala  
 1 5 10

<210> 838  
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<220>  
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 Ala Ala Ser Ile Leu Gln Thr  
 1 5

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<400> 1066

Gly	Leu	Trp	Phe	Gly	Gly	Arg	Leu	Asp	Tyr
1				5				10	

<210> 1067

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1067

His	Tyr	Trp	Met	Lys
1			5	

<210> 1068

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1068

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

Gly

<210> 1069

<211> 18

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1069

Ser Pro Arg Leu Arg Phe Leu Glu Trp Pro Arg Asn Tyr Tyr Gly Met

1	5	10	15
Asp Val			

<210> 1070  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1070  
 Leu Tyr Met Met Val  
 1 5

<210> 1071  
 <211> 15  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1071  
 Tyr Ile Gly Pro Ser Gly Gly Ala Tyr Ala Asp Ser Val Lys Gly  
 1 5 10 15

<210> 1072  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1072  
 Ser Val Arg Gly Leu Thr Phe Asp Tyr  
 1 5

<210> 1073  
 <211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1073  
 Pro Tyr Glu Met Gly  
 1 5

<210> 1074  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in lab

<400> 1074

Arg Ile Ser Pro Ser Gly Gly Met Thr Leu Tyr Ala Asp Ser Val Lys  
1 5 10 15

Gly

<210> 1075

<211> 14

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1075

Met Gly Arg Gly Gly Trp Trp Ala Phe Asp Ala Phe Asp Ile  
1 5 10

<210> 1076

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1076

Trp Tyr Lys Met Val  
1 5

<210> 1077

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1077

Gly Ile Tyr Pro Ser Gly Gly Thr Thr His Tyr Ala Asp Ser Val Lys  
1 5 10 15

Gly

<210> 1078

<211> 14

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1078

Gly Gly Gly Asp Phe Trp Ser Gly Tyr Tyr Pro Phe Asp Tyr

1 5 10

<210> 1079  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1079  
 Gln Ser Ala Leu Thr Gln Asp Pro Ala Val Ser Val Ala Leu Gly Gln  
 1 5 10 15  
 Thr Val Arg Phe Thr Cys Gln Gly Asp Ser Leu Arg Asn Tyr His Pro  
 20 25 30  
 Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr  
 35 40 45  
 Gly Lys Asn Asn Arg Pro Ser Gly Ile Pro Asp Arg Phe Ser Gly Ser  
 50 55 60  
 Thr Ser Gly Asp Thr Ala Ser Leu Thr Ile Thr Gly Ala Gln Ala Glu  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Asn Ser Arg Asp Ser Ser Gly Asn Leu  
 85 90 95  
 Val Phe Phe Ala Gly Gly Thr Lys Leu Thr Val Leu  
 100 105

<210> 1080  
 <211> 106  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1080  
 Gln Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ser Pro Gly Gln  
 1 5 10 15  
 Thr Ala Ser Ile Thr Cys Ser Gly Asp Lys Leu Gly Asp Lys Tyr Ala  
 20 25 30  
 Ser Trp Tyr Gln Gln Lys Pro Asp Gln Ser Pro Val Leu Val Ile Tyr  
 35 40 45  
 Gln Asp Arg Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser  
 50 55 60  
 Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Met  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Ala Trp Asp Ser Asn Thr Val Val  
 85 90 95  
 Phe Gly Gly Gly Thr Lys Leu Thr Val Leu  
 100 105

<210> 1081  
 <211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

&lt;400&gt; 1081

```

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

```

&lt;210&gt; 1082

&lt;211&gt; 106

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1082

```

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

```

&lt;210&gt; 1083

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1083

```

Gln Tyr Glu Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln
 1           5           10           15
Arg Val Thr Ile Ser Cys Ser Gly Ser Ser Ser Asn Ile Gly Ser Asn
      20           25           30
Thr Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu
      35           40           45
Ile Tyr Ala Asn Asn Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser

```

50		55		60
Gly Ser Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln				
65		70		75
Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Val Trp Asp Asp Ser Leu				
	85		90	95
Asn Ala Trp Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu				
	100		105	110

&lt;210&gt; 1084

&lt;211&gt; 107

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1084

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

&lt;210&gt; 1085

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1085

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

&lt;210&gt; 1086

<211> 110  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1086  
 Gln Ser Val Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln  
 1 5 10 15  
 Arg Val Thr Ile Ser Ser Ser Gly Ser Ser Asn Ile Gly Arg Asn  
 20 25 30  
 Ala Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Asn Asn Asn Gln Arg Pro Ser Gly Val Pro Asp Arg Phe Ser  
 50 55 60  
 Ala Ser Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln  
 65 70 75 80  
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu  
 85 90 95  
 Asn Gly Trp Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu  
 100 105 110

<210> 1087  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1087  
 Gln Tyr Glu Leu Thr Gln Ala Pro Ser Val Ser Val Ala Pro Gly Gln  
 1 5 10 15  
 Thr Ala Thr Ile Ser Cys Gly Gly Asn Asn Ile Gly Asp Lys Ser Val  
 20 25 30  
 His Trp Tyr Arg Gln Arg Pro Gly Gln Ala Pro Met Val Val Ile Tyr  
 35 40 45  
 Glu Asp Lys Asn Arg Pro Ala Gly Ile Pro Glu Arg Leu Ser Gly Ser  
 50 55 60  
 Asn Ser Glu Asn Thr Ala Thr Leu Thr Ile Asn Arg Val Glu Ala Glu  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Phe Cys Gln Val Trp Asp Ser Ser Thr Asp His  
 85 90 95  
 His Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu  
 100 105

<210> 1088  
 <211> 106  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1088  
 Gln Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Lys



1				5					10					15		
Thr	Ala	Thr	Ile	Thr	Cys	Gly	Gly	Asn	Asn	Ile	Gly	Thr	Thr	Ser	Val	
			20					25					30			
Gln	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Val	Leu	Ile	Ile	Tyr	
		35					40				45					
Tyr	Gly	Ser	Asp	Arg	Pro	Ser	Arg	Ile	Pro	Glu	Arg	Phe	Ser	Gly	Ser	
	50					55					60					
Asn	Ser	Gly	Asn	Thr	Ala	Thr	Leu	Thr	Ile	Ser	Gly	Ala	Gln	Pro	Val	
65					70					75				80		
Asp	Glu	Ala	Asp	Tyr	Ile	Cys	Gln	Thr	Trp	Val	Lys	Gly	Ala	Gly	Ile	
			85						90					95		
Phe	Gly	Gly	Gly	Thr	Lys	Leu	Thr	Val	Leu							
			100					105								

&lt;210&gt; 1089

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1089

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

&lt;210&gt; 1090

&lt;211&gt; 107

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1090

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

Glu	Asp	Phe	Ala	Thr	Tyr	Phe	Cys	Gln	Gln	Ser	His	Ser	Val	Pro	Leu
				85					90					95	
Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys					
			100					105							

<210> 1091  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1091															
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Gly	Ile	Arg	Asn
			20					25					30		
Asp	Leu	Gly	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Arg	Leu
		35					40					45			
Ile	Tyr	Ala	Ala	Ser	Ser	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Glu	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Gln
65					70					75					80
Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Leu	Gln	His	Asn	Ser	Tyr	Pro
				85					90					95	
Pro	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			100					105							

<210> 1092  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1092															
Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ala	Thr	Leu	Ser	Leu	Ser	Pro
1				5					10					15	
Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys	Arg	Ala	Ser	Gln	Ser	Val	Ser	Ser
			20					25					30		
Asp	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Arg	Leu	Leu
		35					40					45			
Ile	Tyr	Asp	Ala	Phe	Lys	Arg	Ala	Thr	Gly	Ile	Pro	Ala	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu	Glu
65					70					75					80
Pro	Glu	Asp	Phe	Ala	Val	Tyr	Tyr	Cys	Gln	Gln	Arg	Ser	Asn	Trp	Pro
				85					90					95	
Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys				
			100					105							

<210> 1093  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1093

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

&lt;210&gt; 1094

&lt;211&gt; 113

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1094

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

Lys

&lt;210&gt; 1095

&lt;211&gt; 105

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1095

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Gly	Thr	Leu	Ser	Leu	Ser	Pro
1				5					10					15	

Gly	Glu	Arg	Ala	Thr	Leu	Ser	Cys	Arg	Ala	Ser	Gln	Ser	Val	Ser	Ser
			20					25					30		
Ser	Tyr	Leu	Ala	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Arg	Leu
		35					40					45			
Leu	Ile	Tyr	Gly	Ala	Ser	Ser	Arg	Ala	Thr	Gly	Ile	Pro	Asp	Arg	Phe
	50					55					60				
Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser	Ser	Leu
65					70					75					80
Glu	Pro	Glu	Asp	Phe	Ala	Val	Tyr	Tyr	Cys	Gln	Gln	Arg	Ser	Ile	Phe
				85					90					95	
Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Lys							
			100					105							

&lt;210&gt; 1096

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1096

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

&lt;210&gt; 1097

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1097

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

				85					90					95
Leu	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Val	Glu	Ile	Arg			
			100					105						

<210> 1098  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1098

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

<210> 1099  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1099

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

<210> 1100  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1100

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

&lt;210&gt; 1101

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1101

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

&lt;210&gt; 1102

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1102

Gln	Ser	Ala	Leu	Thr	Gln	Pro	Pro	Ser	Val	Ser	Gly	Thr	Pro	Gly	Gln
1				5					10					15	
Lys	Val	Thr	Ile	Ser	Cys	Ser	Gly	Ser	Ser	Ser	Thr	Ile	Gly	Ser	Asn
			20					25					30		
Phe	Val	Asn	Trp	Tyr	Gln	Gln	Leu	Pro	Gly	Thr	Ala	Pro	Lys	Leu	Leu

		35					40				45					
Ile	Tyr	Thr	Asn	Asn	Gln	Arg	Pro	Ser	Gly	Val	Pro	Asp	Arg	Phe	Ser	
	50					55					60					
Gly	Ser	Lys	Ser	Gly	Thr	Ser	Ala	Ser	Leu	Ala	Ile	Ser	Gly	Leu	Gln	
65					70					75					80	
Ser	Glu	Asp	Glu	Ala	Asp	Tyr	Tyr	Cys	Ala	Thr	Trp	Asp	Asp	Asn	Leu	
			85						90					95		
Leu	Gly	Pro	Val	Phe	Gly	Gly	Gly	Thr	Arg	Leu	Ala	Val	Leu			
			100					105					110			

&lt;210&gt; 1103

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1103

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

&lt;210&gt; 1104

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1104

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

<210> 1105  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1105  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Ala Ile Thr Cys Arg Ala Ser Gln Ser Ile Asp Thr  
 20 25 30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Ala Ala Ser Lys Leu Glu Asp Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Thr Gly Thr Asp Phe Thr Leu Thr Ile Arg Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Phe Ala Ser Tyr Phe Cys Gln Gln Ser Tyr Ser Ser Pro  
 85 90 95  
 Gly Ile Thr Phe Gly Pro Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 1106  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1106  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Gln Ala Ser Gln Asp Ile Ser Asp  
 20 25 30  
 Tyr Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Tyr Asp Ala Ser Asn Leu Glu Thr Gly Val Thr Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Tyr Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Ile Ala Thr Tyr Phe Cys Gln Gln Asn Asp Asn Leu Pro  
 85 90 95  
 Phe Thr Phe Gly Pro Gly Thr Lys Val His Ile Lys  
 100 105

<210> 1107  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab



&lt;400&gt; 1107

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

&lt;210&gt; 1108

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1108

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

&lt;210&gt; 1109

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1109

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

Gly	Gly	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Phe	Thr	Ile	Ser	Ser	Leu	Gln
65					70					75					80
Pro	Glu	Asp	Ile	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Asp	Asn	Leu	Pro
				85					90					95	
Tyr	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Leu	Glu	Ile	Lys				
			100					105							

<210> 1110  
 <211> 109  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

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

<210> 1111  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

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

<210> 1112  
 <211> 110

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1112  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu Ser Leu Phe Pro  
 1 5 10 15  
 Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser  
 20 25 30  
 Arg Tyr Leu Ala Trp Tyr Gln Gln Arg Phe Gly Gln Ala Pro Arg Leu  
 35 40 45  
 Leu Ile Tyr Ala Ala Ser Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe  
 50 55 60  
 Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu  
 65 70 75 80  
 Glu Pro Glu Asp Ser Ala Phe Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser  
 85 90 95  
 Pro Pro Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile Lys  
 100 105 110

<210> 1113  
 <211> 107  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1113  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Gly Thr Leu Ser Val Ser Pro  
 1 5 10 15  
 Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe  
 20 25 30  
 Ser Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu  
 35 40 45  
 Ile Tyr Asp Thr Ser Asn Arg Val Ala Gly Ile Pro Ala Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ala Gly Thr Asp Phe Thr Leu Ser Ile Ser Thr Leu Glu  
 65 70 75 80  
 Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln His Arg Ser Asn Trp Pro  
 85 90 95  
 Gly Phe Gly Gln Gly Thr Lys Leu Glu Leu Lys  
 100 105

<210> 1114  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1114  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15

Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Gln	Ala	Ser	Gln	Asp	Ile	Ile	Asn
			20					25					30		
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Lys	Leu	Leu
		35					40					45			
Ile	Tyr	Asp	Ala	Ser	Asn	Leu	Glu	Thr	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Gly	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Phe	Thr	Ile	Ser	Ser	Leu	Gln
65					70					75				80	
Pro	Glu	Asp	Ile	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Tyr	Asp	Asn	Leu	Pro
			85						90					95	
Tyr	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Leu	Glu	Asn	Lys				
			100					105							

&lt;210&gt; 1115

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1115

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

&lt;210&gt; 1116

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1116

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

				85					90					95
Leu	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Val	Glu	Ile	Lys			
			100					105						

<210> 1117  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1117  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Arg Asn  
 20 25 30  
 Asp Leu Gly Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Arg Leu  
 35 40 45  
 Ile Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln His Asn Ser Tyr Pro  
 85 90 95  
 Arg Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 1118  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1118  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ala Thr Leu Ser Val Ser Pro  
 1 5 10 15  
 Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser  
 20 25 30  
 Asn Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu  
 35 40 45  
 Ile Tyr Gly Ala Ser Thr Arg Ala Thr Gly Ile Pro Ala Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln  
 65 70 75 80  
 Ser Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Asn Asn Trp Pro  
 85 90 95  
 Arg Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
 100 105

<210> 1119  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1119

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

&lt;210&gt; 1120

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1120

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

&lt;210&gt; 1121

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1121

Gln	Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val
1				5					10					15	
Gly	Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Gln	Ser	Ile	Gly	Ser
			20					25					30		
Tyr	Leu	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro	Arg	Leu	Leu

		35					40				45				
Ile	Ser	Asp	Ala	Ser	Asn	Leu	Gln	Ser	Gly	Val	Pro	Ser	Arg	Phe	Ser
	50					55					60				
Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ala	Ser	Leu	Gln
65					70					75					80
Pro	Asp	Asp	Phe	Ala	Thr	Tyr	His	Cys	Gln	Gln	Ser	Tyr	Arg	Leu	Phe
			85						90					95	
Pro	Thr	Phe	Gly	Gln	Gly	Thr	Arg	Leu	Glu	Ile	Lys				
			100					105							

&lt;210&gt; 1122

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1122

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

&lt;210&gt; 1123

&lt;211&gt; 108

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1123

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

<210> 1124  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1124  
 Gln Asp Ile Gln Met Thr Gln Ser Pro Ser Ser Leu Ser Ala Ser Val  
 1 5 10 15  
 Gly Asp Arg Val Thr Ile Ser Cys Arg Ser Ser Gln Ser Ile Ala Lys  
 20 25 30  
 Tyr Leu Thr Trp Tyr Gln Gln Lys Ala Gly Lys Ala Pro Lys Leu Leu  
 35 40 45  
 Ile Phe Ala Ala Ser Glu Leu Gln Ser Gly Val Pro Ser Arg Phe Ser  
 50 55 60  
 Gly Ser Gly Ser Gly Thr Asp Tyr Thr Leu Thr Ile Thr Ser Leu Gln  
 65 70 75 80  
 Pro Glu Asp Phe Val Thr Tyr Tyr Cys Gln Gln Thr Tyr Ser Phe Pro  
 85 90 95  
 His Thr Phe Gly Gln Gly Thr Ser Leu Arg Ile Lys  
 100 105

<210> 1125  
 <211> 108  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1125  
 Gln Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln  
 1 5 10 15  
 Thr Ala Lys Ile Thr Cys Ala Gly Asn Asn Ile Gly Ser Lys Ser Val  
 20 25 30  
 His Trp Tyr Gln Gln Arg Pro Gly Gln Ala Pro Val Leu Val Val Tyr  
 35 40 45  
 Asp Asp Ser Asp Arg Pro Pro Gly Ile Pro Glu Arg Leu Ser Gly Ser  
 50 55 60  
 Asn Ser Glu Asn Thr Ala Thr Leu Thr Ile Ser Arg Val Glu Ala Gly  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Val Trp Asp Ser Asp Ser Asp His  
 85 90 95  
 Tyr Val Phe Gly Ser Gly Thr Lys Val Thr Val Leu  
 100 105

<210> 1126  
 <211> 105  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab



&lt;400&gt; 1126

Gln	Tyr	Glu	Leu	Thr	Gln	Pro	Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln
1				5					10					15	
Thr	Ala	Ser	Ile	Thr	Cys	Ser	Gly	Asp	Arg	Leu	Gly	Asp	Lys	Tyr	Ala
			20					25					30		
Ser	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ala	Pro	Val	Leu	Val	Val	Tyr
		35					40					45			
Asp	Asp	Ser	Glu	Arg	Pro	Ser	Gly	Ile	Pro	Glu	Arg	Phe	Ser	Gly	Ser
	50					55					60				
Asn	Ser	Gly	Asn	Thr	Ala	Ala	Leu	Thr	Ile	Gly	Arg	Val	Glu	Ala	Gly
65					70					75					80
Asp	Glu	Ala	Asp	Tyr	Cys	Gln	Val	Trp	Asp	Ser	Ser	Ile	Val	Phe	
			85					90					95		
Gly	Ser	Gly	Thr	Lys	Val	Thr	Val	Leu							
			100					105							

&lt;210&gt; 1127

&lt;211&gt; 110

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1127

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

&lt;210&gt; 1128

&lt;211&gt; 106

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1128

Gln	Tyr	Glu	Leu	Thr	Gln	Pro	Pro	Ser	Val	Ser	Val	Ser	Pro	Gly	Gln
1				5					10					15	
Thr	Ala	Ser	Ile	Thr	Cys	Ser	Gly	Asp	Lys	Leu	Gly	Asp	Lys	Tyr	Ala
			20					25					30		
Cys	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Gln	Ser	Pro	Val	Leu	Val	Ile	Tyr
		35					40					45			
Gln	Asp	Ser	Lys	Arg	Pro	Ser	Gly	Ile	Pro	Glu	Arg	Phe	Ser	Gly	Ser
	50					55					60				

```

Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Met
65          70          75          80
Asp Glu Ala Asp Tyr Tyr Cys Gln Ala Trp Asp Ser Ser Thr Tyr Val
          85          90          95
Phe Gly Thr Gly Thr Lys Val Thr Val Leu
          100          105

```

```

<210> 1129
<211> 110
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1129
Gln Ser Ala Leu Thr Gln Pro Arg Ser Val Ser Gly Ser Pro Gly Gln
1          5          10          15
Ser Val Thr Ile Ser Cys Thr Gly Thr Ser Ser Asp Val Gly Gly His
          20          25          30
Asn Tyr Val Ser Trp Tyr Gln Gln His Pro Gly Lys Ala Pro Lys Leu
          35          40          45
Met Ile Tyr Asp Val Ser Lys Arg Pro Ser Gly Val Pro Asp Arg Phe
          50          55          60
Ser Gly Ser Lys Tyr Gly Asn Thr Ala Ser Leu Thr Ile Ser Gly Leu
65          70          75          80
Gln Ala Glu Asp Glu Ala Asp Phe Tyr Cys Cys Ser Tyr Ala Gly Arg
          85          90          95
Tyr Thr Tyr Val Phe Gly Thr Gly Thr Lys Val Thr Val Leu
          100          105          110

```

```

<210> 1130
<211> 107
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1130
Gln Tyr Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ser Pro Gly Gln
1          5          10          15
Thr Ala Ser Ile Thr Cys Ser Gly Asp Arg Leu Glu Asp Lys Tyr Thr
          20          25          30
Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ser Pro Leu Leu Val Ile Tyr
          35          40          45
Gln Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
          50          55          60
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Met
65          70          75          80
Asp Glu Ala Asp Tyr Phe Cys Gln Ala Trp Asp Ser Ser Ser Ala Tyr
          85          90          95
Val Phe Gly Thr Gly Thr Lys Val Thr Val Leu
          100          105

```

```

<210> 1131
<211> 108

```

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1131  
 Gln Ser Ala Leu Thr Gln Gln Pro Ser Val Ser Val Ala Pro Gly Lys  
 1 5 10 15  
 Thr Ala Thr Ile Thr Cys Gly Gly Asn Ile Gly Ser Lys Ser Val  
 20 25 30  
 His Trp Tyr Gln Gln Arg Pro Gly Gln Ala Pro Val Val Met Phe  
 35 40 45  
 Phe Asp Tyr Asp Arg Pro Ser Gly Ile Pro Asp Arg Phe Ser Gly Ser  
 50 55 60  
 Asn Ser Gly Asp Thr Ala Thr Leu Thr Ile Ser Arg Val Glu Ala Gly  
 65 70 75 80  
 Asp Glu Ala Asp Tyr Tyr Cys Gln Val Trp Asp Ser Arg Thr Asp Arg  
 85 90 95  
 Tyr Val Phe Gly Thr Gly Thr Thr Val Thr Val Leu  
 100 105

<210> 1132  
 <211> 126  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1132  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ile Tyr  
 20 25 30  
 Asp Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Trp Ile Ser Pro Ser Gly Gly Arg Thr Leu Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Thr Trp Asp Tyr Tyr Asp Ser Ser Gly Tyr Phe Asn Asp Ala  
 100 105 110  
 Phe Asp Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser  
 115 120 125

<210> 1133  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1133

```

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

```

```

<210> 1134
<211> 117
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1134
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ala Tyr
      20          25          30
Gln Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35          40          45
Ser Ser Ile Ser Ser Ser Gly Gly Tyr Thr Ser Tyr Ala Asp Ser Val
      50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
      85          90          95
Ala Arg Glu Ser Arg Tyr Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
      100          105          110
Val Thr Val Ser Ser
      115

```

```

<210> 1135
<211> 122
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1135
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Glu Tyr
      20          25          30
Pro Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val

```

		35					40					45					
Ser	Gly	Ile	Gly	Ser	Ser	Gly	Gly	Ser	Thr	Thr	Tyr	Ala	Asp	Ser	Val		
	50					55					60						
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr		
65					70					75					80		
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys		
			85						90					95			
Ala	Arg	Ala	Pro	Leu	Tyr	Ser	Ser	Thr	Ser	Tyr	Ala	Phe	Asp	Ile	Trp		
			100					105					110				
Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser								
		115					120										

&lt;210&gt; 1136

&lt;211&gt; 116

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1136

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

&lt;210&gt; 1137

&lt;211&gt; 120

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1137

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

```

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
      85          90          95
Ala Arg Gly Gln Met Trp Pro Gly Val Ala Phe Glu Met Trp Gly Gln
      100          105          110
Gly Thr Met Val Thr Val Ser Ser
      115          120

```

```

<210> 1138
<211> 120
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1138
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr
      20          25          30
Phe Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35          40          45
Ser Ser Ile Gly Ser Ser Asp Gly Tyr Thr Arg Tyr Ala Asp Ser Val
      50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
      85          90          95
Ala Arg Asp Leu Ser Trp Trp Pro Asp Ala Phe Asp Ile Trp Gly Gln
      100          105          110
Gly Thr Met Val Thr Val Ser Ser
      115          120

```

```

<210> 1139
<211> 104
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1139
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr
      20          25          30
Arg Met Glu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35          40          45
Ser Trp Ile Tyr Ser Ser Gly Gly Ile Thr Asn Tyr Ala Asp Ser Val
      50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Thr Tyr Tyr Cys
      85          90          95
Ala Arg Leu Val Thr Val Ser Ser
      100

```

<210> 1140  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1140  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Val Tyr  
 20 25 30  
 Asp Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Gly Pro Ser Gly Gly Trp Thr Gly Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asp Ser Gly Gly Trp Glu Ala Leu Tyr Tyr Tyr Tyr Tyr Met  
 100 105 110  
 Asp Val Trp Gly Lys Gly Thr Thr Val Thr Val Ser Ser  
 115 120 125

<210> 1141  
 <211> 121  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1141  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Val Tyr  
 20 25 30  
 Phe Met Asp Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Gly Ile Gly Pro Ser Gly Gly Val Thr Ser Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Ala Gly Gln Leu Ala Gln Gly His Tyr Tyr Met Asp Val Trp Gly  
 100 105 110  
 Lys Gly Thr Thr Val Thr Val Ser Ser  
 115 120

<210> 1142  
 <211> 119  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1142

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

&lt;210&gt; 1143

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1143

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

&lt;210&gt; 1144

&lt;211&gt; 122

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1144

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----



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

&lt;210&gt; 1145

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1145

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

&lt;210&gt; 1146

&lt;211&gt; 119

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1146

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly		
1				5					10					15			
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Ser	Tyr		
			20					25					30				
Val	Met	Val	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val		
		35					40					45					

```

Ser Val Ile Ser Pro Ser Gly Gly Val Thr Phe Tyr Ala Asp Ser Val
 50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Asp Arg Arg Ser Asn Ser Leu Phe Asp Pro Trp Gly Gln Gly
          100          105          110
Thr Leu Val Thr Val Ser Ser
          115

```

<210> 1147  
 <211> 119  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

```

<400> 1147
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Thr Tyr
          20          25          30
Gly Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Tyr Ile Trp Pro Ser Gly Gly Leu Thr Trp Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Ser Gly Tyr Ser Tyr Gly Arg Phe Asp Tyr Trp Gly Gln Gly
          100          105          110
Thr Leu Val Thr Val Ser Ser
          115

```

<210> 1148  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

```

<400> 1148
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr
          20          25          30
Asp Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Trp Ile Ser Pro Ser Gly Gly Ser Thr Leu Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys

```

				85					90					95		
Ala	Ser	Ser	Gly	Leu	Tyr	Gly	Ser	Gly	Ser	Tyr	Ala	Ala	Phe	Asp	Val	
			100					105					110			
Trp	Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser						
		115					120									

<210> 1149  
 <211> 117  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1149

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

<210> 1150  
 <211> 129  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1150

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

Ser

<210> 1151  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1151  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Val Tyr  
 20 25 30  
 Trp Met Glu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Ser Ser Ser Gly Gly Leu Thr Ser Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Thr Thr Asp Leu Val Ser Asn Trp Pro Trp Gly Gly Tyr Trp Gly Gln  
 100 105 110  
 Gly Thr Leu Val Thr Val Ser Ser  
 115 120

<210> 1152  
 <211> 125  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1152  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr  
 20 25 30  
 Lys Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Ser Ser Ser Gly Gly Asp Thr Ala Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asp Arg Ala Pro Tyr Tyr Tyr Asp Ser Ser Gly Tyr Tyr Ser  
 100 105 110  
 Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser  
 115 120 125

<210> 1153  
 <211> 119

<212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1153  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr  
 20 25 30  
 Ala Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Ser Pro Ser Gly Gly Tyr Thr Tyr Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Glu Ser Gly Thr Thr Asn Ala Phe Asp Ile Trp Gly Gln Gly  
 100 105 110  
 Thr Met Val Thr Val Ser Ser  
 115

<210> 1154  
 <211> 117  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1154  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Tyr  
 20 25 30  
 His Met Glu Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Tyr Ile Ser Pro Ser Gly Gly Ser Thr His Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Asp Gly Trp Thr Val Pro Arg Asp Trp Gly Gln Gly Thr Leu  
 100 105 110  
 Val Thr Val Ser Ser  
 115

<210> 1155  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

&lt;400&gt; 1155

```

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

```

&lt;210&gt; 1156

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1156

```

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

```

&lt;210&gt; 1157

&lt;211&gt; 115

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1157

```

Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
 1           5           10           15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr

```

			20					25					30				
Ser	Met	Val	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val		
		35					40					45					
Ser	Arg	Ile	Arg	Pro	Ser	Gly	Gly	Gln	Thr	Met	Tyr	Ala	Asp	Ser	Val		
		50				55					60						
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr		
65					70				75					80			
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys		
			85						90					95			
Ala	Thr	Gly	Tyr	Ala	Phe	Asp	Ile	Trp	Gly	Gln	Gly	Thr	Met	Val	Thr		
		100						105					110				
Val	Ser	Ser															
		115															

&lt;210&gt; 1158

&lt;211&gt; 125

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1158

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

&lt;210&gt; 1159

&lt;211&gt; 125

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1159

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

Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
65					70					75					80
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
				85					90					95	
Thr	Thr	Gly	Ile	Ala	Ala	Ala	Gly	Asn	Tyr	Tyr	Tyr	Tyr	Tyr	Gly	Met
			100					105					110		
Asp	Val	Trp	Gly	Gln	Gly	Thr	Thr	Val	Thr	Val	Ser	Ser			
		115					120					125			

&lt;210&gt; 1160

&lt;211&gt; 124

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1160

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

&lt;210&gt; 1161

&lt;211&gt; 119

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1161

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
1				5					10					15	
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Asn	Tyr
			20					25					30		
Ala	Met	Thr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
		35				40						45			
Ser	Ser	Ile	Tyr	Ser	Ser	Gly	Gly	Asp	Thr	Ala	Tyr	Ala	Asp	Tyr	Val
	50				55						60				
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
65					70					75					80
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
				85					90					95	
Ala	Arg	Glu	Tyr	Tyr	Thr	Gly	Trp	Asn	Phe	Asp	Tyr	Trp	Gly	Gln	Gly



100  
Thr Leu Val Thr Val Ser Ser  
115

```
<210> 1162
<211> 122
<212> PRT
<213> Artificial Sequence
```

<220>  
<223> Made in lab

<400> 1162															
Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
1				5					10					15	
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Gln	Tyr
			20					25					30		
Asp	Met	Val	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
		35				40						45			
Ser	Tyr	Ile	Tyr	Ser	Ser	Gly	Gly	His	Thr	Leu	Tyr	Val	Asp	Ser	Val
	50					55					60				
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
65					70				75					80	
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Thr	Tyr	Tyr	Cys
			85					90						95	
Ala	Arg	Ile	Arg	Ser	Ser	Gly	Tyr	Tyr	His	Glu	Val	Leu	Asp	Tyr	Trp
			100					105					110		
Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser						
		115					120								

```
<210> 1163
<211> 123
<212> PRT
<213> Artificial Sequence
```

<220>  
<223> Made in lab

<400> 1163															
Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
1				5					10					15	
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	Thr	Tyr
			20					25					30		
Met	Met	Tyr	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val
		35					40					45			
Ser	Val	Ile	Gly	Pro	Ser	Gly	Gly	Ala	Thr	Gly	Tyr	Ala	Asp	Ser	Val
	50					55					60				
Lys	Gly	Arg	Phe	Thr	Ile	Ser	Arg	Asp	Asn	Ser	Lys	Asn	Thr	Leu	Tyr
65					70				75					80	
Leu	Gln	Met	Asn	Ser	Leu	Arg	Ala	Glu	Asp	Thr	Ala	Val	Tyr	Tyr	Cys
			85						90					95	
Ala	Arg	Ile	Arg	Lys	Ala	Phe	Gly	Tyr	Gly	Ser	Gly	Ser	Leu	Asp	Tyr
			100					105					110		
Trp	Gly	Gln	Gly	Thr	Leu	Val	Thr	Val	Ser	Ser					
		115					120								

<210> 1164

<211> 126  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1164  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr  
 20 25 30  
 Asp Met Gln Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Tyr Ile Gly Pro Ser Gly Gly Asp Thr Asp Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Ser Ser Ser Tyr Tyr Tyr Asp Ser Ser Gly Tyr Tyr His Glu Ala  
 100 105 110  
 Phe Asp Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser  
 115 120 125

<210> 1165  
 <211> 127  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1165  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Tyr Tyr  
 20 25 30  
 Met Met Arg Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Tyr Ile Gly Pro Ser Gly Gly Ala Thr Thr Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Gly Arg Ser Val Lys Tyr Tyr Asp Ser Ser Gly Tyr Leu  
 100 105 110  
 Leu Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser  
 115 120 125

<210> 1166  
 <211> 122  
 <212> PRT  
 <213> Artificial Sequence

<220>

<223> Made in lab

<400> 1166

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

<210> 1167

<211> 122

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1167

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

<210> 1168

<211> 118

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1168

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
1				5					10					15	

```

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser His Tyr
      20              25              30
Trp Met Met Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35              40              45
Ser Ser Ile Val Pro Ser Gly Gly Asp Thr Tyr Tyr Ala Asp Ser Val
      50              55              60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65      70              75              80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
      85              90              95
Thr Thr Asp Pro Val Met Thr Pro Val Asp Tyr Trp Gly Gln Gly Thr
      100             105             110
Leu Val Thr Val Ser Ser
      115

```

```

<210> 1169
<211> 123
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1169
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
1      5              10              15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr
      20              25              30
Phe Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35              40              45
Ser Ser Ile Tyr Pro Ser Gly Gly Ile Thr Lys Tyr Ala Asp Ser Val
      50              55              60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65      70              75              80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Thr Tyr Tyr Cys
      85              90              95
Ala Lys Glu Thr Tyr Tyr Tyr Gly Ser Gly Ser Tyr Ala Phe Asp Ile
      100             105             110
Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
      115             120

```

```

<210> 1170
<211> 117
<212> PRT
<213> Artificial Sequence

```

```

<220>
<223> Made in lab

```

```

<400> 1170
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
1      5              10              15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Gln Tyr
      20              25              30
Tyr Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
      35              40              45
Ser Arg Ile Ser Pro Ser Gly Gly Met Thr Ser Tyr Ala Asp Ser Val

```

50		55		60	
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr					
65		70		75	80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Met Tyr Tyr Cys					
	85		90		95
Ala Arg His Lys Tyr Gly Gly Pro Asp Phe Trp Gly Gln Gly Thr Leu					
	100		105		110
Val Thr Val Ser Ser					
	115				

&lt;210&gt; 1171

&lt;211&gt; 115

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1171

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

&lt;210&gt; 1172

&lt;211&gt; 113

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1172

Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly					
1		5		10	15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr					
	20		25		30
Thr Met Ala Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val					
	35		40		45
Ser Tyr Ile Ser Pro Ser Gly Gly Phe Thr Gly Tyr Ala Asp Ser Val					
	50		55		60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr					
65		70		75	80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys					
	85		90		95

Ala Thr Trp Gly Asp Pro Trp Gly Gln Gly Thr Leu Val Thr Val Ser  
100 105 110

Ser

<210> 1173

<211> 126

<212> PRT

<213> Artificial Sequence

 $\langle 220 \rangle$ 

<223> Made in lab

<400> 1173

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

<210> 1174

<211> 116

&lt;212&gt; PRT

<213> Artificial Sequence

<220>

<223> Made in lab

<400> 1174

[illegible]

<210> 1175  
 <211> 124  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1175  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr  
 20 25 30  
 His Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Tyr Pro Ser Gly Gly Phe Thr Ala Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Glu Ser Ala Tyr Tyr Tyr Asp Ser Ser Pro Pro Ala Phe Asp  
 100 105 110  
 Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser  
 115 120

<210> 1176  
 <211> 117  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1176  
 Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
 1 5 10 15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Thr Tyr  
 20 25 30  
 Ala Met Tyr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
 35 40 45  
 Ser Ser Ile Tyr Ser Ser Gly Gly Ala Thr Trp Tyr Ala Asp Ser Val  
 50 55 60  
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr  
 65 70 75 80  
 Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Ser Thr Phe Asp Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu  
 100 105 110  
 Val Thr Val Ser Ser  
 115

<210> 1177  
 <211> 121  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1177

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

&lt;210&gt; 1178

&lt;211&gt; 128

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1178

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

&lt;210&gt; 1179

&lt;211&gt; 120

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1179

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly
-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----	-----



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

&lt;210&gt; 1180

&lt;211&gt; 117

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1180

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

&lt;210&gt; 1181

&lt;211&gt; 127

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in lab

&lt;400&gt; 1181

Glu	Val	Gln	Leu	Leu	Glu	Ser	Gly	Gly	Gly	Leu	Val	Gln	Pro	Gly	Gly		
1				5					10					15			
Ser	Leu	Arg	Leu	Ser	Cys	Ala	Ala	Ser	Gly	Phe	Thr	Phe	Ser	His	Tyr		
			20					25					30				
Trp	Met	Lys	Trp	Val	Arg	Gln	Ala	Pro	Gly	Lys	Gly	Leu	Glu	Trp	Val		
		35					40					45					

```

Ser Gly Ile Ser Ser Ser Gly Gly Gln Thr Asp Tyr Ala Asp Ser Val
  50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
  65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
          85          90          95
Ala Arg Ser Pro Arg Leu Arg Phe Leu Glu Trp Pro Arg Asn Tyr Tyr
          100          105          110
Gly Met Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser
          115          120          125

```

<210> 1182  
 <211> 116  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

```

<400> 1182
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
  1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Leu Tyr
          20          25          30
Met Met Val Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Tyr Ile Gly Pro Ser Gly Gly Ala Tyr Ala Asp Ser Val Lys Gly
          50          55          60
Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu Gln
          65          70          75          80
Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala Arg
          85          90          95
Ser Val Arg Gly Leu Thr Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val
          100          105          110
Thr Val Ser Ser
          115

```

<210> 1183  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

```

<400> 1183
Glu Val Gln Leu Leu Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly
  1          5          10          15
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Pro Tyr
          20          25          30
Glu Met Gly Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
          35          40          45
Ser Arg Ile Ser Pro Ser Gly Gly Met Thr Leu Tyr Ala Asp Ser Val
          50          55          60
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
          65          70          75          80
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys

```

				85					90					95			
Ala	Arg	Met	Gly	Arg	Gly	Gly	Trp	Trp	Ala	Phe	Asp	Ala	Phe	Asp	Ile		
			100						105					110			
Trp	Gly	Gln	Gly	Thr	Met	Val	Thr	Val	Ser	Ser							
			115						120								

<210> 1184  
 <211> 123  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

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

<210> 1185  
 <211> 23  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

Met	Asp	Phe	Gln	Val	Gln	Ile	Phe	Ser	Phe	Leu	Leu	Ile	Ser	Ala	Ser		
1				5					10					15			
Val	Ile	Met	Ser	Arg	Gly	Xaa											
			20														

<210> 1186  
 <211> 20  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

Met	Glu	Ala	Pro	Ala	Gln	Leu	Leu	Phe	Leu	Leu	Leu	Leu	Trp	Leu	Pro		
1				5					10					15			

Asp Thr Thr Gly  
20

<210> 1187  
<211> 15  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1187  
Arg Ala Ser Glu Ser Val His Asn Phe Gly Ile Ser Phe Met Asn  
1 5 10 15

<210> 1188  
<211> 7  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1188  
Thr Ala Ser Asn Gln Gly Ser  
1 5

<210> 1189  
<211> 9  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1189  
His Gln Gly Lys Glu Val Pro Trp Thr  
1 5

<210> 1190  
<211> 15  
<212> PRT  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1190  
Arg Ala Ser Glu Ser Val Gly Asn Phe Gly Ile Ser Phe Val Asn  
1 5 10 15

<210> 1191  
<211> 7  
<212> PRT  
<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1191

Ala Ala Ser Asn Gln Gly Ser

1 5

<210> 1192

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1192

Gln Gln Ser Lys Glu Val Pro Tyr Thr

1 5

<210> 1193

<211> 6

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1193

Asn Thr Tyr Ala Met Asn

1 5

<210> 1194

<211> 19

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1194

Arg Thr Val Thr Lys Ser Asn Lys Tyr Ala Thr Tyr Tyr Ala Asp Ser

1 5 10 15

Val Ser Asp

<210> 1195

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1195

Glu Asp Tyr Tyr Gly Thr Leu Asp Tyr

1 5

<210> 1196

<211> 5  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1196  
 Ser Tyr Trp Met His  
 1 5

<210> 1197  
 <211> 17  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1197  
 Glu Ile Asn Pro Ser Asn Gly Arg Thr Asn Tyr Asn Glu Arg Phe Lys  
 1 5 10 15  
 Ser

<210> 1198  
 <211> 9  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1198  
 Ser Asp Thr Lys Ala Thr Cys Asp Tyr  
 1 5

<210> 1199  
 <211> 111  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1199  
 Glu Ile Val Leu Thr Gln Ser Pro Pro Ser Leu Ala Val Ser Leu Gly  
 1 5 10 15  
 Gln Arg Ala Thr Ile Ser Cys Arg Ala Ser Glu Ser Val His Asn Phe  
 20 25 30  
 Gly Ile Ser Phe Met Asn Trp Phe Gln Gln Lys Ser Gly Gln Pro Pro  
 35 40 45  
 Lys Leu Leu Ile Tyr Thr Ala Ser Asn Gln Gly Ser Gly Val Pro Ala  
 50 55 60  
 Arg Phe Ser Gly Arg Gly Ser Gly Thr Asp Phe Ser Leu Ile Ile His  
 65 70 75 80  
 Pro Val Glu Glu Asp Asp Thr Ala Met Tyr Phe Cys His Gln Gly Lys

				85					90					95	
Glu	Val	Pro	Trp	Thr	Phe	Gly	Gly	Gly	Thr	Lys	Leu	Glu	Ile	Lys	
				100				105					110		

<210> 1200  
 <211> 111  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1200  
 Asp Ile Val Leu Thr Gln Ser Pro Ala Ser Leu Ala Val Ser Leu Gly  
 1 5 10 15  
 Gln Arg Ala Thr Ile Ser Cys Arg Ala Ser Glu Ser Val Gly Asn Phe  
 20 25 30  
 Gly Ile Ser Phe Val Asn Trp Phe Gln Gln Lys Pro Gly Gln Pro Pro  
 35 40 45  
 Lys Leu Leu Ile Tyr Ala Ala Ser Asn Gln Gly Ser Gly Val Pro Ala  
 50 55 60  
 Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Ser Leu Asn Ile His  
 65 70 75 80  
 Pro Met Glu Glu Asp Asp Thr Ala Met Tyr Phe Cys Gln Gln Ser Lys  
 85 90 95  
 Glu Val Pro Trp Thr Phe Gly Gly Gly Thr Lys Leu Glu Ile Lys  
 100 105 110

<210> 1201  
 <211> 111  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1201  
 Glu Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly  
 1 5 10 15  
 Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Glu Ser Val His Asn Phe  
 20 25 30  
 Gly Ile Ser Phe Met Asn Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro  
 35 40 45  
 Arg Leu Leu Ile Tyr Thr Ala Ser Asn Gln Gly Ser Gly Ile Pro Ala  
 50 55 60  
 Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser  
 65 70 75 80  
 Ser Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys His Gln Gly Lys  
 85 90 95  
 Glu Val Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys  
 100 105 110

<210> 1202  
 <211> 111  
 <212> PRT  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1202

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

&lt;210&gt; 1203

&lt;211&gt; 111

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1203

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

&lt;210&gt; 1204

&lt;211&gt; 111

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1204

Asp	Ile	Gln	Met	Thr	Gln	Ser	Pro	Ser	Ser	Leu	Ser	Ala	Ser	Val	Gly
1				5					10					15	
Asp	Arg	Val	Thr	Ile	Thr	Cys	Arg	Ala	Ser	Glu	Ser	Val	Gly	Asn	Phe
			20					25					30		
Gly	Ile	Ser	Phe	Val	Asn	Trp	Tyr	Gln	Gln	Lys	Pro	Gly	Lys	Ala	Pro



		35					40					45					
Lys	Leu	Leu	Ile	Tyr	Ala	Ala	Ser	Asn	Gln	Gly	Ser	Gly	Val	Pro	Ser		
	50					55					60						
Arg	Phe	Ser	Gly	Ser	Gly	Ser	Gly	Thr	Asp	Phe	Thr	Leu	Thr	Ile	Ser		
65					70				75						80		
Ser	Leu	Gln	Pro	Glu	Asp	Phe	Ala	Thr	Tyr	Tyr	Cys	Gln	Gln	Ser	Lys		
				85					90					95			
Glu	Val	Pro	Tyr	Thr	Phe	Gly	Gln	Gly	Thr	Lys	Leu	Glu	Ile	Lys			
			100					105					110				

&lt;210&gt; 1205

&lt;211&gt; 120

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1205

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

&lt;210&gt; 1206

&lt;211&gt; 118

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1206

Gln	Val	Gln	Leu	Gln	Gln	Pro	Gly	Ala	Glu	Leu	Val	Lys	Pro	Gly	Ala		
1				5					10					15			
Ser	Val	Lys	Leu	Ser	Cys	Lys	Ala	Ser	Gly	Tyr	Ile	Phe	Thr	Ser	Tyr		
			20					25				30					
Trp	Met	His	Trp	Val	Lys	Gln	Arg	Pro	Gly	Gln	Gly	Leu	Glu	Trp	Ile		
		35				40					45						
Gly	Glu	Ile	Asn	Pro	Ser	Asn	Gly	Arg	Thr	Asn	Tyr	Asn	Glu	Arg	Phe		
	50				55				60								
Lys	Ser	Lys	Ala	Thr	Leu	Thr	Val	Asp	Lys	Ser	Ser	Ser	Thr	Ala	Tyr		
65				70				75							80		
Ile	Gln	Leu	Ser	Ser	Leu	Thr	Ser	Asp	Asp	Ser	Ala	Val	Tyr	Tyr	Cys		
			85					90						95			

Ser Arg Ser Asp Thr Lys Ala Thr Cys Asp Tyr Trp Gly Gln Gly Thr  
                   100                  105                  110  
 Thr Leu Thr Val Ser Ser  
                   115

<210> 1207  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1207  
 Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Thr Tyr  
                   20                  25                  30  
 Ala Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Gly Arg Thr Val Thr Lys Ser Asn Lys Tyr Ala Thr Tyr Tyr Ala Asp  
                   50                  55                  60  
 Ser Val Ser Asp Arg Phe Thr Ile Ser Arg Asp Asp Ser Lys Asn Ser  
   65                  70                  75                  80  
 Leu Tyr Leu Gln Met Asn Ser Leu Lys Thr Glu Asp Thr Ala Val Tyr  
                   85                  90                  95  
 Tyr Cys Ala Arg Glu Asp Tyr Tyr Gly Thr Leu Asp Tyr Trp Gly Gln  
                   100                  105                  110  
 Gly Thr Leu Val Thr Val Ser Ser  
                   115                  120

<210> 1208  
 <211> 120  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1208  
 Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly  
   1                  5                  10                  15  
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Thr Tyr  
                   20                  25                  30  
 Ala Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val  
                   35                  40                  45  
 Gly Arg Thr Val Thr Lys Ser Asn Lys Tyr Ala Thr Tyr Tyr Ala Asp  
                   50                  55                  60  
 Ser Val Ser Asp Arg Phe Thr Ile Ser Arg Asp Asp Ser Lys Asn Ser  
   65                  70                  75                  80  
 Leu Tyr Leu Gln Met Asn Ser Leu Lys Thr Glu Asp Thr Ala Val Tyr  
                   85                  90                  95  
 Tyr Cys Ala Arg Glu Asp Tyr Tyr Gly Thr Leu Asp Tyr Trp Gly Gln  
                   100                  105                  110  
 Gly Thr Leu Val Thr Val Ser Ser  
                   115                  120

<210> 1209  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1209  
 Gln Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys Pro Gly Ala  
 1 5 10 15  
 Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Ile Phe Thr Ser Tyr  
 20 25 30  
 Trp Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met  
 35 40 45  
 Gly Glu Ile Asn Pro Ser Asn Gly Arg Thr Asn Tyr Asn Glu Arg Phe  
 50 55 60  
 Lys Ser Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Thr Ala Tyr  
 65 70 75 80  
 Met Glu Leu Ser Arg Leu Arg Ser Asp Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Ser Asp Thr Lys Ala Thr Cys Asp Tyr Trp Gly Gln Gly Thr  
 100 105 110  
 Thr Val Thr Val Ser Ser  
 115

<210> 1210  
 <211> 118  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1210  
 Gln Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys Pro Gly Ala  
 1 5 10 15  
 Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Ile Phe Thr Ser Tyr  
 20 25 30  
 Trp Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met  
 35 40 45  
 Gly Glu Ile Asn Pro Ser Asn Gly Arg Thr Asn Tyr Asn Glu Arg Phe  
 50 55 60  
 Lys Ser Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Thr Ala Tyr  
 65 70 75 80  
 Met Glu Leu Ser Arg Leu Arg Ser Asp Asp Thr Ala Val Tyr Tyr Cys  
 85 90 95  
 Ala Arg Ser Asp Thr Lys Ala Thr Cys Asp Tyr Trp Gly Gln Gly Thr  
 100 105 110  
 Thr Val Thr Val Ser Ser  
 115

<210> 1211  
 <211> 333  
 <212> DNA  
 <213> Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1211

gagattgtgc	tgacccaatc	tccaccttct	ttggctgtgt	ctctggggca	gagggccacc	60
atctcctgca	gagccagcga	aagtgttcat	aattttggca	ttagttttat	gaactggttt	120
caacagaaat	caggacagcc	acccaaactc	ctcatctata	ctgcatccaa	ccaaggatcc	180
ggggtccctg	ccaggttttag	tggcagaggg	tctgggacag	acttcagtct	catcatccac	240
cctgtggagg	aagatgatac	tgcaatgtat	ttctgtcacc	aaggtaagga	ggttccgtgg	300
acgttcggtg	gaggcaccaa	gctagaaatc	aaa			333

&lt;210&gt; 1212

&lt;211&gt; 333

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1212

gacattgtgc	tgacccaatc	tccagcttct	ttggctgtgt	ctctagggca	gagggccacc	60
atctcctgca	gagccagcga	aagtgttggt	aattttggca	ttagttttgt	gaactggttc	120
caacagaaac	caggacagcc	acccaaactc	ctcatctatg	ctgcctccaa	ccaaggatcc	180
ggggtccctg	ccaggttttag	tggcagtggt	tctgggacag	acttcagcct	caacatccat	240
cctatggagg	aggatgatac	tgcaatgtat	ttctgtcagc	aaagtaagga	ggttccgtac	300
acgttcggag	gggggaccaa	gctggaaata	aaa			333

&lt;210&gt; 1213

&lt;211&gt; 333

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1213

gaaattgtgt	tgacacagtc	tccagccacc	ctgtctttgt	ctccagggga	aagagccacc	60
ctctcctgca	gagccagcga	aagtgttcat	aattttggca	ttagttttat	gaactgggtac	120
caacagaaac	ctggccaggc	tcccaggctc	ctcatctata	ctgcatccaa	ccaaggatcc	180
ggcatccag	ccaggttcag	tggcagtggt	tctgggacag	acttcactct	caccatcagc	240
agcctagagc	ctgaagattt	tgcagtttat	tactgtcacc	aaggtaagga	ggttccgtgg	300
acgttcggcc	aagggaccaa	ggtggaaatc	aaa			333

&lt;210&gt; 1214

&lt;211&gt; 333

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1214

gacattgtgc	tgacccagtc	tccagcctcc	ttggccgtgt	ctccaggaca	gagggccacc	60
atcacctgca	gagccagcga	aagtgttcat	aattttggca	ttagttttat	gaactgggtat	120
cagcagaaac	caggacaacc	tcctaaactc	ctgatttaca	ctgcatccaa	ccaaggatcc	180
ggggtccag	ccaggttcag	cggcagtggt	tctgggaccg	atttcaccct	cacaattaat	240
cctgtggaag	ctaatagatac	tgcaaattat	tactgtcacc	aaggtaagga	ggttccgtgg	300

acgttcggcc aagggacca ggtggaaatc aaa

333

<210> 1215

<211> 333

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1215

gaaattgtgt	tgacacagtc	tccagccacc	ctgtctttgt	ctccagggga	aagagccacc	60
ctctcctgca	gagccagcga	aagtgttggt	aattttggca	ttagttttgt	gaactgggtac	120
caacagaaac	ctggccaggc	tcccaggctc	ctcatctatg	ctgcctccaa	ccaaggatcc	180
ggcatccag	ccaggttcag	tggcagtggg	tctgggacag	acttcactct	caccatcagc	240
agcctagagc	ctgaagattt	tgcagtttat	tactgtcagc	aaagtaagga	ggttccgtac	300
acttttggcc	aggggacca	gctggagatc	aag			333

<210> 1216

<211> 334

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1216

gacatccaga	tgaccagtc	tccatcctcc	ctgtctgcat	ctgtaggaga	cagagtcacc	60
atcacttgca	gagccagcga	aagtgttggt	aattttggca	ttagttttgt	gaactgggtat	120
cagcagaaac	cagggaaagc	ccctaagctc	ctgatctatg	ctgcctccaa	ccaaggatcc	180
gggggtccat	caaggttcag	tggcagtggg	tctgggacag	atttcactct	caccatcagc	240
agctctgaac	ctgaagattt	tgcaacttac	tactgtcagc	aaagtaagga	ggttccgtac	300
acttttggcc	aggggacca	gctggagatc	aagt			334

<210> 1217

<211> 360

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1217

gaggtgcagc	ttgttgagtc	tgggtggagga	ttggtacagc	ctaaaggga	attgagactc	60
tcatgtgccg	cctctggatt	cagcttcaat	acctatgcc	tgaactgggt	ccgccaggct	120
ccaggaaaga	gtttggagtg	ggttgctcgc	acagtgacta	aaagtaataa	gtatgcaaca	180
tattatgcag	attcagtgcg	tgacagagtc	accatctcca	gagaggattc	acaaagcatg	240
ctttatctgc	aatgacca	cttgaaaact	gaggacacag	ccatgtatta	ctgtgtgaga	300
gaagattact	acggcactct	ggactactgg	ggtcaaggaa	cctcagtcac	cgtctcctca	360

<210> 1218

<211> 354

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1218  
caggtccaac tgcagcagcc tggggctgaa ctggtgaagc ctggggcttc tgtgaagctg 60  
tcctgcaagg cttctggcta catcttcacc agctactgga tgcactgggt gaagcagagg 120  
cctggacaag gccttgagtg gattggagag attaatccta gcaacggtcg tactaactac 180  
aatgagaggt tcaagagcaa ggccacactg actgtagaca aatcctccag cacagcctac 240  
attcaactca gcagcctgac atctgatgac tctgcggtct attactgttc aagatcggat 300  
actaaggcta cgtgtgacta ctggggccaa ggcaccactc tcacagtctc ctca 354

<210> 1219  
<211> 360  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1219  
gaggtgcagc tgggtggagtc tgggggagggc ttggtccagc ctggaggggc cctgagactc 60  
tcctgtgcag cctctggatt caccttcaat acctatgcca tgaactgggt ccgccaggct 120  
ccaggggaagg ggctggagtg ggttggccgc acagtgacta aaagtaataa gtatgcaaca 180  
tattatgcag attcagtgag tgacagattc accatctcaa gagatgattc aaagaactca 240  
ctgtatctgc aaatgaacag cctgaaaacc gaggacacgg ccgtgtatta ctgtgctaga 300  
gaagattact acggcactct ggactactgg ggccaaggaa ccctgggtcac cgtctcctca 360

<210> 1220  
<211> 360  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1220  
gaggtgcagc tgggtggagtc tgggggagggc ttggtccagc ctggaggggc cctgagactc 60  
tcctgtgcag cctctggatt caccttcaat acctatgcca tgaactgggt ccgccaggct 120  
ccaggggaagg ggctggagtg ggttggccgc acagtgacta aaagtaataa gtatgcaaca 180  
tattatgcag attcagtgag tgacagattc accatctcaa gagatgattc aaagaactca 240  
ctgtatctgc aaatgaacag cctgaaaacc gaggacacgg ccgtgtatta ctgtgctaga 300  
gaagattact acggcactct ggactactgg ggccaaggaa ccctgggtcac cgtctcctca 360

<210> 1221  
<211> 354  
<212> DNA  
<213> Artificial Sequence

<220>  
<223> Made in a lab

<400> 1221  
caggtgcagc tgggtgcagtc tggggctgag gtgaagaagc ctggggcctc agtgaaggtc 60  
tcctgcaagg cttctggcta catcttcacc agctactgga tgcactgggt gcgacaggcc 120  
cctggacaag ggcttgagtg gatgggagag attaatccta gcaacggtcg tactaactac 180  
aatgagaggt tcaagagcag ggtcaccatg accagggaca cgtccatcag cacagcctac 240  
atggagctga gcaggctgag atctgacgac acggccgtgt attactgtgc gagatcggat 300  
actaaggcta cgtgtgacta ctggggccaa gggaccacgg tcaccgtctc ctct 354

<210> 1222  
 <211> 354  
 <212> DNA  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1222  
 caggtgcagc tgggtgcagtc tggggctgag gtgaagaagc ctggggcctc agtgaaggtc 60  
 tcctgcaagg cttctggcta catcttcacc agctactgga tgcactgggt gcgacaggcc 120  
 cctggacaag ggcttgagtg gatgggagag attaatccta gcaacggtcg tactaactac 180  
 aatgagaggt tcaagagcag ggtcaccatg accagggaca cgtccatcag cacagcctac 240  
 atggagctga gcaggctgag atctgacgac acggccgtgt attactgtgc gagatcggat 300  
 actaaggcta cgtgtgacta ctggggccaa gggaccacgg tcaccgtctc ctct 354

<210> 1223  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1223  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Gly Thr Gln Met Ala Gly  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1224  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1224  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Gly Thr Gln Lys Ala Arg  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1225  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1225  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Gly Thr Gln Met Ala Arg  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1226  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1226  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Arg Thr Gln Lys Ala Gly  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1227  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1227  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Arg Thr Gln Met Ala Gly  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1228  
 <211> 21  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1228  
 Gln Arg His Asn Asn Ser Ser Leu Asn Thr Arg Thr Gln Met Ala Arg  
 1 5 10 15  
 His Ser Pro Asn Ser  
 20

<210> 1229  
 <211> 22  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in lab

<400> 1229  
 Met Asp Phe Gln Val Gln Ile Phe Ser Phe Leu Leu Ile Ser Ala Ser  
 1 5 10 15  
 Val Ile Met Ser Arg Gly  
 20



<210> 1230  
 <211> 723  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1230

Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	Ala	Pro	Glu	Pro	Gly	Ser
1				5					10					15	
Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	Thr	Ala	Gln	Met	Cys	Cys
			20					25					30		
Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	Val	Phe	Cys	Thr	Lys	Thr
		35					40					45			
Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	Ser	Thr	Tyr	Thr	Gln	Leu
	50					55					60				
Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	Gly	Ser	Arg	Cys	Ser	Ser
65					70					75					80
Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	Glu	Gln	Asn	Arg	Ile	Cys
				85					90					95	
Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	Ser	Lys	Gln	Glu	Gly	Cys
			100					105					110		
Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	Pro	Gly	Phe	Gly	Val	Ala
		115					120					125			
Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	Cys	Lys	Pro	Cys	Ala	Pro
	130					135					140				
Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	Asp	Ile	Cys	Arg	Pro	His
145					150					155					160
Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	Asn	Ala	Ser	Met	Asp	Ala
			165						170					175	
Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	Met	Ala	Pro	Gly	Ala	Val
			180					185					190		
His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	Gln	His	Thr	Gln	Pro	Thr
		195					200					205			
Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	Phe	Leu	Leu	Pro	Met	Gly
	210					215					220				
Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	Asp	Thr	Gly	Glu	Pro	Lys
225				230						235					240
Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Ala
			245						250					255	
Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr
			260				265						270		
Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val
		275					280						285		
Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val
	290					295					300				
Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser
305				310						315					320
Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu
			325						330					335	
Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala	Leu	Pro	Ala
			340				345						350		
Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro	Arg	Glu	Pro
		355					360					365			
Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr	Lys	Asn	Gln
	370					375					380				

Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala  
 385 390 395 400  
 Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr  
 405 410 415  
 Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu  
 420 425 430  
 Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser  
 435 440 445  
 Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser  
 450 455 460  
 Leu Ser Pro Gly Gln Arg His Asn Asn Ser Ser Leu Asn Thr Gly Thr  
 465 470 475 480  
 Gln Met Ala Gly His Ser Pro Asn Ser Ala Pro Trp Gln Cys Ala Pro  
 485 490 495  
 Cys Ser Ala Glu Lys Leu Ala Leu Cys Pro Pro Val Ser Ala Ser Cys  
 500 505 510  
 Ser Glu Val Thr Arg Ser Ala Gly Cys Gly Cys Cys Pro Met Cys Ala  
 515 520 525  
 Leu Pro Leu Gly Ala Ala Cys Gly Val Ala Thr Ala Arg Cys Ala Arg  
 530 535 540  
 Gly Leu Ser Cys Arg Ala Leu Pro Gly Glu Gln Gln Pro Leu His Ala  
 545 550 555 560  
 Leu Thr Arg Gly Gln Gly Ala Cys Val Gln Glu Ser Asp Ala Ser Ala  
 565 570 575  
 Pro His Ala Ala Glu Ala Gly Ser Pro Glu Ser Pro Glu Ser Thr Glu  
 580 585 590  
 Ile Thr Glu Glu Glu Leu Leu Asp Asn Phe His Leu Met Ala Pro Ser  
 595 600 605  
 Glu Glu Asp His Ser Ile Leu Trp Asp Ala Ile Ser Thr Tyr Asp Gly  
 610 615 620  
 Ser Lys Ala Leu His Val Thr Asn Ile Lys Lys Trp Lys Glu Pro Cys  
 625 630 635 640  
 Arg Ile Glu Leu Tyr Arg Val Val Glu Ser Leu Ala Lys Ala Gln Glu  
 645 650 655  
 Thr Ser Gly Glu Glu Ile Ser Lys Phe Tyr Leu Pro Asn Cys Asn Lys  
 660 665 670  
 Asn Gly Phe Tyr His Ser Arg Gln Cys Glu Thr Ser Met Asp Gly Glu  
 675 680 685  
 Ala Gly Leu Cys Trp Cys Val Tyr Pro Trp Asn Gly Lys Arg Ile Pro  
 690 695 700  
 Gly Ser Pro Glu Ile Arg Gly Asp Pro Asn Cys Gln Ile Tyr Phe Asn  
 705 710 715 720  
 Val Gln Asn

<210> 1231

<211> 2235

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1231

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cgaggattgc	ccgcccaggt	ggcatttaca	ccctacgccc	cggagcccg	gagcacatgc	120
cggctcagag	aatactatga	ccagacagct	cagatgtgct	gcagcaaata	ctcgccgggc	180

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caacatgcaa aagtcttctg taccaagacc tcggacaccg tgtgtgactc ctgtgaggac 240
agcacataca cccagctctg gaactgggtt cccgagtgtg tgagctgtgg ctcccgtgt 300
agctctgacc aggtggaaac tcaagcctgc actcgggaac agaaccgcat ctgcacctgc 360
aggcccggtt ggtactgctg gctgagcaag caggaggggt gccggctgtg cgcgccgtg 420
cgcaagtgcc gcccggtt cggcgtggcc agaccaggaa ctgaaacatc agacgtggtg 480
tgcaagccct gtgccccggg gacgttctcc aacacgactt catccacgga tatttgagg 540
ccccaccaga tctgtaacgt ggtggccatc cctgggaatg caagcatgga tgcagtctgc 600
acgtccacgt cccccaccg gagtatggcc ccaggggcag tacacttacc ccagccagt 660
tccacacgat cccaacacac gcagccaact ccagaacca gactgctcc aagcacctcc 720
ttctgtctcc caatgggccc cagcccccca gctgaaggga gactggcgga caccggtgag 780
cccaaattct ctgacaaaac tcacacatgc ccaccgtgcc cagcacctga agccgcgggt 840
gcaccgtcag tcttctctct cccccaaaa cccaaggaca ccctcatgat ctcccggacc 900
cctgaggtca catgctggtt ggtggacgtg agccacgaag accctgaggt caagttcaac 960
tggtacgtgg acggcgtgga ggtgcataat gccaaagaaa agccgcggga ggagcagtac 1020
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aaggcgtacg cgtgcgcggt ctccaacaaa gccctcccag ccccatcga gaaaaccatc 1140
tccaaagcca aagggcagcc ccgagaacca caggtgtaca ccctgcccc atcccgggat 1200
gagctgacca agaaccaggt cagcctgacc tgctgtgtca aaggcttcta tccaagcgac 1260
atcgccgtgg agtgggagag caatgggcag ccggagaaca actacaagac cagcctccc 1320
gtgctggact ccgacggctc tttcttctc tacagcaagc tcaccgtgga caagagcagg 1380
tggcagcagg ggaacgtctt ctcatgctcc gtgatgcatg aggtctctga caaccactac 1440
acgcagaaga gcctctccct gtctccgggt cagaggcaca acaattcttc cctgaatata 1500
ggaactcaga tggcaggtca ttctccgaat tctgctccgt ggcagtgcgc gccctgctcc 1560
gccgagaagc tcgcgctctg cccgccggtg tccgcctcgt gctcggaggt cacccggtcc 1620
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agacagtgtg agacatccat ggatggagag gcgggactct gctggtgcgt ctacccttgg 2160
aatgggaaga ggatccctgg gtctccagag atcaggggag accccaactg ccagatatat 2220
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<210> 1232

<211> 1156

<212> PRT

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1232

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Met Glu Ala Pro Ala Gln Leu Leu Phe Leu Leu Leu Leu Trp Leu Pro
 1           5           10           15
Asp Thr Thr Gly Glu Ile Cys Gly Pro Gly Ile Asp Ile Arg Asn Asp
      20           25           30
Tyr Gln Gln Leu Lys Arg Leu Glu Asn Cys Thr Val Ile Glu Gly Tyr
      35           40           45
Leu His Ile Leu Leu Ile Ser Lys Ala Glu Asp Tyr Arg Ser Tyr Arg
      50           55           60
Phe Pro Lys Leu Thr Val Ile Thr Glu Tyr Leu Leu Leu Phe Arg Val
      65           70           75           80
Ala Gly Leu Glu Ser Leu Gly Asp Leu Phe Pro Asn Leu Thr Val Ile
      85           90           95

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Arg	Gly	Trp	Lys	Leu	Phe	Tyr	Asn	Tyr	Ala	Leu	Val	Ile	Phe	Glu	Met
			100					105					110		
Thr	Asn	Leu	Lys	Asp	Ile	Gly	Leu	Tyr	Asn	Leu	Arg	Asn	Ile	Thr	Arg
		115					120					125			
Gly	Ala	Ile	Arg	Ile	Glu	Lys	Asn	Ala	Asp	Leu	Cys	Tyr	Leu	Ser	Thr
	130					135					140				
Val	Asp	Trp	Ser	Leu	Ile	Leu	Asp	Ala	Val	Ser	Asn	Asn	Tyr	Ile	Val
145					150					155					160
Gly	Asn	Lys	Pro	Pro	Lys	Glu	Cys	Gly	Asp	Leu	Cys	Pro	Gly	Thr	Met
			165						170					175	
Glu	Glu	Lys	Pro	Met	Cys	Glu	Lys	Thr	Thr	Ile	Asn	Asn	Glu	Tyr	Asn
			180					185					190		
Tyr	Arg	Cys	Trp	Thr	Thr	Asn	Arg	Cys	Gln	Lys	Met	Cys	Pro	Ser	Thr
		195					200					205			
Cys	Gly	Lys	Arg	Ala	Cys	Thr	Glu	Asn	Asn	Glu	Cys	Cys	His	Pro	Glu
	210					215					220				
Cys	Leu	Gly	Ser	Cys	Ser	Ala	Pro	Asp	Asn	Asp	Thr	Ala	Cys	Val	Ala
225					230					235					240
Cys	Arg	His	Tyr	Tyr	Tyr	Ala	Gly	Val	Cys	Val	Pro	Ala	Cys	Pro	Pro
			245					250					255		
Asn	Thr	Tyr	Arg	Phe	Glu	Gly	Trp	Arg	Cys	Val	Asp	Arg	Asp	Phe	Cys
			260					265				270			
Ala	Asn	Ile	Leu	Ser	Ala	Glu	Ser	Ser	Asp	Ser	Glu	Gly	Phe	Val	Ile
		275				280						285			
His	Asp	Gly	Glu	Cys	Met	Gln	Glu	Cys	Pro	Ser	Gly	Phe	Ile	Arg	Asn
	290					295					300				
Gly	Ser	Gln	Ser	Met	Tyr	Cys	Ile	Pro	Cys	Glu	Gly	Pro	Cys	Pro	Lys
305					310					315					320
Val	Cys	Glu	Glu	Glu	Lys	Lys	Thr	Lys	Thr	Ile	Asp	Ser	Val	Thr	Ser
			325						330					335	
Ala	Gln	Met	Leu	Gln	Gly	Cys	Thr	Ile	Phe	Lys	Gly	Asn	Leu	Leu	Ile
			340					345					350		
Asn	Ile	Arg	Arg	Gly	Asn	Asn	Ile	Ala	Ser	Glu	Leu	Glu	Asn	Phe	Met
		355					360					365			
Gly	Leu	Ile	Glu	Val	Val	Thr	Gly	Tyr	Val	Lys	Ile	Arg	His	Ser	His
	370					375					380				
Ala	Leu	Val	Ser	Leu	Ser	Phe	Leu	Lys	Asn	Leu	Arg	Leu	Ile	Leu	Gly
385					390					395					400
Glu	Glu	Gln	Leu	Glu	Gly	Asn	Tyr	Ser	Phe	Tyr	Val	Leu	Asp	Asn	Gln
			405						410				415		
Asn	Leu	Gln	Gln	Leu	Trp	Asp	Trp	Asp	His	Arg	Asn	Leu	Thr	Ile	Lys
			420					425					430		
Ala	Gly	Lys	Met	Tyr	Phe	Ala	Phe	Asn	Pro	Lys	Leu	Cys	Val	Ser	Glu
		435					440					445			
Ile	Tyr	Arg	Met	Glu	Glu	Val	Thr	Gly	Thr	Lys	Gly	Arg	Gln	Ser	Lys
	450					455					460				
Gly	Asp	Ile	Asn	Thr	Arg	Asn	Asn	Gly	Glu	Arg	Ala	Ser	Cys	Glu	Ser
465					470					475					480
Asp	Val	Leu	His	Phe	Thr	Ser	Thr	Thr	Thr	Ser	Lys	Asn	Arg	Ile	Ile
			485						490					495	
Ile	Thr	Trp	His	Arg	Tyr	Arg	Pro	Pro	Asp	Tyr	Arg	Asp	Leu	Ile	Ser
			500					505					510		
Phe	Thr	Val	Tyr	Tyr	Lys	Glu	Ala	Pro	Phe	Lys	Asn	Val	Thr	Glu	Tyr
		515					520					525			
Asp	Gly	Gln	Asp	Ala	Cys	Gly	Ser	Asn	Ser	Trp	Asn	Met	Val	Asp	Val
	530					535					540				
Asp	Leu	Pro	Pro	Asn	Lys	Asp	Val	Glu	Pro	Gly	Ile	Leu	Leu	His	Gly

545					550					555				560	
Leu	Lys	Pro	Trp	Thr	Gln	Tyr	Ala	Val	Tyr	Val	Lys	Ala	Val	Thr	Leu
				565					570					575	
Thr	Met	Val	Glu	Asn	Asp	His	Ile	Arg	Gly	Ala	Lys	Ser	Glu	Ile	Leu
			580					585					590		
Tyr	Ile	Arg	Thr	Asn	Ala	Ser	Val	Pro	Ser	Ile	Pro	Leu	Asp	Val	Leu
		595					600					605			
Ser	Ala	Ser	Asn	Ser	Ser	Ser	Gln	Leu	Ile	Val	Lys	Trp	Asn	Pro	Pro
	610						615				620				
Ser	Leu	Pro	Asn	Gly	Asn	Leu	Ser	Tyr	Tyr	Ile	Val	Arg	Trp	Gln	Arg
625					630						635				640
Gln	Pro	Gln	Asp	Gly	Tyr	Leu	Tyr	Arg	His	Asn	Tyr	Cys	Ser	Lys	Asp
				645					650					655	
Lys	Ile	Pro	Ile	Arg	Lys	Tyr	Ala	Asp	Gly	Thr	Ile	Asp	Ile	Glu	Glu
			660					665					670		
Val	Thr	Glu	Asn	Pro	Lys	Thr	Glu	Val	Cys	Gly	Gly	Glu	Lys	Gly	Pro
		675					680					685			
Cys	Cys	Ala	Cys	Pro	Lys	Thr	Glu	Ala	Glu	Lys	Gln	Ala	Glu	Lys	Glu
	690					695					700				
Glu	Ala	Glu	Tyr	Arg	Lys	Val	Phe	Glu	Asn	Phe	Leu	His	Asn	Ser	Ile
705					710					715					720
Phe	Val	Pro	Arg	Pro	Glu	Arg	Lys	Arg	Arg	Asp	Val	Met	Gln	Val	Ala
				725					730					735	
Asn	Thr	Thr	Met	Ser	Ser	Arg	Ser	Arg	Asn	Thr	Thr	Ala	Ala	Asp	Thr
			740					745					750		
Tyr	Asn	Ile	Thr	Asp	Pro	Glu	Glu	Leu	Glu	Thr	Glu	Tyr	Pro	Phe	Phe
	755						760					765			
Glu	Ser	Arg	Val	Asp	Asn	Lys	Glu	Arg	Thr	Val	Ile	Ser	Asn	Leu	Arg
	770					775					780				
Pro	Phe	Thr	Leu	Tyr	Arg	Ile	Asp	Ile	His	Ser	Cys	Asn	His	Glu	Ala
785					790					795					800
Glu	Lys	Leu	Gly	Cys	Ser	Ala	Ser	Asn	Phe	Val	Phe	Ala	Arg	Thr	Met
				805					810					815	
Pro	Ala	Glu	Gly	Ala	Asp	Asp	Ile	Pro	Gly	Pro	Val	Thr	Trp	Glu	Pro
			820					825					830		
Arg	Pro	Glu	Asn	Ser	Ile	Phe	Leu	Lys	Trp	Pro	Glu	Pro	Glu	Asn	Pro
		835					840					845			
Asn	Gly	Leu	Ile	Leu	Met	Tyr	Glu	Ile	Lys	Tyr	Gly	Ser	Gln	Val	Glu
	850					855					860				
Asp	Gln	Arg	Glu	Cys	Val	Ser	Arg	Gln	Glu	Tyr	Arg	Lys	Tyr	Gly	Gly
865					870					875					880
Ala	Lys	Leu	Asn	Arg	Leu	Asn	Pro	Gly	Asn	Tyr	Thr	Ala	Arg	Ile	Gln
			885						890					895	
Ala	Thr	Ser	Leu	Ser	Gly	Asn	Gly	Ser	Trp	Thr	Asp	Pro	Val	Phe	Phe
		900						905					910		
Tyr	Val	Gln	Ala	Lys	Thr	Gly	Tyr	Glu	Asn	Ser	Ser	Glu	Pro	Lys	Ser
		915					920					925			
Ser	Asp	Lys	Thr	His	Thr	Ser	Pro	Pro	Ser	Ser	Ala	Pro	Glu	Leu	Leu
	930					935					940				
Gly	Gly	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr	Leu
945					950					955					960
Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val	Ser
				965					970					975	
His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val	Glu
			980					985					990		
Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser	Thr
		995					1000					1005			

Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn  
 1010 1015 1020  
 Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro  
 1025 1030 1035 1040  
 Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln  
 1045 1050 1055  
 Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu Leu Thr Lys Asn Gln Val  
 1060 1065 1070  
 Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val  
 1075 1080 1085  
 Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro  
 1090 1095 1100  
 Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr  
 1105 1110 1115 1120  
 Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val  
 1125 1130 1135  
 Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu  
 1140 1145 1150  
 Ser Pro Gly Lys  
 1155

<210> 1233  
 <211> 961  
 <212> PRT  
 <213> Artificial Sequence

<220>  
 <223> Made in a lab

<400> 1233  
 Met Glu Ala Pro Ala Gln Leu Leu Phe Leu Leu Leu Leu Trp Leu Pro  
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 Asp Thr Thr Gly Glu Ile Cys Gly Pro Gly Ile Asp Ile Arg Asn Asp  
 20 25 30  
 Tyr Gln Gln Leu Lys Arg Leu Glu Asn Cys Thr Val Ile Glu Gly Tyr  
 35 40 45  
 Leu His Ile Leu Leu Ile Ser Lys Ala Glu Asp Tyr Arg Ser Tyr Arg  
 50 55 60  
 Phe Pro Lys Leu Thr Val Ile Thr Glu Tyr Leu Leu Leu Phe Arg Val  
 65 70 75 80  
 Ala Gly Leu Glu Ser Leu Gly Asp Leu Phe Pro Asn Leu Thr Val Ile  
 85 90 95  
 Arg Gly Trp Lys Leu Phe Tyr Asn Tyr Ala Leu Val Ile Phe Glu Met  
 100 105 110  
 Thr Asn Leu Lys Asp Ile Gly Leu Tyr Asn Leu Arg Asn Ile Thr Arg  
 115 120 125  
 Gly Ala Ile Arg Ile Glu Lys Asn Ala Asp Leu Cys Tyr Leu Ser Thr  
 130 135 140  
 Val Asp Trp Ser Leu Ile Leu Asp Ala Val Ser Asn Asn Tyr Ile Val  
 145 150 155 160  
 Gly Asn Lys Pro Pro Lys Glu Cys Gly Asp Leu Cys Pro Gly Thr Met  
 165 170 175  
 Glu Glu Lys Pro Met Cys Glu Lys Thr Thr Ile Asn Asn Glu Tyr Asn  
 180 185 190  
 Tyr Arg Cys Trp Thr Thr Asn Arg Cys Gln Lys Met Cys Pro Ser Thr  
 195 200 205  
 Cys Gly Lys Arg Ala Cys Thr Glu Asn Asn Glu Cys Cys His Pro Glu

210		215		220
Cys Leu Gly Ser Cys Ser	Ala Pro Asp Asn Asp	Thr Ala Cys Val Ala		
225	230	235	240	
Cys Arg His Tyr Tyr Tyr	Ala Gly Val Cys Val	Pro Ala Cys Pro Pro		
	245	250	255	
Asn Thr Tyr Arg Phe Glu Gly Trp Arg	Cys Val Asp Arg Asp Phe Cys			
	260	265	270	
Ala Asn Ile Leu Ser Ala Glu Ser Ser Asp Ser Glu Gly Phe Val Ile				
	275	280	285	
His Asp Gly Glu Cys Met Gln Glu Cys Pro Ser Gly Phe Ile Arg Asn				
	290	295	300	
Gly Ser Gln Ser Met Tyr Cys Ile Pro Cys Glu Gly Pro Cys Pro Lys				
305	310	315	320	
Val Cys Glu Glu Glu Lys Lys Thr Lys Thr Ile Asp Ser Val Thr Ser				
	325	330	335	
Ala Gln Met Leu Gln Gly Cys Thr Ile Phe Lys Gly Asn Leu Leu Ile				
	340	345	350	
Asn Ile Arg Arg Gly Asn Asn Ile Ala Ser Glu Leu Glu Asn Phe Met				
	355	360	365	
Gly Leu Ile Glu Val Val Thr Gly Tyr Val Lys Ile Arg His Ser His				
	370	375	380	
Ala Leu Val Ser Leu Ser Phe Leu Lys Asn Leu Arg Leu Ile Leu Gly				
385	390	395	400	
Glu Glu Gln Leu Glu Gly Asn Tyr Ser Phe Tyr Val Leu Asp Asn Gln				
	405	410	415	
Asn Leu Gln Gln Leu Trp Asp Trp Asp His Arg Asn Leu Thr Ile Lys				
	420	425	430	
Ala Gly Lys Met Tyr Phe Ala Phe Asn Pro Lys Leu Cys Val Ser Glu				
	435	440	445	
Ile Tyr Arg Met Glu Glu Val Thr Gly Thr Lys Gly Arg Gln Ser Lys				
	450	455	460	
Gly Asp Ile Asn Thr Arg Asn Asn Gly Glu Arg Ala Ser Cys Glu Ser				
465	470	475	480	
Asp Val Leu His Phe Thr Ser Thr Thr Thr Ser Lys Asn Arg Ile Ile				
	485	490	495	
Ile Thr Trp His Arg Tyr Arg Pro Pro Asp Tyr Arg Asp Leu Ile Ser				
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Phe Thr Val Tyr Tyr Lys Glu Ala Pro Phe Lys Asn Val Thr Glu Tyr				
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Asp Gly Gln Asp Ala Cys Gly Ser Asn Ser Trp Asn Met Val Asp Val				
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Asp Leu Pro Pro Asn Lys Asp Val Glu Pro Gly Ile Leu Leu His Gly				
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Leu Lys Pro Trp Thr Gln Tyr Ala Val Tyr Val Lys Ala Val Thr Leu				
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Thr Met Val Glu Asn Asp His Ile Arg Gly Ala Lys Ser Glu Ile Leu				
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Tyr Ile Arg Thr Asn Ala Ser Val Pro Ser Ile Pro Leu Asp Val Leu				
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Ser Ala Ser Asn Ser Ser Ser Gln Leu Ile Val Lys Trp Asn Pro Pro				
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Ser Leu Pro Asn Gly Asn Leu Ser Tyr Tyr Ile Val Arg Trp Gln Arg				
625	630	635	640	
Gln Pro Gln Asp Gly Tyr Leu Tyr Arg His Asn Tyr Cys Ser Lys Asp				
	645	650	655	
Lys Ile Pro Ile Arg Lys Tyr Ala Asp Gly Thr Ile Asp Ile Glu Glu				
	660	665	670	

Val Thr Glu Asn Pro Lys Thr Glu Val Cys Gly Gly Glu Lys Gly Pro  
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 Cys Cys Ala Cys Pro Lys Thr Glu Ala Glu Lys Gln Ala Glu Lys Glu  
 690 695 700  
 Glu Ala Glu Tyr Arg Lys Val Phe Glu Asn Phe Leu His Asn Ser Ile  
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 Phe Val Pro Arg Pro Glu Ser Asp Gln Glu Pro Lys Ser Cys Asp Lys  
 725 730 735  
 Thr His Thr Ser Pro Pro Cys Ser Ala Pro Glu Leu Leu Gly Gly Pro  
 740 745 750  
 Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser  
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 Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu Asp  
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 Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His Asn  
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 Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val  
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 Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu  
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 Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys  
 835 840 845  
 Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr  
 850 855 860  
 Leu Pro Pro Ser Arg Asp Glu Leu Thr Lys Asn Gln Val Ser Leu Thr  
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 Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu  
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 Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu  
 900 905 910  
 Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys  
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 Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu  
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 Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly  
 945 950 955 960  
 Lys

<210> 1234

<211> 3468

<212> DNA

<213> Artificial Sequence

<220>

<223> Made in a lab

<400> 1234

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gggaataagc	cccaaagga	atgtggggac	ctgtgtccag	ggacctgga	ggagaagccg	540



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&lt;210&gt; 1235

&lt;211&gt; 2883

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1235

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aaa						2883

&lt;210&gt; 1236

&lt;211&gt; 633

&lt;212&gt; PRT

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1236

Leu	Pro	Ala	Gln	Val	Ala	Phe	Thr	Pro	Tyr	Ala	Pro	Glu	Pro	Gly	Ser
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Thr	Cys	Arg	Leu	Arg	Glu	Tyr	Tyr	Asp	Gln	Thr	Ala	Gln	Met	Cys	Cys
			20					25					30		
Ser	Lys	Cys	Ser	Pro	Gly	Gln	His	Ala	Lys	Val	Phe	Cys	Thr	Lys	Thr
		35					40					45			
Ser	Asp	Thr	Val	Cys	Asp	Ser	Cys	Glu	Asp	Ser	Thr	Tyr	Thr	Gln	Leu
	50					55					60				
Trp	Asn	Trp	Val	Pro	Glu	Cys	Leu	Ser	Cys	Gly	Ser	Arg	Cys	Ser	Ser
65					70					75					80
Asp	Gln	Val	Glu	Thr	Gln	Ala	Cys	Thr	Arg	Glu	Gln	Asn	Arg	Ile	Cys
				85					90					95	
Thr	Cys	Arg	Pro	Gly	Trp	Tyr	Cys	Ala	Leu	Ser	Lys	Gln	Glu	Gly	Cys
			100					105					110		
Arg	Leu	Cys	Ala	Pro	Leu	Arg	Lys	Cys	Arg	Pro	Gly	Phe	Gly	Val	Ala
		115					120					125			
Arg	Pro	Gly	Thr	Glu	Thr	Ser	Asp	Val	Val	Cys	Lys	Pro	Cys	Ala	Pro
	130					135					140				
Gly	Thr	Phe	Ser	Asn	Thr	Thr	Ser	Ser	Thr	Asp	Ile	Cys	Arg	Pro	His
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Gln	Ile	Cys	Asn	Val	Val	Ala	Ile	Pro	Gly	Asn	Ala	Ser	Met	Asp	Ala
			165						170					175	
Val	Cys	Thr	Ser	Thr	Ser	Pro	Thr	Arg	Ser	Met	Ala	Pro	Gly	Ala	Val
			180					185					190		
His	Leu	Pro	Gln	Pro	Val	Ser	Thr	Arg	Ser	Gln	His	Thr	Gln	Pro	Thr
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Pro	Glu	Pro	Ser	Thr	Ala	Pro	Ser	Thr	Ser	Phe	Leu	Leu	Pro	Met	Gly
	210					215					220				
Pro	Ser	Pro	Pro	Ala	Glu	Gly	Ser	Thr	Gly	Asp	Thr	Gly	Glu	Pro	Lys
225					230					235					240
Ser	Ser	Asp	Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Ala
			245						250					255	
Ala	Gly	Ala	Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr
			260					265					270		
Leu	Met	Ile	Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val
		275					280					285			
Ser	His	Glu	Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val
	290					295					300				
Glu	Val	His	Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser
305					310					315					320
Thr	Tyr	Arg	Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu
			325						330					335	
Asn	Gly	Lys	Ala	Tyr	Ala	Cys	Ala	Val	Ser	Asn	Lys	Ala	Leu	Pro	Ala
			340					345					350		
Pro	Ile	Glu	Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro	Arg	Glu	Pro
		355					360					365			
Gln	Val	Tyr	Thr	Leu	Pro	Pro	Ser	Arg	Asp	Glu	Leu	Thr	Lys	Asn	Gln
	370					375					380				
Val	Ser	Leu	Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser	Asp	Ile	Ala
385					390					395					400
Val	Glu	Trp	Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr	Lys	Thr	Thr
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[illegible]

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<210> 1237
<211> 450
<212> PRT
<213> Artificial Sequence
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<220>  
<223> Made in a lab

<400> 1237															
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Ala	Asp	Leu	Asp	Lys	Cys	Met	Asp	Cys	Ala	Ser	Cys	Arg	Ala	Arg	Pro
			20					25					30		
His	Ser	Asp	Phe	Cys	Leu	Gly	Cys	Ala	Ala	Ala	Pro	Pro	Ala	Pro	Phe
		35					40					45			
Arg	Leu	Leu	Trp	Thr	Gly	Glu	Pro	Lys	Ser	Ser	Asp	Lys	Thr	His	Thr
	50					55					60				
Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Ala	Ala	Gly	Ala	Pro	Ser	Val	Phe
65					70					75				80	
Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile	Ser	Arg	Thr	Pro
				85					90					95	
Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val	Ser	His	Glu	Asp	Pro	Glu	Val
			100					105					110		
Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His	Asn	Ala	Lys	Thr
		115					120					125			
Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg	Val	Val	Ser	Val
		130				135					140				
Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys	Ala	Tyr	Ala	Cys
145					150					155					160

Ala Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser  
165 170 175  
Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro  
180 185 190  
Ser Arg Asp Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val  
195 200 205  
Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly  
210 215 220  
Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp  
225 230 235 240  
Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp  
245 250 255  
Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His  
260 265 270  
Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Gln Arg His  
275 280 285  
Asn Asn Ser Ser Leu Asn Thr Gly Thr Gln Met Ala Gly His Ser Pro  
290 295 300  
Asn Ser Thr Ile Pro Pro His Val Gln Lys Ser Val Asn Asn Asp Met  
305 310 315 320  
Ile Val Thr Asp Asn Asn Gly Ala Val Lys Phe Pro Gln Leu Cys Lys  
325 330 335  
Phe Cys Asp Val Arg Phe Ser Thr Cys Asp Asn Gln Lys Ser Cys Met  
340 345 350  
Ser Asn Cys Ser Ile Thr Ser Ile Cys Glu Lys Pro Gln Glu Val Cys  
355 360 365  
Val Ala Val Trp Arg Lys Asn Asp Glu Asn Ile Thr Leu Glu Thr Val  
370 375 380  
Cys His Asp Pro Lys Leu Pro Tyr His Asp Phe Ile Leu Glu Asp Ala  
385 390 395 400  
Ala Ser Pro Lys Cys Ile Met Lys Glu Lys Lys Lys Pro Gly Glu Thr  
405 410 415  
Phe Phe Met Cys Ser Cys Ser Ser Asp Glu Cys Asn Asp Asn Ile Ile  
420 425 430  
Phe Ser Glu Glu Tyr Asn Thr Ser Asn Pro Asp Leu Leu Leu Val Ile  
435 440 445  
Phe Gln  
450

&lt;210&gt; 1238

&lt;211&gt; 1968

&lt;212&gt; DNA

&lt;213&gt; Artificial Sequence

&lt;220&gt;

&lt;223&gt; Made in a lab

&lt;400&gt; 1238

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