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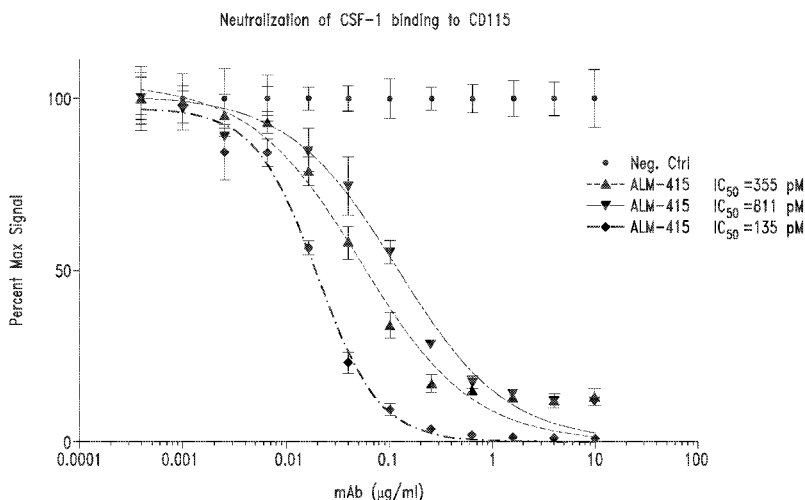


FIG. 10

(57) **Abstract:** The present invention provides anti-CD115 monoclonal antibodies and related compositions, which may be used in any of a variety of therapeutic and diagnostic methods for the treatment of cancer, autoimmune, and other diseases.

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ANTI-CD115 ANTIBODIES

CROSS REFERENCE TO RELATED APPLICATIONS

[0001] This application claims the benefit under 35 U.S.C. § 119(e) of U.S. Provisional Application No. 62/220,147, filed September 17, 2015 and U.S. Provisional Application No. 62/219,578, filed September 16, 2015, each of which is incorporated by reference herein in its entirety.

SEQUENCE LISTING

[0002] The Sequence Listing associated with this application is provided in text format in lieu of a paper copy, and is hereby incorporated by reference into the specification. The name of the text file containing the Sequence Listing is ABLX_007_02WO_ST25.txt. The text file is 525 KB, was created on September 16, 2016, and is being submitted electronically via EFS-Web.

BACKGROUND

Technical Field

[0003] The present invention relates generally to anti-CD115 antibodies, compositions and methods of using same. Such antibodies are useful, for example, for treating a variety of diseases, such as oncological and immunological diseases.

Description of the Related Art

[0004] CD115, also known as Colony-Stimulating Factor 1 Receptor (CSF1R) and macrophage-colony stimulating factor receptor (M-CSFR), is a cell surface receptor tyrosine kinase belonging to the platelet-derived growth factor family. CD115 has two structurally unrelated ligands, namely CSF-1 (M-CSF) and IL-34. CD115 is expressed by hematopoietic stem cells, myeloid cells, including monocytes, macrophages, osteoclasts, dendritic cells, and microglia, neural progenitor cells, and epithelial cells, including Paneth cells (Stanley and Chitu, Cold Spring Harb Perspect Biol 2014;6:a021857).

[0005] Dysregulation of CD115, and/or its ligands, is associated with proliferative diseases and disorders (e.g., neoplasms, tumors and metastases), as well as immunological and neurological diseases and disorders. The present invention provides chimeric and fully human anti-CD115 antibodies, including CD115 antagonists.

SUMMARY OF THE INVENTION

[0006] The present invention relates to anti-CD115 antibodies. More specifically, it relates to chimeric anti-CD115 antibodies generated from an AlivamaB Mouse, fully human anti-CD115 antibodies produced therefrom, and methods of use thereof.

[0007] One aspect of the invention provides an isolated anti-CD115 antibody, or an antigen-binding fragment thereof, comprising i) a heavy chain variable region comprising a VHCDR1 selected from any of SEQ ID NOs:436-543, a VHCDR2 selected from any of SEQ ID NOs:868-975, and a VHCDR3 selected from any of SEQ ID NOs:1300-1407 and ii) a light chain variable region comprising a VLCDR1 selected from any of SEQ ID NOs:652-759, a VLCDR2 selected from any of SEQ ID NOs:1084-1191, and a VLCDR3 selected from any of SEQ ID NOs:1516-1623.

[0008] In one embodiment, the VHCDR1, VHCDR2, and VHCDR3 of the anti-CD115 antibody, or antigen-binding fragment thereof, comprise SEQ ID NOs:450, 882, and 1314, respectively. In one embodiment, the VLCDR1, VLCDR2, and VLCDR3 comprise SEQ ID NOs:666, 1098, and 1530, respectively. In another embodiment, the VH is selected from any one of SEQ ID NOs:109-216. In yet another embodiment, the VL is selected from any one of SEQ ID NOs:325-432. In one embodiment, the VH comprises SEQ ID NO:123. In another embodiment, the VL comprises SEQ ID NO:339. In another embodiment, the VH comprises SEQ ID NO:123, and the VL comprises SEQ ID NO:339.

[0009] In one embodiment, the anti-CD115 antibody, or antigen-binding fragment thereof, is human. In one embodiment, the antibody is chimeric. In certain embodiments, the antibody is selected from a single-variable domain antibody, single chain antibody, a scFv, a bispecific antibody, a multi-specific antibody, a Fab, a F(ab')₂, and a whole antibody.

[0010] One aspect of the invention provides a recombinant polynucleotide encoding the anti-CD115 antibody, or antigen-binding fragment thereof, described above. Another aspect of the invention provides an expression vector comprising the recombinant polynucleotide. In another aspect of the invention provides an isolated host cell that comprises the expression vector. One aspect of the invention provides a composition comprising an anti-CD115 antibody, or antigen-binding fragment thereof, described herein and a physiologically acceptable carrier.

BRIEF DESCRIPTION OF THE FIGURES

[0011] Figure 1 shows ELISA can detect an increase in p-MCSFR. For each grouping of bars, the lysate ratio is neat, 1:2, 1:4, 1:8, 1:16 and 1:32 from left to right.

- [0012] Figure 2 shows Bin1 α -CD115 mAbs may block M-CSF and IL34 induced CD115 phosphorylation and Bin 3 35A may block all phosphorylation.
- [0013] Figure 3 shows CD115 phosphorylation (p-CD115 or p-MCSFR) measured by ELISA.
- [0014] Figures 4A and 4B show inhibition of m-CSF- and IL-34-induced phosphorylation of CD115 (MCSFR).
- [0015] Figure 5 shows binding of anti-CD115 IgG κ mAbs to CD115 expressed on OCI-AML5 cells.
- [0016] Figure 6 shows an assay for detecting inhibition of m-CSF (CSF-1) induced phosphorylation on CD115 expressing AML5 cells by anti-CD115 IgG κ mAb.
- [0017] Figure 7 shows binding of anti-CD115 IgG λ mAbs to OCI-AML5 cells.
- [0018] Figure 8 shows anti-CD115 IgG λ mAb inhibition of CSF-1 induced phosphorylation of CD115.
- [0019] Figure 9 shows the IC₅₀ of selected anti-CD115 mAbs. For each grouping of bars, no M-CSF, 10, 2, 0.4, 0.08, 0.016, 0.0032, and 0.00064 ng/ml are shown from left to right.
- [0020] Figure 10 shows neutralization of CSF-1 binding to CD115 by anti-CD115 mAbs.
- [0021] Figure 11 shows neutralization of CSF-1 induced phosphorylation of CD115 by anti-CD115 mAbs.
- [0022] Figure 12 shows internalization of anti-CD115 mAbs. From left to right, examples within the panel of anti-CD115 mAbs exhibiting no internalization, weak internalization, mid internalization, strong internalization and very strong internalization are depicted.
- [0023] Figure 13 shows conversion of anti-CD115 mAb ALM-423 to fully human antibodies.
- [0024] Figure 14 shows neutralization of CSF-1 binding to CD115 by fully human anti-CD115 mAbs.
- [0025] Figure 15 shows neutralization of pTyr formation on CD115 by fully human anti-CD115 mAbs.
- [0026] Figure 16 shows some anti-CD115 antibodies that are antagonists of CSF-1 induce p-tyr formation on CD115.

BRIEF DESCRIPTION OF THE SEQUENCES

- [0027] SEQ ID NOs:1-108 are polynucleotide sequences encoding VH regions of the anti-CD115 antibodies listed in Table 2.
- [0028] SEQ ID NOs:109-216 are amino acid sequences of VH regions of the anti-CD115 antibodies listed in Table 2.
- [0029] SEQ ID NOs:217-324 are polynucleotide sequences encoding VL regions of the anti-CD115 antibodies listed in Table 2.
- [0030] SEQ ID NOs:325-432 are amino acid sequences of VL regions of the anti-CD115 antibodies listed in Table 2.
- [0031] SEQ ID NO:433 is an IgG specific primer.
- [0032] SEQ ID NO:434 is an Ig λ specific primer.
- [0033] SEQ ID NO:435 is an Ig κ specific primer.
- [0034] SEQ ID NOs:436-543 are amino acid sequences of the VHCDR1 of the anti-CD115 antibodies listed in Table 2.
- [0035] SEQ ID NOs:544-651 are polynucleotide sequences encoding the VHCDR1 of the anti-CD115 antibodies listed in Table 2.
- [0036] SEQ ID NOs:652-759 are amino acid sequences of the VLCDR1 of the anti-CD115 antibodies listed in Table 2.
- [0037] SEQ ID NOs:760-867 are polynucleotide sequences encoding the VLCDR1 of the anti-CD115 antibodies listed in Table 2.
- [0038] SEQ ID NOs:868-975 are amino acid sequences of the VHCDR2 of the anti-CD115 antibodies listed in Table 2.
- [0039] SEQ ID NOs:976-1083 are polynucleotide sequences encoding the VHCDR2 of the anti-CD115 antibodies listed in Table 2.
- [0040] SEQ ID NOs:1084-1191 are amino acid sequences of the VLCDR2 of the anti-CD115 antibodies listed in Table 2.
- [0041] SEQ ID NOs:1192-1299 are polynucleotide sequences encoding the VLCDR2 of the anti-CD115 antibodies listed in Table 2.
- [0042] SEQ ID NOs:1300-1407 are amino acid sequences of the VHCDR3 of the anti-CD115 antibodies listed in Table 2.
- [0043] SEQ ID NOs:1408-1515 are polynucleotide sequences encoding the VHCDR3 of the anti-CD115 antibodies listed in Table 2.

[0044] SEQ ID NOs:1516-1623 are amino acid sequences of the VLCDR3 of the anti-CD115 antibodies listed in Table 2.

[0045] SEQ ID NOs:1624-1731 are polynucleotide sequences encoding the VLCDR3 of the anti-CD115 antibodies listed in Table 2.

DETAILED DESCRIPTION

[0046] The present disclosure relates to anti-CD115 antibodies. Ablexis has used its proprietary AlivaMab Mouse technology (*See* WO 2010/039900 and WO 2011/123708, incorporated herein in their entirety) to generate panels of monoclonal antibodies (mAbs) against human CD115. Antibodies that potently neutralize CD115 signaling induced by CSF-1 were identified within the panel of CD115 AlivaMab antibodies. In one embodiment, anti-CD115 AlivaMab antibodies potently neutralize CD115 signaling induced by IL-34. In one embodiment, anti-CD115 AlivaMab antibodies that potently neutralize CD115 signaling induced by both CSF-1 and IL-34. CD115 (colony-stimulating factor 1 receptor, CSF1R, C-FMS) is a member of the receptor tyrosine kinase superfamily. For a review of CD115 biology, refer to Stanley and Chitu, *Cold Spring Harb. Perspect. Biol.* 2014 Jun 2;6(6).

[0047] Embodiments of the invention pertain to the use of anti-CD115 antibodies, or antigen-binding fragments thereof, for the diagnosis, assessment and treatment of diseases and disorders associated with CD115, CSF-1 and/or IL-34 or aberrant expression thereof. The subject antibodies are used in the treatment or prevention of neoplasms and/or the treatment or prevention of autoimmune and/or inflammatory diseases, among other diseases.

[0048] Portions of variable regions from the AlivaMab antibodies may include all or a combination of the complementarity determining regions (CDRs) of the VH and/or VL. The variable regions may be formatted with constant regions, either native or modified for various desired effector functions, in a standard antibody structure (two heavy chains with two light chains). The variable regions may also be formatted as multi-specific antibodies, e.g., bispecific antibodies binding to two different epitopes on CD115 or to two different antigens, one of which is CD115. The variable regions may also be formatted as antibody fragments, e.g., single-domain antibodies comprising a single VH or VL, Fabs or Fab'2. The antibodies may also be used as antibody-drug conjugates, or carry other additions such as small molecule toxins, biologic toxins, cytokines, oligopeptides, or RNAs to increase therapeutic modality and/or increase safety.

[0049] The practice of the present invention will employ, unless indicated specifically to the contrary, conventional methods of virology, immunology, microbiology, molecular

biology and recombinant DNA techniques within the skill of the art, many of which are described below for the purpose of illustration. Such techniques are explained fully in the literature. See, e.g., *Current Protocols in Molecular Biology* or *Current Protocols in Immunology*, John Wiley & Sons, New York, N.Y.(2009); Ausubel et al., *Short Protocols in Molecular Biology*, 3rd ed., Wiley & Sons, 1995; Sambrook and Russell, *Molecular Cloning: A Laboratory Manual* (3rd Edition, 2001); Maniatis et al., *Molecular Cloning: A Laboratory Manual* (1982); *DNA Cloning: A Practical Approach*, vol. I & II (D. Glover, ed.); *Oligonucleotide Synthesis* (N. Gait, ed., 1984); *Nucleic Acid Hybridization* (B. Hames & S. Higgins, eds., 1985); *Transcription and Translation* (B. Hames & S. Higgins, eds., 1984); *Animal Cell Culture* (R. Freshney, ed., 1986); Perbal, *A Practical Guide to Molecular Cloning* (1984) and other like references.

[0050] Before describing certain embodiments in detail, it is to be understood that this invention is not limited to particular compositions or biological systems, which can vary. It is also to be understood that the terminology used herein is for the purpose of describing particular illustrative embodiments only, and is not intended to be limiting. The terms used in this specification generally have their ordinary meaning in the art, within the context of this invention and in the specific context where each term is used. Certain terms are discussed below or elsewhere in the specification, to provide additional guidance to the practitioner in describing the compositions and methods of the invention and how to make and use them. The scope and meaning of any use of a term will be apparent from the specific context in which the term is used. As such, the definitions set forth herein are intended to provide illustrative guidance in ascertaining particular embodiments of the invention, without limitation to particular compositions or biological systems.

[0051] As used in the present disclosure and the appended claims, the singular forms “a,” “an” and “the” include plural references unless the content clearly dictates otherwise.

[0052] Throughout the present disclosure and the appended claims, unless the context requires otherwise, the word “comprise”, or variations such as “comprises” or “comprising”, will be understood to imply the inclusion of a stated element or group of elements but not the exclusion of any other element or group of elements.

[0053] The terms “antibody” and “immunoglobulin” (Ig) are used interchangeably herein. An antibody may be either membrane bound or secreted. As used herein, the term encompasses not only intact, or “whole”, polyclonal or monoclonal antibodies, but also fragments thereof (such as single-variable domain (VH, VL or combination thereof) antibodies, Fab, Fab', F(ab')₂, Fv), single chain (ScFv), synthetic variants thereof, naturally

occurring variants, fusion proteins comprising an antibody portion with an antigen-binding fragment of the required specificity, humanized antibodies, chimeric antibodies, and any other modified configuration of the immunoglobulin molecule that comprises an antigen-binding site or fragment (epitope recognition site) of the required specificity.

[0054] Antibody, or Ig, molecules are typically comprised of two identical heavy chains and two identical light chains linked together through disulfide bonds. Both heavy chains (IgH) and light chains (IgL) contain a variable (V) region or domain and a constant (C) region or domain. The portion of the IgH locus encoding the V region comprises multiple copies of variable (V), diversity (D), and joining (J) gene segments. The portion of the IgL loci encoding the V region comprises multiple copies of V and J gene segments. The V region encoding portion of the IgH and IgL loci undergo gene segment rearrangement, e.g., different combinations of a V, (D) and J gene segments arrange to form the IgH and IgL variable regions, to develop diverse antigen specificity in antibodies. Each variable region comprises three complementarity-determining regions (CDRs) interspersed between the less variable framework regions (FRs). The heavy chain comprises VHCDR1, VHCDR2, and VHCDR3. The light chain comprises VLCDR1, VLCDR2, and VLCDR3. The secreted form of the IgH C region is made up of three C domains, CH1, CH2, CH3, optionally CH4 (C μ), and a hinge region except for C μ , which lacks a hinge region. The membrane-bound form of the IgH C region also has membrane and intra-cellular domains. The IgH constant region determines the isotype of the antibody, e.g. IgM, IgD, IgG1, IgG2, IgG3, IgG4, IgA and IgE. It will be appreciated that non-human mammals, such as an AlivaMab Mouse, encoding multiple Ig isotypes will be able to undergo isotype class switching. There are two types of human IgL, Ig κ and Ig λ .

[0055] The term "antigen-binding fragment" as used herein refers to a polypeptide fragment that contains at least one CDR of an immunoglobulin heavy and/or light chain that binds to CD115. In this regard, an antigen-binding fragment of the antibodies may comprise 1, 2, 3, 4, 5, or all 6 CDRs of a VH and VL sequence set forth herein from anti-CD115 antibodies described herein. An antigen-binding fragment of the CD115-specific antibodies described herein is capable of binding to CD115. In certain embodiments, an antigen-binding fragment or an antibody comprising an antigen-binding fragment, prevents or inhibits CSF-1 and/or IL-34 binding to CD115 and subsequent signaling events. In other embodiments, an anti-CD115 antibody, or an antigen-binding fragment thereof, prevents signaling events mediated by CD115 by preventing dimerization of CD115, including dimerization that is

induced by CSF-1 or IL-34 binding or that may happen spontaneously under certain conditions of expression CD115. In certain embodiments, the antigen-binding fragment binds specifically to and/or inhibits or modulates the biological activity of human CD115.

[0056] In certain embodiments, antibodies and antigen-binding fragments thereof as described herein include a heavy chain and a light chain CDR set, respectively interposed between a heavy chain and a light chain framework region (FR) set that provide conformational support to the CDRs and define the spatial relationship of the CDRs relative to each other. As used herein, the term "CDR set" refers to the three hypervariable regions of a heavy or light chain V region. Proceeding from the N terminus of a heavy or light chain, these regions are denoted as "CDR1," "CDR2," and "CDR3" respectively. An antigen-binding site, therefore, includes six CDRs, comprising the CDR set from each of a heavy and a light chain V region.

[0057] A "Fab" domain or fragment comprises the N-terminal portion of the IgH, which includes the V region and the CH1 domain of the IgH, and the entire IgL. A "F(ab')₂" domain comprises the Fab domain and a portion of the hinge region, wherein the 2 IgH are linked together via disulfide linkage in the middle hinge region. Both the Fab and F(ab')₂ are "antigen-binding fragments." The C-terminal portion of the IgH, comprising the CH2 and CH3 domains, is the "Fc" domain. The Fc domain is the portion of the Ig recognized by cell receptors, such as the FcR, and to which the complement-activating protein, C1q, binds. The lower hinge region, which is encoded in the 5' portion of the CH2 exon, provides flexibility within the antibody for binding to FcR receptors. An "Fv" fragment includes a non-covalent VH::VL heterodimer including an antigen-binding site. In certain embodiments, single chain Fv (scFv) antibodies are contemplated. A scFv is a covalently linked VH::VL heterodimer which is expressed from a gene fusion including VH- and VL-encoding genes linked by a peptide-encoding linker (Huston et al. (1988) Proc. Nat. Acad. Sci. USA 85(16):5879-5883).

[0058] Where bispecific antibodies are to be used, these may be conventional bispecific antibodies, which can be manufactured in a variety of ways (Holliger, P. and Winter G. Current Opinion Biotechnol. 4, 446-449 (1993)), e.g., prepared chemically or from hybrid hybridomas, or may be any of the bispecific antibody fragments mentioned above.

[0059] As used herein "chimeric antibody" refers to an antibody encoded by a polynucleotide sequence containing polynucleotide sequences from two or more species, e.g., human and mouse.

[0060] As used herein "chimeric Ig chain" refers to an Ig heavy chain or an Ig light chain encoded by a polynucleotide sequence containing polynucleotide sequences from two

or more species, e.g., human and mouse. For example, a chimeric Ig heavy chain may comprise a human VH domain, DH domain, JH domain, CH1 domain, and upper hinge region and mouse CH2 and CH3 domains. In one embodiment, the middle hinge region is mouse. In one embodiment, the middle hinge region is human. In one embodiment, the middle hinge region is chimeric.

[0061] “Polypeptide,” “peptide” or “protein” are used interchangeably herein to describe a chain of amino acids that are linked together by chemical bonds. A polypeptide or protein may be an IgH, IgL, V domain, C domain, or an antibody.

[0062] The strength, or affinity of immunological binding interactions can be expressed in terms of the dissociation constant (K_D) of the interaction, wherein a smaller K_D represents a greater affinity. Immunological binding properties of selected polypeptides can be quantified using methods well known in the art. One such method entails measuring the rates of antigen-binding site/antigen complex formation and dissociation, wherein those rates depend on the concentrations of the complex partners, the affinity of the interaction, and on geometric parameters that equally influence the rate in both directions. Thus, both the “on rate constant” (K_{on}) and the “off rate constant” (K_{off}) can be determined by calculation of the concentrations and the actual rates of association and dissociation. The ratio of K_{off}/K_{on} enables cancellation of all parameters not related to affinity, and is thus equal to the dissociation constant, K_D . See, generally, Davies et al. (1990) Annual Rev. Biochem. 59:439-473.

[0063] “Polynucleotide” refers to a chain of nucleic acids that are linked together by chemical bonds. Polynucleotides include, but are not limited to, DNA, cDNA, RNA, mRNA, and gene sequences and segments. Polynucleotides may be isolated from a living source such as a eukaryotic cell, prokaryotic cell or virus, or may be derived through in vitro manipulation by using standard techniques of molecular biology, or by DNA synthesis, or by a combination of a number of techniques.

[0064] As used herein, the term “vector” refers to a nucleic acid molecule into which another nucleic acid fragment can be integrated without loss of the vector's ability to replicate. Vectors may originate from a virus, a plasmid or the cell of a higher organism. Vectors are utilized to introduce foreign or recombinant DNA into a host cell, wherein the vector is replicated.

[0065] A polynucleotide agent can be contained in a vector, which can facilitate manipulation of the polynucleotide, including introduction of the polynucleotide into a target cell. The vector can be a cloning vector, which is useful for maintaining the polynucleotide,

or can be an expression vector, which contains, in addition to the polynucleotide, regulatory elements useful for expressing the polynucleotide and, where the polynucleotide encodes an RNA, for expressing the encoded RNA in a particular cell, either for subsequent translation of the RNA into a polypeptide or for subsequent trans regulatory activity by the RNA in the cell. An expression vector can contain the expression elements necessary to achieve, for example, sustained transcription of the encoding polynucleotide, or the regulatory elements can be operatively linked to the polynucleotide prior to its being cloned into the vector.

[0066] An expression vector (or the polynucleotide) generally contains or encodes a promoter sequence, which can provide constitutive or, if desired, inducible or tissue specific or developmental stage specific expression of the encoding polynucleotide, a poly-A recognition sequence, and a ribosome recognition site or internal ribosome entry site, or other regulatory elements such as an enhancer, which can be tissue specific. The vector also can contain elements required for replication in a prokaryotic or eukaryotic host system or both, as desired. Such vectors, which include plasmid vectors and viral vectors such as bacteriophage, baculovirus, retrovirus, lentivirus, adenovirus, vaccinia virus, alpha virus and adeno-associated virus vectors, are well known and can be purchased from a commercial source (Promega, Madison Wis.; Stratagene, La Jolla Calif.; GIBCO/BRL, Gaithersburg Md.) or can be constructed by one skilled in the art (see, for example, Meth. Enzymol., Vol. 185, Goeddel, ed. (Academic Press, Inc., 1990); Jolly, Canc. Gene Ther. 1:51-64, 1994; Flotte, J. Bioenerg. Biomemb 25:37-42, 1993; Kirshenbaum et al., J. Clin. Invest 92:381-387, 1993; each of which is incorporated herein by reference).

[0067] The term "construct" as used herein refers to a sequence of DNA artificially constructed by genetic engineering, recombineering or synthesis. In one embodiment, the DNA constructs are linearized prior to recombination. In another embodiment, the DNA constructs are not linearized prior to recombination.

[0068] The terms "inhibit", "neutralize", and "antagonize" are used interchangeably herein and encompass anti-CD115 antibodies that block, inhibit, and/or decrease the activity of CD115. Examples of CD115 activity include kinase function and ligand binding, e.g., binding to CSF-1 and/or IL-34.

[0069] The term "treating" with regard to a subject, refers to improving at least one symptom of the subject's disease or disorder. Treating includes curing, improving, or at least partially ameliorating the disease or disorder.

[0070] As used herein, the term "disorder" refers to, and is used interchangeably with, the terms disease, condition, or illness.

[0071] The term “pharmaceutically acceptable carrier” refers generally to any material (e.g., carrier, excipient, or stabilizer) that may accompany a therapeutic agent and is nontoxic to the subject or patient being exposed thereto.

[0072] The term “administering,” as used herein, refers to any mode of transferring, delivering, introducing, or transporting a pharmaceutical composition or other agent, such as an anti-CD115 antibody, to a subject. Such modes include oral administration, topical contact, intravenous, intraperitoneal, intramuscular, intranasal, or subcutaneous administration.

[0073] The term “inhibit” or “neutralize” or “block” may relate generally to the ability of one or more anti-CD115 antibodies of the invention to decrease a biological activity of CD115, such as intracellular signaling and/or ligand binding. The inhibition/blocking of CSF-1 and/or IL-34 to CD115 preferably reduces or alters the normal level or type of cell signaling that occurs when CSF-1 and/or IL-34 binds to CD115 without inhibition or blocking. Inhibition and blocking are also intended to include any measurable decrease in the binding of CSF-1 and/or IL-34 to CD115 when in contact with an anti CD115 antibody as disclosed herein as compared to the ligand not in contact with an anti CD115 antibody, e.g., the blocking of CSF-1 and/or IL-34 to CD115 by at least about a 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, 15%, 16%, 17%, 18%, 19%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% decrease, including all integers in between. In one embodiment, a neutralizing anti-CD115 antibody inhibits binding of CSF-1 and/or IL-34 to CD115 by at least about a 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 11%, 12%, 13%, 14%, 15%, 16%, 17%, 18%, 19%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or 100% decrease, including all integers in between.

[0074] An antibody, or antigen-binding fragment thereof, is said to “specifically bind,” “immunologically bind,” and/or is “immunologically reactive” to CD115 if it reacts at a detectable level (within, for example, an ELISA assay) with CD115, and does not react detectably with unrelated polypeptides under similar conditions. Antibodies are considered to specifically bind to a target polypeptide when the binding affinity is at least 1×10^{-7} M or, preferably, at least 1×10^{-8} M. In one embodiment, the antibody, or antigen-binding fragment thereof, specifically binds human CD115.

[0075] Each embodiment in this specification is to be applied mutatis mutandis to every other embodiment unless expressly stated otherwise.

[0076] Standard techniques may be used for recombinant DNA, oligonucleotide synthesis, and tissue culture and transformation (e.g., electroporation, lipofection). Enzymatic reactions and purification techniques may be performed according to manufacturer's specifications or as commonly accomplished in the art or as described herein. These and related techniques and procedures may be generally performed according to conventional methods well known in the art and as described in various general and more specific references that are cited and discussed throughout the present specification. Unless specific definitions are provided, the nomenclature utilized in connection with, and the laboratory procedures and techniques of, molecular biology, analytical chemistry, synthetic organic chemistry, and medicinal and pharmaceutical chemistry described herein are those well-known and commonly used in the art. Standard techniques may be used for recombinant technology, molecular biological, microbiological, chemical syntheses, chemical analyses, pharmaceutical preparation, formulation, and delivery, and treatment of patients.

CD115

[0077] CD115 is expressed by a variety of cells, including, but not limited to, hematopoietic stem cells (HSCs); myeloid cells, including monocytes, macrophages, osteoclasts, dendritic cells, and microglia; neural progenitor cells; and epithelial cells, including Paneth cells (Stanley and Chitu, Cold Spring Harb Perspect Biol 2014;6:a021857). Dysregulation of CD115, and/or its ligands, is associated with proliferative diseases and disorders (e.g., neoplasms, tumors and metastases), as well as immunological and neurological diseases and disorders, making it an important therapeutic target.

Anti-CD115 Antibodies

[0078] AlivaMab Mouse anti-CD115 antibodies were generated using both AlivaMab Mouse Kappa mice and AlivaMab Mouse Lambda mice (also referred to herein interchangeably as AlivaMab Kappa Mice and AlivaMab Lambda Mice, respectively). Antibodies produced by AlivaMab Kappa Mice comprise a chimeric immunoglobulin heavy (IgH) chain and a human immunoglobulin kappa (Igκ) light chain. Antibodies produced by AlivaMab Lambda Mice comprise a chimeric IgH chain and a human immunoglobulin lambda (Igλ) light chain. The chimeric IgH chain of the AlivaMab Mouse antibodies comprises a human variable region comprising a human variable heavy (VH) domain, a human diversity heavy (DH) domain, and a human joining heavy (JH) domain, a human constant heavy 1 (CH1) domain, a human upper hinge region (except for C_μ, which is

naturally missing an upper hinge region), a mouse middle hinge region, a mouse CH2 domain, and a mouse CH3 domain. Upon identification of a lead candidate antibody, e.g., an anti-CD115 antibody, the human heavy chain variable region is readily appended to a fully human constant region while maintaining the antigen-binding characteristics of the parent chimeric antibody that were developed *in vivo* in the AlivaMab Mouse. In one embodiment, the human heavy chain variable region, CH1 and, optionally, upper hinge region of the chimeric antibody are appended to human hinge, a human CH2 domain and a human CH3 domain in order to produce a fully human antibody.

[0079] Accordingly, in one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, of the invention is chimeric. In one embodiment, the chimeric anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a chimeric IgH chain and a human Ig κ chain. In one embodiment, the chimeric anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a chimeric IgH chain and a human Ig λ chain. In one embodiment, the chimeric anti-CD115 antibody is human and mouse. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, of the invention is human. In one embodiment, the human anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a human IgH chain and a human Ig κ chain. In one embodiment, the human anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a human IgH chain and a human Ig λ chain. In one embodiment, the isotype of the anti-CD115 antibody is selected from IgM, IgD, IgG1, IgG2, IgG3, IgG4, IgA and IgE. In one embodiment, the isotype of the anti-CD115 antibody is selected from IgG1, IgG2, IgG3, and IgG4.

[0080] In one embodiment, the anti-CD115 antibody binds an Fc receptor (FcR) selected from an Fc γ R, an Fc ϵ R, and an Fc α R. In one embodiment, the anti-CD115 antibody binds an Fc γ R selected from Fc γ RI (CD64), Fc γ RII (CD32), and Fc γ RIII (CD16), including isoforms thereof. In one embodiment, the Fc region of the anti-CD115 antibody comprises a mutation so that it preferentially binds a particular Fc γ R (see, e.g., U.S. 6,737,056 and U.S. 2015/0031862).

[0081] In one aspect of the invention, the CDRs of an anti-CD115 antibody, or antigen-binding fragment thereof, may be mixed and matched between the CDRs of antibody clones described herein. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, comprises a VHCDR1 comprising any one of SEQ ID NOs:436-543, a VHCDR2 comprising any one of SEQ ID NOs:868-975, and a VHCDR3 comprising any one of SEQ ID NOs:1300-1407. In one embodiment, the VHCDR1, VHCDR2 and VHCDR3 are

selected from three different anti-CD115 clones disclosed herein. In one embodiment, the VHCDR1, VHCDR2 and VHCDR3 are selected from two different anti-CD115 clones disclosed herein.

[0082] In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VLCDR1 comprising any one of SEQ ID NOs:652-759, a VLCDR2 comprising any one of SEQ ID NOs:1084-1191, and a VLCDR3 comprising any one of SEQ ID NOs:1516-1623. In one embodiment, the VLCDR1, VLCDR2 and VLCDR3 are selected from three different anti-CD115 clones disclosed herein. In one embodiment, the VLCDR1, VLCDR2 and VLCDR3 are selected from two different anti-CD115 clones disclosed herein.

[0083] In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises 1) comprises a VHCDR1 comprising any one of SEQ ID NOs: 436-543, a VHCDR2 comprising any one of SEQ ID NOs: 868-975, and a VHCDR3 comprising any one of SEQ ID NOs: 300-1407, and 2) a VLCDR1 comprising any one of SEQ ID NOs: 652-759, a VLCDR2 comprising any one of SEQ ID NOs: 1084-1191, and a VLCDR3 comprising any one of SEQ ID NOs: 1516-1623.

[0084] In one aspect of the invention, the CDRs of an anti-CD115 antibody, or antigen-binding fragment thereof, are from the same anti-CD115 antibody clone disclosed herein. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2 and a VHCDR3 from the same anti-CD115 clone disclosed herein. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2, and a VHCDR3 of a VH selected from any one of SEQ ID NOs:109-216. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2, and a VHCDR3 comprising the corresponding sequences listed in Table 3.

[0085] Accordingly, in one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2, and a VHCDR3 selected from: SEQ ID NOs:436, 868, and 1300; SEQ ID NOs:437, 869, and 1301; SEQ ID NOs:438, 870, and 1302; SEQ ID NOs:439, 871, and 1303; SEQ ID NOs:440, 872, and 1304; SEQ ID NOs:441, 873, and 1305; SEQ ID NOs:442, 874, and 1306; SEQ ID NOs:443, 875, and 1307; SEQ ID NOs:444, 876, and 1308; SEQ ID NOs:445, 877, and 1309; SEQ ID NOs:446, 878, and 1310; SEQ ID NOs:447, 879, and 1311; SEQ ID NOs:448, 880, and 1312; SEQ ID NOs:449, 881, and 1313; SEQ ID NOs:450, 882, and 1314; SEQ ID NOs:451, 883, and 1315; SEQ ID NOs:452, 884, and 1316; SEQ ID NOs:453, 885, and 1317; SEQ ID NOs:454, 886, and 1318; SEQ ID NOs:455, 887, and 1319; SEQ ID NOs:456, 888, and 1320; SEQ ID

NOs:457, 889, and 1321; SEQ ID NOs:458, 890, and 1322; SEQ ID NOs:459, 891, and 1323; SEQ ID NOs:460, 892, and 1324; SEQ ID NOs:461, 893, and 1325; SEQ ID NOs:462, 894, and 1326; SEQ ID NOs:463, 895, and 1327; SEQ ID NOs:464, 896, and 1328; SEQ ID NOs:465, 897, and 1329; SEQ ID NOs:466, 898, and 1330; SEQ ID NOs:467, 899, and 1331; SEQ ID NOs:468, 900, and 1332; SEQ ID NOs:469, 901, and 1333; SEQ ID NOs:470, 902, and 1334; SEQ ID NOs:471, 903, and 1335; SEQ ID NOs:472, 904, and 1336; SEQ ID NOs:473, 905, and 1337; SEQ ID NOs:474, 906, and 1338; SEQ ID NOs:475, 907, and 1339; SEQ ID NOs:476, 908, and 1340; SEQ ID NOs:477, 909, and 1341; SEQ ID NOs:478, 910, and 1342; SEQ ID NOs:479, 911, and 1343; SEQ ID NOs:480, 912, and 1344; SEQ ID NOs:481, 913, and 1345; SEQ ID NOs:482, 914, and 1346; SEQ ID NOs:483, 915, and 1347; SEQ ID NOs:484, 916, and 1348; SEQ ID NOs:485, 917, and 1349; SEQ ID NOs:486, 918, and 1350; SEQ ID NOs:487, 919, and 1351; SEQ ID NOs:488, 920, and 1352; SEQ ID NOs:489, 921, and 1353; SEQ ID NOs:490, 922, and 1354; SEQ ID NOs:491, 923, and 1355; SEQ ID NOs:492, 924, and 1356; SEQ ID NOs:493, 925, and 1357; SEQ ID NOs:494, 926, and 1358; SEQ ID NOs:495, 927, and 1359; SEQ ID NOs:496, 928, and 1360; SEQ ID NOs:497, 929, and 1361; SEQ ID NOs:498, 930, and 1362; SEQ ID NOs:499, 931, and 1363; SEQ ID NOs:500, 932, and 1364; SEQ ID NOs:501, 933, and 1365; SEQ ID NOs:502, 934, and 1366; SEQ ID NOs:503, 935, and 1367; SEQ ID NOs:504, 936, and 1368; SEQ ID NOs:505, 937, and 1369; SEQ ID NOs:506, 938, and 1370; SEQ ID NOs:507, 939, and 1371; SEQ ID NOs:508, 940, and 1372; SEQ ID NOs:509, 941, and 1373; SEQ ID NOs:510, 942, and 1374; SEQ ID NOs:511, 943, and 1375; SEQ ID NOs:512, 944, and 1376; SEQ ID NOs:513, 945, and 1377; SEQ ID NOs:514, 946, and 1378; SEQ ID NOs:515, 947, and 1379; SEQ ID NOs:516, 948, and 1380; SEQ ID NOs:517, 949, and 1381; SEQ ID NOs:518, 950, and 1382; SEQ ID NOs:519, 951, and 1383; SEQ ID NOs:520, 952, and 1384; SEQ ID NOs:521, 953, and 1385; SEQ ID NOs:522, 954, and 1386; SEQ ID NOs:523, 955, and 1387; SEQ ID NOs:524, 956, and 1388; SEQ ID NOs:525, 957, and 1389; SEQ ID NOs:526, 958, and 1390; SEQ ID NOs:527, 959, and 1391; SEQ ID NOs:528, 960, and 1392; SEQ ID NOs:529, 961, and 1393; SEQ ID NOs:530, 962, and 1394; SEQ ID NOs:531, 963, and 1395; SEQ ID NOs:532, 964, and 1396; SEQ ID NOs:533, 965, and 1397; SEQ ID NOs:534, 966, and 1398; SEQ ID NOs:535, 967, and 1399; SEQ ID NOs:536, 968, and 1400; SEQ ID NOs:537, 969, and 1401; SEQ ID NOs:538, 970, and 1402; SEQ ID NOs:539, 971, and 1403; SEQ ID NOs:540, 972, and 1404; SEQ ID NOs:541, 973, and 1405; SEQ ID NOs:542, 974, and 1406; and SEQ ID NOs:543, 975, and 1407.

[0086] In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VLCDR1, a VLCDR2 and a VLCDR3 from the same anti-CD115 clone disclosed herein. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VLCDR1, a VLCDR2, and a VLCDR3 of a VL selected from any one of SEQ ID NOs:325-432. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VLCDR1, a VLCDR2, and a VLCDR3 comprising the corresponding sequences listed in Table 3.

[0087] Accordingly, in one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VLCDR1, a VLCDR2, and a VLCDR3 selected from: SEQ ID NOs:652, 1084, and 1516; SEQ ID NOs:653, 1085, and 1517; SEQ ID NOs:654, 1086, and 1518; SEQ ID NOs:655, 1087, and 1519; SEQ ID NOs:656, 1088, and 1520; SEQ ID NOs:657, 1089, and 1521; SEQ ID NOs:658, 1090, and 1522; SEQ ID NOs:659, 1091, and 1523; SEQ ID NOs:660, 1092, and 1524; SEQ ID NOs:661, 1093, and 1525; SEQ ID NOs:662, 1094, and 1526; SEQ ID NOs:663, 1095, and 1527; SEQ ID NOs:664, 1096, and 1528; SEQ ID NOs:665, 1097, and 1529; SEQ ID NOs:666, 1098, and 1530; SEQ ID NOs:667, 1099, and 1531; SEQ ID NOs:668, 1100, and 1532; SEQ ID NOs:669, 1101, and 1533; SEQ ID NOs:670, 1102, and 1534; SEQ ID NOs:671, 1103, and 1535; SEQ ID NOs:672, 1104, and 1536; SEQ ID NOs:673, 1105, and 1537; SEQ ID NOs:674, 1106, and 1538; SEQ ID NOs:675, 1107, and 1539; SEQ ID NOs:676, 1108, and 1540; SEQ ID NOs:677, 1109, and 1541; SEQ ID NOs:678, 1110, and 1542; SEQ ID NOs:679, 1111, and 1543; SEQ ID NOs:680, 1112, and 1544; SEQ ID NOs:681, 1113, and 1545; SEQ ID NOs:682, 1114, and 1546; SEQ ID NOs:683, 1115, and 1547; SEQ ID NOs:684, 1116, and 1548; SEQ ID NOs:685, 1117, and 1549; SEQ ID NOs:686, 1118, and 1550; SEQ ID NOs:687, 1119, and 1551; SEQ ID NOs:688, 1120, and 1552; SEQ ID NOs:689, 1121, and 1553; SEQ ID NOs:690, 1122, and 1554; SEQ ID NOs:691, 1123, and 1555; SEQ ID NOs:692, 1124, and 1556; SEQ ID NOs:693, 1125, and 1557; SEQ ID NOs:694, 1126, and 1558; SEQ ID NOs:695, 1127, and 1559; SEQ ID NOs:696, 1128, and 1560; SEQ ID NOs:697, 1129, and 1561; SEQ ID NOs:698, 1130, and 1562; SEQ ID NOs:699, 1131, and 1563; SEQ ID NOs:700, 1132, and 1564; SEQ ID NOs:701, 1133, and 1565; SEQ ID NOs:702, 1134, and 1566; SEQ ID NOs:703, 1135, and 1567; SEQ ID NOs:704, 1136, and 1568; SEQ ID NOs:705, 1137, and 1569; SEQ ID NOs:706, 1138, and 1570; SEQ ID NOs:707, 1139, and 1571; SEQ ID NOs:708, 1140, and 1572; SEQ ID NOs:709, 1141, and 1573; SEQ ID NOs:710, 1142, and 1574; SEQ ID NOs:711, 1143, and 1575; SEQ ID NOs:712, 1144, and 1576; SEQ ID NOs:713, 1145, and 1577; SEQ ID NOs:714, 1146, and

1578; SEQ ID NOs:715, 1147, and 1579; SEQ ID NOs:716, 1148, and 1580; SEQ ID NOs:717, 1149, and 1581; SEQ ID NOs:718, 1150, and 1582; SEQ ID NOs:719, 1151, and 1583; SEQ ID NOs:720, 1152, and 1584; SEQ ID NOs:721, 1153, and 1585; SEQ ID NOs:722, 1154, and 1586; SEQ ID NOs:723, 1155, and 1587; SEQ ID NOs:724, 1156, and 1588; SEQ ID NOs:725, 1157, and 1589; SEQ ID NOs:726, 1158, and 1590; SEQ ID NOs:727, 1159, and 1591; SEQ ID NOs:728, 1160, and 1592; SEQ ID NOs:729, 1161, and 1593; SEQ ID NOs:730, 1162, and 1594; SEQ ID NOs:731, 1163, and 1595; SEQ ID NOs:732, 1164, and 1596; SEQ ID NOs:733, 1165, and 1597; SEQ ID NOs:734, 1166, and 1598; SEQ ID NOs:735, 1167, and 1599; SEQ ID NOs:736, 1168, and 1600; SEQ ID NOs:737, 1169, and 1601; SEQ ID NOs:738, 1170, and 1602; SEQ ID NOs:739, 1171, and 1603; SEQ ID NOs:740, 1172, and 1604; SEQ ID NOs:741, 1173, and 1605; SEQ ID NOs:742, 1174, and 1606; SEQ ID NOs:743, 1175, and 1607; SEQ ID NOs:744, 1176, and 1608; SEQ ID NOs:745, 1177, and 1609; SEQ ID NOs:746, 1178, and 1610; SEQ ID NOs:747, 1179, and 1611; SEQ ID NOs:748, 1180, and 1612; SEQ ID NOs:749, 1181, and 1613; SEQ ID NOs:750, 1182, and 1614; SEQ ID NOs:751, 1183, and 1615; SEQ ID NOs:752, 1184, and 1616; SEQ ID NOs:753, 1185, and 1617; SEQ ID NOs:754, 1186, and 1618; SEQ ID NOs:755, 1187, and 1619; SEQ ID NOs:756, 1188, and 1620; SEQ ID NOs:757, 1189, and 1621; SEQ ID NOs:758, 1190, and 1622; and SEQ ID NOs:759, 1191, and 1623.

[0088] In another aspect of the invention, the CDRs of an anti-CD115 antibody, or antigen-binding fragment thereof, are selected from the corresponding VH and VL of a single clone described herein. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises 1) a VHCDR1, a VHCDR2, and a VHCDR3 selected from the VHCDR1, VHCDR2 and VHCDR3 of one VH selected from any one of SEQ ID NOs: 109-216 and 2) a VLCDR1, a VLCDR2, and a VLCDR3 selected from the VLCDR1, VLCDR2 and VLCDR3 of one VL selected from any one of SEQ ID NOs:325-432. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2, a VHCDR3, a VLCDR1, a VLCDR2, and a VLCDR3 within the corresponding VH and VL amino acid sequences of a single clone as set forth in Table 3.

[0089] Accordingly, in one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VHCDR1, a VHCDR2, a VHCDR3, a VLCDR1, a VLCDR2, and a VLCDR3 selected from: SEQ ID NOs:436, 868, 1300, 652, 1084, and 1516; SEQ ID NOs:437, 869, 1301, 653, 1085, and 1517; SEQ ID NOs:438, 870, 1302, 654, 1086, and 1518; SEQ ID NOs:439, 871, 1303, 655, 1087, and 1519; SEQ ID NOs:440, 872, 1304,

656, 1088, and 1520; SEQ ID NOs:441, 873, 1305, 657, 1089, and 1521; SEQ ID NOs:442, 874, 1306, 658, 1090, and 1522; SEQ ID NOs:443, 875, 1307, 659, 1091, and 1523; SEQ ID NOs:444, 876, 1308, 660, 1092, and 1524; SEQ ID NOs:445, 877, 1309, 661, 1093, and 1525; SEQ ID NOs:446, 878, 1310, 662, 1094, and 1526; SEQ ID NOs:447, 879, 1311, 663, 1095, and 1527; SEQ ID NOs:448, 880, 1312, 664, 1096, and 1528; SEQ ID NOs:449, 881, 1313, 665, 1097, and 1529; SEQ ID NOs:450, 882, 1314, 666, 1098, and 1530; SEQ ID NOs:451, 883, 1315, 667, 1099, and 1531; SEQ ID NOs:452, 884, 1316, 668, 1100, and 1532; SEQ ID NOs:453, 885, 1317, 669, 1101, and 1533; SEQ ID NOs:454, 886, 1318, 670, 1102, and 1534; SEQ ID NOs:455, 887, 1319, 671, 1103, and 1535; SEQ ID NOs:456, 888, 1320, 672, 1104, and 1536; SEQ ID NOs:457, 889, 1321, 673, 1105, and 1537; SEQ ID NOs:458, 890, 1322, 674, 1106, and 1538; SEQ ID NOs:459, 891, 1323, 675, 1107, and 1539; SEQ ID NOs:460, 892, 1324, 676, 1108, and 1540; SEQ ID NOs:461, 893, 1325, 677, 1109, and 1541; SEQ ID NOs:462, 894, 1326, 678, 1110, and 1542; SEQ ID NOs:463, 895, 1327, 679, 1111, and 1543; SEQ ID NOs:464, 896, 1328, 680, 1112, and 1544; SEQ ID NOs:465, 897, 1329, 681, 1113, and 1545; SEQ ID NOs:466, 898, 1330, 682, 1114, and 1546; SEQ ID NOs:467, 899, 1331, 683, 1115, and 1547; SEQ ID NOs:468, 900, 1332, 684, 1116, and 1548; SEQ ID NOs:469, 901, 1333, 685, 1117, and 1549; SEQ ID NOs:470, 902, 1334, 686, 1118, and 1550; SEQ ID NOs:471, 903, 1335, 687, 1119, and 1551; SEQ ID NOs:472, 904, 1336, 688, 1120, and 1552; SEQ ID NOs:473, 905, 1337, 689, 1121, and 1553; SEQ ID NOs:474, 906, 1338, 690, 1122, and 1554; SEQ ID NOs:475, 907, 1339, 691, 1123, and 1555; SEQ ID NOs:476, 908, 1340, 692, 1124, and 1556; SEQ ID NOs:477, 909, 1341, 693, 1125, and 1557; SEQ ID NOs:478, 910, 1342, 694, 1126, and 1558; SEQ ID NOs:479, 911, 1343, 695, 1127, and 1559; SEQ ID NOs:480, 912, 1344, 696, 1128, and 1560; SEQ ID NOs:481, 913, 1345, 697, 1129, and 1561; SEQ ID NOs:482, 914, 1346, 698, 1130, and 1562; SEQ ID NOs:483, 915, 1347, 699, 1131, and 1563; SEQ ID NOs:484, 916, 1348, 700, 1132, and 1564; SEQ ID NOs:485, 917, 1349, 701, 1133, and 1565; SEQ ID NOs:486, 918, 1350, 702, 1134, and 1566; SEQ ID NOs:487, 919, 1351, 703, 1135, and 1567; SEQ ID NOs:488, 920, 1352, 704, 1136, and 1568; SEQ ID NOs:489, 921, 1353, 705, 1137, and 1569; SEQ ID NOs:490, 922, 1354, 706, 1138, and 1570; SEQ ID NOs:491, 923, 1355, 707, 1139, and 1571; SEQ ID NOs:492, 924, 1356, 708, 1140, and 1572; SEQ ID NOs:493, 925, 1357, 709, 1141, and 1573; SEQ ID NOs:494, 926, 1358, 710, 1142, and 1574; SEQ ID NOs:495, 927, 1359, 711, 1143, and 1575; SEQ ID NOs:496, 928, 1360, 712, 1144, and 1576; SEQ ID NOs:497, 929, 1361, 713, 1145, and 1577; SEQ ID NOs:498, 930, 1362, 714, 1146, and 1578; SEQ ID NOs:499, 931, 1363, 715, 1147, and 1579; SEQ ID

NOs:500, 932, 1364, 716, 1148, and 1580; SEQ ID NOs:501, 933, 1365, 717, 1149, and 1581; SEQ ID NOs:502, 934, 1366, 718, 1150, and 1582; SEQ ID NOs:503, 935, 1367, 719, 1151, and 1583; SEQ ID NOs:504, 936, 1368, 720, 1152, and 1584; SEQ ID NOs:505, 937, 1369, 721, 1153, and 1585; SEQ ID NOs:506, 938, 1370, 722, 1154, and 1586; SEQ ID NOs:507, 939, 1371, 723, 1155, and 1587; SEQ ID NOs:508, 940, 1372, 724, 1156, and 1588; SEQ ID NOs:509, 941, 1373, 725, 1157, and 1589; SEQ ID NOs:510, 942, 1374, 726, 1158, and 1590; SEQ ID NOs:511, 943, 1375, 727, 1159, and 1591; SEQ ID NOs:512, 944, 1376, 728, 1160, and 1592; SEQ ID NOs:513, 945, 1377, 729, 1161, and 1593; SEQ ID NOs:514, 946, 1378, 730, 1162, and 1594; SEQ ID NOs:515, 947, 1379, 731, 1163, and 1595; SEQ ID NOs:516, 948, 1380, 732, 1164, and 1596; SEQ ID NOs:517, 949, 1381, 733, 1165, and 1597; SEQ ID NOs:518, 950, 1382, 734, 1166, and 1598; SEQ ID NOs:519, 951, 1383, 735, 1167, and 1599; SEQ ID NOs:520, 952, 1384, 736, 1168, and 1600; SEQ ID NOs:521, 953, 1385, 737, 1169, and 1601; SEQ ID NOs:522, 954, 1386, 738, 1170, and 1602; SEQ ID NOs:523, 955, 1387, 739, 1171, and 1603; SEQ ID NOs:524, 956, 1388, 740, 1172, and 1604; SEQ ID NOs:525, 957, 1389, 741, 1173, and 1605; SEQ ID NOs:526, 958, 1390, 742, 1174, and 1606; SEQ ID NOs:527, 959, 1391, 743, 1175, and 1607; SEQ ID NOs:528, 960, 1392, 744, 1176, and 1608; SEQ ID NOs:529, 961, 1393, 745, 1177, and 1609; SEQ ID NOs:530, 962, 1394, 746, 1178, and 1610; SEQ ID NOs:531, 963, 1395, 747, 1179, and 1611; SEQ ID NOs:532, 964, 1396, 748, 1180, and 1612; SEQ ID NOs:533, 965, 1397, 749, 1181, and 1613; SEQ ID NOs:534, 966, 1398, 750, 1182, and 1614; SEQ ID NOs:535, 967, 1399, 751, 1183, and 1615; SEQ ID NOs:536, 968, 1400, 752, 1184, and 1616; SEQ ID NOs:537, 969, 1401, 753, 1185, and 1617; SEQ ID NOs:538, 970, 1402, 754, 1186, and 1618; SEQ ID NOs:539, 971, 1403, 755, 1187, and 1619; SEQ ID NOs:540, 972, 1404, 756, 1188, and 1620; SEQ ID NOs:541, 973, 1405, 757, 1189, and 1621; SEQ ID NOs:542, 974, 1406, 758, 1190, and 1622; and SEQ ID NOs:543, 975, 1407, 759, 1191, and 1623.

[0090] In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, comprises a VH comprising any one of SEQ ID NOs: 109-216. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, comprises a VL comprising any one of SEQ ID NOs:325-432. In one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a corresponding VH and VL of a single clone as set forth in Table 3.

[0091] Accordingly, in one embodiment, an anti-CD115 antibody, or an antigen-binding fragment thereof, comprises a VH and a VL selected from: SEQ ID NOs:109 and

325; SEQ ID NOs:110 and 326; SEQ ID NOs:111 and 327; SEQ ID NOs:112 and 328; SEQ ID NOs:113 and 329; SEQ ID NOs:114 and 330; SEQ ID NOs:115 and 331; SEQ ID NOs:116 and 332; SEQ ID NOs:117 and 333; SEQ ID NOs:118 and 334; SEQ ID NOs:119 and 335; SEQ ID NOs:120 and 336; SEQ ID NOs:121 and 337; SEQ ID NOs:122 and 338; SEQ ID NOs:123 and 339; SEQ ID NOs:124 and 340; SEQ ID NOs:125 and 341; SEQ ID NOs:126 and 342; SEQ ID NOs:127 and 343; SEQ ID NOs:128 and 344; SEQ ID NOs:129 and 345; SEQ ID NOs:130 and 346; SEQ ID NOs:131 and 347; SEQ ID NOs:132 and 348; SEQ ID NOs:133 and 349; SEQ ID NOs:134 and 350; SEQ ID NOs:135 and 351; SEQ ID NOs:136 and 352; SEQ ID NOs:137 and 353; SEQ ID NOs:138 and 354; SEQ ID NOs:139 and 355; SEQ ID NOs:140 and 356; SEQ ID NOs:141 and 357; SEQ ID NOs:142 and 358; SEQ ID NOs:143 and 359; SEQ ID NOs:144 and 360; SEQ ID NOs:145 and 361; SEQ ID NOs:146 and 362; SEQ ID NOs:147 and 363; SEQ ID NOs:148 and 364; SEQ ID NOs:149 and 365; SEQ ID NOs:150 and 366; SEQ ID NOs:151 and 367; SEQ ID NOs:152 and 368; SEQ ID NOs:153 and 369; SEQ ID NOs:154 and 370; SEQ ID NOs:155 and 371; SEQ ID NOs:156 and 372; SEQ ID NOs:157 and 373; SEQ ID NOs:158 and 374; SEQ ID NOs:159 and 375; SEQ ID NOs:160 and 376; SEQ ID NOs:161 and 377; SEQ ID NOs:162 and 378; SEQ ID NOs:163 and 379; SEQ ID NOs:164 and 380; SEQ ID NOs:165 and 381; SEQ ID NOs:166 and 382; SEQ ID NOs:167 and 383; SEQ ID NOs:168 and 384; SEQ ID NOs:169 and 385; SEQ ID NOs:170 and 386; SEQ ID NOs:171 and 387; SEQ ID NOs:172 and 388; SEQ ID NOs:173 and 389; SEQ ID NOs:174 and 390; SEQ ID NOs:175 and 391; SEQ ID NOs:176 and 392; SEQ ID NOs:177 and 393; SEQ ID NOs:178 and 394; SEQ ID NOs:179 and 395; SEQ ID NOs:180 and 396; SEQ ID NOs:181 and 397; SEQ ID NOs:182 and 398; SEQ ID NOs:183 and 399; SEQ ID NOs:184 and 400; SEQ ID NOs:185 and 401; SEQ ID NOs:186 and 402; SEQ ID NOs:187 and 403; SEQ ID NOs:188 and 404; SEQ ID NOs:189 and 405; SEQ ID NOs:190 and 406; SEQ ID NOs:191 and 407; SEQ ID NOs:192 and 408; SEQ ID NOs:193 and 409; SEQ ID NOs:194 and 410; SEQ ID NOs:195 and 411; SEQ ID NOs:196 and 412; SEQ ID NOs:197 and 413; SEQ ID NOs:198 and 414; SEQ ID NOs:199 and 415; SEQ ID NOs:200 and 416; SEQ ID NOs:201 and 417; SEQ ID NOs:202 and 418; SEQ ID NOs:203 and 419; SEQ ID NOs:204 and 420; SEQ ID NOs:205 and 421; SEQ ID NOs:206 and 422; SEQ ID NOs:207 and 423; SEQ ID NOs:208 and 424; SEQ ID NOs:209 and 425; SEQ ID NOs:210 and 426; SEQ ID NOs:211 and 427; SEQ ID NOs:212 and 428; SEQ ID NOs:213 and 429; SEQ ID NOs:214 and 430; SEQ ID NOs:215 and 431; and SEQ ID NOs:216 and 432.

[0092] In one embodiment, an anti-CD115 antibody is a whole antibody. In one embodiment, an anti-CD115 antibody is a single chain antibody. In one embodiment, an anti-CD115 antibody is a scFv. In one embodiment, an anti-CD115 antibody is a Fab. In one embodiment, an anti-CD115 antibody is a F(ab')₂. In one embodiment, an anti-CD115 antibody is a Fv.

[0093] In one embodiment, an anti-CD115 antibody is a bispecific antibody. In one embodiment, a bispecific anti-CD115 antibody specifically recognizes two different epitopes of CD115. In one embodiment, a bispecific anti-CD115 comprises a first CDR set comprising the VHCDR1, VHCDR2, VHCDR3, VLCDR1, VLCDR2, and VLCDR3 from a first anti-CD115 antibody clone disclosed herein and a second CDR set comprising the VHCDR1, VHCDR2, VHCDR3, VLCDR1, VLCDR2, and VLCDR3 of a second anti-CD115 antibody clone disclosed herein. In one embodiment, a bispecific anti-CD115 comprises a corresponding first VH and first VL of a first anti-CD115 antibody clone disclosed herein and a corresponding second VH and second VL of a second anti-CD115 antibody clone disclosed herein. In one embodiment, a bispecific anti-CD115 antibody specifically recognizes CD115 and another antigen.

Neutralizing Anti-CD115 Antibodies

[0094] One aspect of the present invention provides anti-CD115 antibodies, and antigen-binding fragments thereof, that are CD115 antagonists. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, neutralizes or inhibits one or more ligands of CD115 from binding CD115. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, inhibits CSF-1 from binding CD115. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, inhibits IL-34 from binding CD115. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, inhibits CSF-1 and IL-34 from binding CD115. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, prevents dimerization of CD115, including dimerization that is induced by CSF-1 or IL-34 binding or that may happen spontaneously under certain conditions of expression CD115. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, inhibits CD115 signaling. In one embodiment, an antagonist anti-CD115 antibody, or antigen-binding fragment thereof, inhibits ligand-induced phosphorylation of CD115.

Polynucleotides

[0095] One aspect of the present invention provides a polynucleotide sequence that encodes an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein. In one embodiment, the polynucleotide is a recombinant polynucleotide. In one embodiment, the polynucleotide is cDNA.

[0096] In one embodiment, a polynucleotide sequence encodes a CDR of an anti-CD115 antibody disclosed herein. In one embodiment, the polynucleotide comprises a VHCDR1 polynucleotide sequence selected from any one of SEQ ID NOs:544-651. In one embodiment, the polynucleotide comprises a VHCDR2 polynucleotide sequence selected from any one of SEQ ID NOs:976-1083. In one embodiment, the polynucleotide comprises a VHCDR3 polynucleotide sequence selected from any one of SEQ ID NOs:1408-1515. In one embodiment, the polynucleotide comprises a VLCDR1 polynucleotide sequence selected from any one of SEQ ID NOs:760-867. In one embodiment, the polynucleotide comprises a VLCDR2 polynucleotide sequence selected from any one of SEQ ID NOs:1192-1299. In one embodiment, the polynucleotide comprises a VLCDR3 polynucleotide sequence selected from any one of SEQ ID NOs:1624-1731.

[0097] In one embodiment, a polynucleotide sequence encodes a VH of an anti-CD115 antibody disclosed herein. In one embodiment, the polynucleotide comprises a VH polynucleotide sequence selected from any one of SEQ ID NOs:1-108. In one embodiment, a polynucleotide sequence encodes a VL of an anti-CD115 antibody disclosed herein. In one embodiment, the polynucleotide comprises a VL polynucleotide sequence selected from any one of SEQ ID NOs:217-324. In one embodiment, a polynucleotide sequence encodes a VH and a VL of an anti-CD115 antibody disclosed herein.

[0098] One embodiment of the invention provides a vector comprising a polynucleotide sequence encoding an anti-CD115 antibody, or an antigen-binding fragment thereof, disclosed herein. In one embodiment, the vector is an expression vector. In one embodiment, the vector is a cloning vector. One embodiment of the invention provides a host cell comprising the vector.

Methods of Use

[0099] The AlivaMab antibodies against CD115, and in particular fully human antibodies incorporating all or portions of the heavy chain and light chain variable regions from the AlivaMab antibodies, may have utility for the treatment of human disease including, but not limited to, diseases in oncology and autoimmunity and inflammation. As the understanding of CD115 biology and disease association becomes better known, it is

expected that opportunities for human clinical therapeutic indications may expand. In particular, oncological, immunological, and neurological diseases and disorders are contemplated.

[00100] An anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein may be used in research, diagnostic, and/or therapeutic methods. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein is used to treat diseases and disorders associated with CD115, CSF-1 and/or IL-34. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein is used to treat diseases and disorders associated with CD115 overexpression. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein is used to treat diseases and disorders associated with CSF-1 overexpression. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein is used to treat diseases and disorders associated with IL-34 overexpression. In one embodiment, an anti-CD115 antibody, or antigen-binding fragment thereof, disclosed herein is used to treat diseases and disorders associated with aberrant CD115 signaling.

[00101] Embodiments of the invention pertain to the use of anti-CD115 antibodies, or antigen-binding fragments thereof, for the diagnosis and prognosis of diseases and disorders associated with CD115, CSF-1 and/or IL-34 or aberrant expression thereof.

Modified Anti-CD115 Antibodies and Compositions

[00102] Anti-CD115 antibodies of the present invention, and antigen-binding fragments and variants thereof, may also be conjugated or operably linked to another compound (e.g., therapeutic agent, label, or tag), referred to herein as a conjugate. The conjugate may be a cytotoxic agent, a chemotherapeutic agent, a cytokine, an anti-angiogenic agent, a tyrosine kinase inhibitor, a toxin, a radioisotope, or other therapeutically active agent. Chemotherapeutic agents, cytokines, anti-angiogenic agents, tyrosine kinase inhibitors, and other therapeutic agents are contemplated. In one embodiment, the antibody is conjugated or operably linked to a toxin, including but not limited to small molecule toxins and enzymatically active toxins of bacterial, fungal, plant, animal or synthetic origin, including fragments and/or variants thereof.

[00103] There are many linking groups known in the art for making antibody conjugates, including, for example, those disclosed in U.S. Pat. No. 5,208,020 or EP Patent 0425235 B1, and Chari et al., *Cancer Research* 52: 127-131 (1992). The linking groups include disulfide groups, thioether groups, acid labile groups, photolabile groups, peptidase

labile groups, or esterase labile groups, as disclosed in the above-identified patents, disulfide and thioether groups being preferred.

[00104] The present invention further relates to pharmaceutical compositions and methods of use. The pharmaceutical compositions of the present invention include an antibody, or fragment thereof, in a pharmaceutically acceptable carrier. Pharmaceutical compositions may be administered *in vivo* for the treatment or prevention of a disease or disorder. Furthermore, pharmaceutical compositions comprising an antibody, or a fragment thereof, of the present invention may include one or more agents for use in combination, or may be administered in conjunction with one or more agents. Agents for use in combination with an anti-CD115 antibody disclosed herein include, but are not limited to cytotoxic agents, chemotherapeutic agents, cytokines, anti-angiogenic agents, tyrosine kinase inhibitors, toxins, and radioisotopes.

[00105] The present invention also provides kits relating to any of the antibodies, or fragments thereof, and/or methods described herein. Kits of the present invention may include diagnostic or therapeutic agents. A kit of the present invention may further provide instructions for use of a composition or antibody and packaging. A kit of the present invention may include devices, reagents, containers or other components. Furthermore, a kit of the present invention may also require the use of an apparatus, instrument or device, including a computer.

EXAMPLES

[00106] The Examples below utilize a CD115 phosphorylation assay in order to detect phospho-CD115. SR cells (confirmed CD115 expression by FACS) were serum starved overnight (1% FBS). Cells were treated with M-CSF or IL-34 in the presence of CD115 mAbs for 30 minutes on ice. Cells were lysed in buffer containing phosphatase and protease inhibitors. Lysates were run on R&D systems p-MCSFR DUOSET, which is an ELISA comprising validated phospho-CD115-specific antibody pairs. Exemplary results from a p-CD115 (MCSFR) ELISA using SR cells are shown in Figures 1 and 3.

EXAMPLE 1

PREPARATION OF MONOCLONAL ANTIBODIES TO CD115

[00107] Monoclonal antibodies were prepared in accordance with a general method as described in "Antibodies: A Laboratory Manual" (Harlow and Lane 1988 CSH Press). Eight-

week old AlivaMab Kappa Mice and eight-week old AlivaMab Lambda Mice mice were immunized using a RIMMS protocol. 50 ug of human CD115 extracellular domain (Sino Biological, China 10161-H08H) was mixed with 40 ul (first immunization), 20 ul (immunizations 2-4) or 0 ul (final immunization) Gerbu MM adjuvant (C-C Biotech, Valley Center, CA #3001-6030) and PBS was added to a final volume of 100 ul. The 50 ug mixture was injected in 20 ul portions in 5 locations per mouse; right and left flanks and right and left shoulder/armpit subcutaneously, and the remaining 20 ul intraperitoneally. This was done 5 times per mouse on days, 1, 4, 7, 9, and 11. On Day 14 mice were sacrificed and terminal materials were collected. Spleens and lymph nodes were prepared and fused with CRL-2016 myeloma cells (ATCC) using a PEG based method as generally described in "Antibodies: A Laboratory Manual" (Harlow and Lane 1988 CSH Press) to establish hybridomas.

[00108] Hybridomas were grown in 384-well tissue culture plates and supernatants from individual wells were screened by ELISA for production of antibodies recognizing huCD115. Positive wells were then transferred to 48-well plates, expanded, and supernatants were collected for huCD115 binding confirmation by ELISA. Positive supernatants were also counter-screened against a non-related histidine-tagged protein. Fifty to sixty hybridoma lines each from AlivaMab Kappa Mice and AlivaMab Lambda Mice confirmed to bind CD115 specifically by ELISA were picked at random and single-cell cloned into 96-well plates. One hundred and eight (108) hybridoma lines were cloned. They were grown into colonies and the supernatant from these individual colonies was screened by ELISA to re-confirm monoclonal antibody binding to huCD115. These supernatants were then screened by FACS to confirm binding to native CD115 on OCI-AML5 cells (DSMZ #ACC-247, Table 1 shows results for select antibodies, and Figure 7). Seventy-five hybridoma clones were confirmed to produce mAb that bound to CD115-expressing OCI-AML5 cells (Figure 5).

Table 1. Summary of Screening for Binding to CD115 on cell Surface

HYBRIDOMA	FACS Binding
CCL-247A	+
CCK-423A	+
CCK-415A	+
CCK-416A	+
CCK-541A	+
CCK-424A	+
CCK-507A	+
CCK-461A	+

CCK-421A	+
CCL-331A	+
CCK-422A	+
CCK-437A	+
CCL-327A	+
CCK-522A	+
CCL-309A	+
CCL-321A	+
CCL-332A	+
CCL-217A	+
CCL-328A	+
CCL-221A	+
CCK-402A	+
CCL-238A	+
CCL-245A	+
CCK-417A	+
CCL-215A	+
CCL-346A	+
CCL-213A	+
CCL-205A	+
CCL-216A	+
CCL-211A	+
CCL-204A	+
CCL-325A	+
CCL-337A	+
CCL-249A	+

EXAMPLE 2

SEQUENCES OF ANTI-CD115 VH AND VL

[0109] Total RNA was extracted from hybridomas producing anti-CD115 monoclonal antibodies using the Qiagen RNeasy Mini kit (Cat No. 74104), followed by 5' RACE, using the 5' RACE system kit (Life Technologies, US cat # 18734-058) with the following 3' gene specific primers IgG 5'- GGTTCGGGGAAGTAGTCCTTGACC -3' (SEQ ID NO:433) IgL 5' – CTGTAGCTTCTGTGGGACTTCCACTGCTC -3' (SEQ ID NO:434) IgK 5' – CCGATTGGAGGGCGTTATCCAC -3' (SEQ ID NO:435). RACE products were gel purified and cloned into pCR4-TOPO using TOPO TA cloning kit for sequencing with One Shot Top 10 chemically competent *E. coli* (Life Technologies, US Cat # K4575-01). Sequencing of vector containing colonies was performed by Sequetech (Mountain View, CA)

using M13F or M13R sequencing primers. The reported nucleotide sequences start at the first nucleotide in the first codon for the amino terminal amino acid in framework 1. The reported polypeptide sequences are based on an *in silico* translation of the nucleic acid sequence and start at the first amino acid at the amino terminus of framework 1.

Table 2. Anti-CD115 mAb Amino Acid (aa) and Polynucleotide (nt) Sequences

Clone	VH		VHCDR1		VHCDR2		VHCDR3		VL		VLCDR1		VLCDR2		VLCDR3	
	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:	aa SEQ ID NO:	nt SEQ ID NO:
CCK-401A	109	1	436	544	868	976	1300	1408	325	217	652	760	1084	1192	1516	1624
CCK-402A	110	2	437	545	869	977	1301	1409	326	218	653	761	1085	1193	1517	1625
CCK-406A	111	3	438	546	870	978	1302	1410	327	219	654	762	1086	1194	1518	1626
CCK-407A	112	4	439	547	871	979	1303	1411	328	220	655	763	1087	1195	1519	1627
CCK-408A	113	5	440	548	872	980	1304	1412	329	221	656	764	1088	1196	1520	1628
CCK-410A	114	6	441	549	873	981	1305	1413	330	222	657	765	1089	1197	1521	1629
CCK-412A	115	7	442	550	874	982	1306	1414	331	223	658	766	1090	1198	1522	1630
CCK-414A	116	8	443	551	875	983	1307	1415	332	224	659	767	1091	1199	1523	1631
CCK-415A	117	9	444	552	876	984	1308	1416	333	225	660	768	1092	1200	1524	1632
CCK-416A	118	10	445	553	877	985	1309	1417	334	226	661	769	1093	1201	1525	1633
CCK-417A	119	11	446	554	878	986	1310	1418	335	227	662	770	1094	1202	1526	1634
CCK-418A	120	12	447	555	879	987	1311	1419	336	228	663	771	1095	1203	1527	1635
CCK-421A	121	13	448	556	880	988	1312	1420	337	229	664	772	1096	1204	1528	1636
CCK-422A	122	14	449	557	881	989	1313	1421	338	230	665	773	1097	1205	1529	1637
CCK-423A	123	15	450	558	882	990	1314	1422	339	231	666	774	1098	1206	1530	1638
CCK-424A	124	16	451	559	883	991	1315	1423	340	232	667	775	1099	1207	1531	1639
CCK-425A	125	17	452	560	884	992	1316	1424	341	233	668	776	1100	1208	1532	1640
CCK-434A	126	18	453	561	885	993	1317	1425	342	234	669	777	1101	1209	1533	1641
CCK-435A	127	19	454	562	886	994	1318	1426	343	235	670	778	1102	1210	1534	1642
CCK-436A	128	20	455	563	887	995	1319	1427	344	236	671	779	1103	1211	1535	1643
CCK-437A	129	21	456	564	888	996	1320	1428	345	237	672	780	1104	1212	1536	1644

CCK-455A	130	22	457	565	889	997	1321	1429	346	238	673	781	1105	1213	1537	1645
CCK-456A	131	23	458	566	890	998	1322	1430	347	239	674	782	1106	1214	1538	1646
CCK-458A	132	24	459	567	891	999	1323	1431	348	240	675	783	1107	1215	1539	1647
CCK-459A	133	25	460	568	892	1000	1324	1432	349	241	676	784	1108	1216	1540	1648
CCK-460A	134	26	461	569	893	1001	1325	1433	350	242	677	785	1109	1217	1541	1649
CCK-461A	135	27	462	570	894	1002	1326	1434	351	243	678	786	1110	1218	1542	1650
CCK-464A	136	28	463	571	895	1003	1327	1435	352	244	679	787	1111	1219	1543	1651
CCK-465A	137	29	464	572	896	1004	1328	1436	353	245	680	788	1112	1220	1544	1652
CCK-467A	138	30	465	573	897	1005	1329	1437	354	246	681	789	1113	1221	1545	1653
CCK-468A	139	31	466	574	898	1006	1330	1438	355	247	682	790	1114	1222	1546	1654
CCK-501A	140	32	467	575	899	1007	1331	1439	356	248	683	791	1115	1223	1547	1655
CCK-503A	141	33	468	576	900	1008	1332	1440	357	249	684	792	1116	1224	1548	1656
CCK-505A	142	34	469	577	901	1009	1333	1441	358	250	685	793	1117	1225	1549	1657
CCK-507A	143	35	470	578	902	1010	1334	1442	359	251	686	794	1118	1226	1550	1658
CCK-511A	144	36	471	579	903	1011	1335	1443	360	252	687	795	1119	1227	1551	1659
CCK-513A	145	37	472	580	904	1012	1336	1444	361	253	688	796	1120	1228	1552	1660
CCK-514A	146	38	473	581	905	1013	1337	1445	362	254	689	797	1121	1229	1553	1661
CCK-516A	147	39	474	582	906	1014	1338	1446	363	255	690	798	1122	1230	1554	1662
CCK-519A	148	40	475	583	907	1015	1339	1447	364	256	691	799	1123	1231	1555	1663
CCK-522A	149	41	476	584	908	1016	1340	1448	365	257	692	800	1124	1232	1556	1664
CCK-525A	150	42	477	585	909	1017	1341	1449	366	258	693	801	1125	1233	1557	1665
CCK-526A	151	43	478	586	910	1018	1342	1450	367	259	694	802	1126	1234	1558	1666
CCK-533A	152	44	479	587	911	1019	1343	1451	368	260	695	803	1127	1235	1559	1667
CCK-539A	153	45	480	588	912	1020	1344	1452	369	261	696	804	1128	1236	1560	1668
CCK-541A	154	46	481	589	913	1021	1345	1453	370	262	697	805	1129	1237	1561	1669
CCK-542A	155	47	482	590	914	1022	1346	1454	371	263	698	806	1130	1238	1562	1670
CCK-543A	156	48	483	591	915	1023	1347	1455	372	264	699	807	1131	1239	1563	1671

CCL-201A	157	49	484	592	916	1024	1348	1456	373	265	700	808	1132	1240	1564	1672
CCL-203A	158	50	485	593	917	1025	1349	1457	374	266	701	809	1133	1241	1565	1673
CCL-204A	159	51	486	594	918	1026	1350	1458	375	267	702	810	1134	1242	1566	1674
CCL-205A	160	52	487	595	919	1027	1351	1459	376	268	703	811	1135	1243	1567	1675
CCL-206A	161	53	488	596	920	1028	1352	1460	377	269	704	812	1136	1244	1568	1676
CCL-207A	162	54	489	597	921	1029	1353	1461	378	270	705	813	1137	1245	1569	1677
CCL-208A	163	55	490	598	922	1030	1354	1462	379	271	706	814	1138	1246	1570	1678
CCL-209A	164	56	491	599	923	1031	1355	1463	380	272	707	815	1139	1247	1571	1679
CCL-211A	165	57	492	600	924	1032	1356	1464	381	273	708	816	1140	1248	1572	1680
CCL-212A	166	58	493	601	925	1033	1357	1465	382	274	709	817	1141	1249	1573	1681
CCL-213A	167	59	494	602	926	1034	1358	1466	383	275	710	818	1142	1250	1574	1682
CCL-215A	168	60	495	603	927	1035	1359	1467	384	276	711	819	1143	1251	1575	1683
CCL-216A	169	61	496	604	928	1036	1360	1468	385	277	712	820	1144	1252	1576	1684
CCL-217A	170	62	497	605	929	1037	1361	1469	386	278	713	821	1145	1253	1577	1685
CCL-218A	171	63	498	606	930	1038	1362	1470	387	279	714	822	1146	1254	1578	1686
CCL-220A	172	64	499	607	931	1039	1363	1471	388	280	715	823	1147	1255	1579	1687
CCL-221A	173	65	500	608	932	1040	1364	1472	389	281	716	824	1148	1256	1580	1688
CCL-223A	174	66	501	609	933	1041	1365	1473	390	282	717	825	1149	1257	1581	1689
CCL-225A	175	67	502	610	934	1042	1366	1474	391	283	718	826	1150	1258	1582	1690
CCL-226A	176	68	503	611	935	1043	1367	1475	392	284	719	827	1151	1259	1583	1691
CCL-229A	177	69	504	612	936	1044	1368	1476	393	285	720	828	1152	1260	1584	1692
CCL-231A	178	70	505	613	937	1045	1369	1477	394	286	721	829	1153	1261	1585	1693
CCL-235A	179	71	506	614	938	1046	1370	1478	395	287	722	830	1154	1262	1586	1694
CCL-238A	180	72	507	615	939	1047	1371	1479	396	288	723	831	1155	1263	1587	1695
CCL-245A	181	73	508	616	940	1048	1372	1480	397	289	724	832	1156	1264	1588	1696
CCL-247A	182	74	509	617	941	1049	1373	1481	398	290	725	833	1157	1265	1589	1697
CCL-249A	183	75	510	618	942	1050	1374	1482	399	291	726	834	1158	1266	1590	1698

CCL-252A	184	76	511	619	943	1051	1375	1483	400	292	727	835	1159	1267	1591	1699
CCL-253A	185	77	512	620	944	1052	1376	1484	401	293	728	836	1160	1268	1592	1700
CCL-255A	186	78	513	621	945	1053	1377	1485	402	294	729	837	1161	1269	1593	1701
CCL-301A	187	79	514	622	946	1054	1378	1486	403	295	730	838	1162	1270	1594	1702
CCL-303A	188	80	515	623	947	1055	1379	1487	404	296	731	839	1163	1271	1595	1703
CCL-305A	189	81	516	624	948	1056	1380	1488	405	297	732	840	1164	1272	1596	1704
CCL-309A	190	82	517	625	949	1057	1381	1489	406	298	733	841	1165	1273	1597	1705
CCL-310A	191	83	518	626	950	1058	1382	1490	407	299	734	842	1166	1274	1598	1706
CCL-311A	192	84	519	627	951	1059	1383	1491	408	300	735	843	1167	1275	1599	1707
CCL-312A	193	85	520	628	952	1060	1384	1492	409	301	736	844	1168	1276	1600	1708
CCL-313A	194	86	521	629	953	1061	1385	1493	410	302	737	845	1169	1277	1601	1709
CCL-314A	195	87	522	630	954	1062	1386	1494	411	303	738	846	1170	1278	1602	1710
CCL-315A	196	88	523	631	955	1063	1387	1495	412	304	739	847	1171	1279	1603	1711
CCL-320A	197	89	524	632	956	1064	1388	1496	413	305	740	848	1172	1280	1604	1712
CCL-321A	198	90	525	633	957	1065	1389	1497	414	306	741	849	1173	1281	1605	1713
CCL-322A	199	91	526	634	958	1066	1390	1498	415	307	742	850	1174	1282	1606	1714
CCL-324A	200	92	527	635	959	1067	1391	1499	416	308	743	851	1175	1283	1607	1715
CCL-325A	201	93	528	636	960	1068	1392	1500	417	309	744	852	1176	1284	1608	1716
CCL-327A	202	94	529	637	961	1069	1393	1501	418	310	745	853	1177	1285	1609	1717
CCL-328A	203	95	530	638	962	1070	1394	1502	419	311	746	854	1178	1286	1610	1718
CCL-329A	204	96	531	639	963	1071	1395	1503	420	312	747	855	1179	1287	1611	1719
CCL-331A	205	97	532	640	964	1072	1396	1504	421	313	748	856	1180	1288	1612	1720
CCL-332A	206	98	533	641	965	1073	1397	1505	422	314	749	857	1181	1289	1613	1721
CCL-333A	207	99	534	642	966	1074	1398	1506	423	315	750	858	1182	1290	1614	1722
CCL-335A	208	100	535	643	967	1075	1399	1507	424	316	751	859	1183	1291	1615	1723
CCL-337A	209	101	536	644	968	1076	1400	1508	425	317	752	860	1184	1292	1616	1724
CCL-338A	210	102	537	645	969	1077	1401	1509	426	318	753	861	1185	1293	1617	1725

CCL-339A	211	103	538	646	970	1078	1402	1510	427	319	754	862	1186	1294	1618	1726
CCL-340A	212	104	539	647	971	1079	1403	1511	428	320	755	863	1187	1295	1619	1727
CCL-341A	213	105	540	648	972	1080	1404	1512	429	321	756	864	1188	1296	1620	1728
CCL-344A	214	106	541	649	973	1081	1405	1513	430	322	757	865	1189	1297	1621	1729
CCL-346A	215	107	542	650	974	1082	1406	1514	431	323	758	866	1190	1298	1622	1730
CCL-349A	216	108	543	651	975	1083	1407	1515	432	324	759	867	1191	1299	1623	1731

EXAMPLE 3
EPI TOPE BINNING

[0110] A competition ELISA was performed to establish competitive binding bins. ELISA plates were coated with 1 ug/ml huCD115 protein (Sino Biological, China 10161-H08H) and blocked with Superblock (Thermo Scientific #37518). After washing, wells were incubated with a mouse monoclonal antibody representing one of six unique competition bins and for some of which, exhibit different activities in blocking CSF-1 and/or IL-34 binding to CD115 (Table 7) (these mouse mAbs were generated by hybridoma by immunizing wild-type mice as described in Example 1 as part of a comparator CD115 antibody generation program as part of the first tests of the newly-created AlivaMab Mouse technology). After 1 hour the wells were washed and incubated with individual clonal anti-huCD115 AlivaMab hybridoma supernatants. After another hour the wells were washed and incubated with a specific secondary antibody that either recognized human kappa LC or human lambda LC depending on which AlivaMab Mouse supernatants were being detected (Southern Biotech Goat X hu kappa LC #2061-05 or Bethyl Goat X hu lambda LC #A80-116P) and detected with Supersignal ELISA Pico Chemiluminescent substrate (Thermo Scientific - Product# 37069) (Tables 5 and 6). Individual AlivaMab Mouse antibodies that were able to bind in the presence of a mouse antibody are considered to be in a unique epitope bin from that particular mouse antibody. Individual AlivaMab Mouse antibodies that were unable to bind in the presence of a mouse antibody are considered to be in the same epitope bin as that particular mouse antibody. In this way multiple epitope bins were defined for huCD115 binding antibodies (Tables 3 and 6).

Table 3. Multiple Epitope Bins

	IgGκ	IgGλ	TOTAL
BIN 1	4	0	4
BIN 2	18	16	34
BIN 3	6	28	34
BIN 4	6	8	14
BIN 5	1	5	6
BIN 6	7	2	9
BIN 7	2	0	2
BIN 8	6	0	6

Table 4. Binning of AlivaMab Kappa Mouse Anti-CD115 mAbs

	1A		1B		1C		1D		1E		3		4		5	
Bin 1	522	505	417	462	511	526	542	519	412	402	503	435	437	401	406	
TMR 95A	7	1	3	3	3	9	23	36	3	131	82	86	84	94	87	
Bin 1.5	24	10	19	23	2	3	10	12	110	126	89	75	78	93	88	
TMR 44B	33	0	6	12	99	68	94	110	13	131	97	79	86	96	96	
Bin 1.7	34	53	110	93	77	92	7	6	70	145	95	85	95	104	103	
TMR 20A	83	110	110	77	76	82	111	110	78	11	23	96	101	98	99	
TMR 35A	67	111	113	89	91	84	76	118	91	147	96	17	51	106	109	
Bin 4	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	
NO COMP	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	

Table 4. (continued)

	1A		1B																			
455	458	522	585	417	462	422	424	461	421	525	415	418	599	516	454	416	456	514	467	466	464	
TMR 95A	0	6	7	1	3	3	6	16	15	9	11	10	19	20	29	21	27	42	46	34	57	
TMR 44B	28	17	24	10	19	23	31	53	41	55	56	46	3	21	10	23	53	27	4	22	13	0
TMR 24A	-1	2	33	0	6	12	11	29	27	18	13	31	35	39	32	37	42	53	58	58	39	
TMR 20A	58	22	34	53	110	93	107	98	110	104	80	97	89	82	70	115	94	73	84	96	86	87
TMR 35A	54	66	80	110	110	77	111	102	110	100	88	96	82	85	82	110	107	64	75	83	100	85
TMR 100A	79	79	67	111	113	89	111	98	110	99	100	93	106	85	87	89	104	96	103	102	98	96
NO COMP	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100	100

Table 4. (continued)

1C				1D				1E				
511	526	460	465	429	418	425	542	519	541	549	468	412

TMR 95A	3	9	23	41	38	35	21	23	36	42	1	61	3
TMR 44B	2	3	48	8	45	60	6	10	12	15	2	24	110
TMR 24A	99	68	61	81	75	75	78	94	110	109	111	85	13
TMR 20A	77	92	85	78	84	91	4	7	6	34	59	53	70
TMR 35A	76	82	86	88	112	86	88	111	110	111	89	104	78
TMR 100A	91	84	94	92	107	94	98	76	118	93	100	95	91
NO COM	100	100	100	100	100	100	100	100	100	100	100	100	100

Table 4. (continued)

	3			4			5							
TMR 95A	482	589	587	459	599	488	495	497	481	486	414	496	581	519
TMR 44B	131	82	74	77	97	84	86	84	94	87	86	81	91	77
TMR 24A	126	89	87	84	92	94	75	78	93	88	87	91	106	86
TMR 20A	131	97	106	86	112	86	79	86	96	96	92	87	95	90
TMR 35A	145	95	112	90	102	97	85	95	104	103	106	96	114	92
TMR 100A	11	23	2	3	4	6	96	101	98	99	91	92	106	89
NO COM	147	96	123	95	109	103	17	51	106	109	101	100	116	93
	100	100	100	100	100	100	100	100	100	100	100	100	100	100

Table 5. Epitope Competition Bins in Panels of anti-CD115 mAbs

TMR	BIN 1		BIN 2		BIN 3		BIN 4		BIN 5		BIN 6		BIN 7		BIN 8	
	CCK	505A	CCK	417A	CCK	511A	CCK	542A	CCK	412A	CCK	402A	CCK	435A	CCK	401A
	0	0	0	0	0	0	0	0	0	1	1	1	1	1	1	1
	522A	462A	462A	462A	526A	519A	519A	542A	321A	42A	503A	437A	437A	437A	406A	406A

Table 6. Anti-CD115 mAb Epitope Bins

HYBRIDOMA	BIN	HYBRIDOMA	BIN
CCK-401A	5	CCL-201A	1C
CCK-402A	3	CCL-203A	1C
CCK-406A	5	CCL-204A	1C
CCK-407A	3	CCL-205A	1C
CCK-408A	3	CCL-206A	1B
CCK-410A	1C	CCL-207A	1C
CCK-412A	1E	CCL-208A	1D
CCK-414A	5	CCL-209A	1D
CCK-415A	1B	CCL-211A	1C
CCK-416A	1B	CCL-212A	1B
CCK-417A	1B	CCL-213A	1B
CCK-418A	1B	CCL-215A	1C
CCK-421A	1B	CCL-216A	1C
CCK-422A	1B	CCL-217A	1C
CCK-423A	1C	CCL-218A	1B
CCK-424A	1B	CCL-220A	1C
CCK-425A	1D	CCL-221A	1E
CCK-434A	1B	CCL-223A	1C
CCK-435A	4	CCL-225A	1B
CCK-436A	5	CCL-226A	1C
CCK-437A	4	CCL-229A	1B
CCK-455A	1A	CCL-231A	1E
CCK-456A	1B	CCL-235A	1B
CCK-458A	1A	CCL-238A	1C
CCK-459A	3	CCL-245A	1C
CCK-460A	1D	CCL-247A	1C
CCK-461A	1B	CCL-249A	1C
CCK-464A	1B	CCL-252A	1B
CCK-465A	1C	CCL-253A	1C
CCK-467A	1B	CCL-255A	1C
CCK-468A	1C	CCL-301A	1D
CCK-501A	5	CCL-303A	1C
CCK-503A	3	CCL-305A	1D
CCK-505A	1A	CCL-309A	1E
CCK-507A	3	CCL-310A	1C
CCK-511A	1C	CCL-311A	1B
CCK-513A	5	CCL-312A	3
CCK-514A	1B	CCL-313A	3
CCK-516A	1B	CCL-314A	1B
CCK-519A	1D	CCL-315A	1B
CCK-522A	1A	CCL-320A	1B
CCK-525A	1B	CCL-321A	1E
CCK-526A	1C	CCL-322A	1C
CCK-533A	3	CCL-324A	1B
CCK-539A	1B	CCL-325A	1C
CCK-541A	1D	CCL-327A	1D

CCK-542A	1D	CCL-328A	1D
CCK-543A	1D	CCL-329A	1C
		CCL-331A	1E
		CCL-332A	1C
		CCL-333A	1C
		CCL-335A	1B
		CCL-337A	1C
		CCL-338A	1B
		CCL-339A	1C
		CCL-340A	1C
		CCL-341A	1D
		CCL-344A	1D
		CCL-346A	1C
		CCL-349A	1B

[0111] Based on functional characterization of the bin-defining mouse mAbs, some antibodies within epitope bins 1A or 1C (defined by dual IL-34- and CSF-1-neutralizing mouse mAb, TMR24A) or bin 3 (defined by dual IL-34- and CSF-1-neutralizing mouse mAb, TMR35A) may neutralize P-TYR formation induced by both CSF-1 and IL-34 (Figure 2). Some antibodies within epitope bin 1C (defined by only CSF-1-neutralizing and IL-34 non-neutralizing mouse mAb, TMR20A) may neutralize only CSF-1-induced P-TYR formation on CD115. However, some bin 3 mAbs may neutralize both IL-34 and CSF-1 induced P-TYR formation on CD115. As summarized in Table 7 below, some of the reference wild-type mouse mAbs can block both cytokines from different locations on the receptor (e.g., bin 1 and bin3), some mAbs block M-CSF while not blocking IL-34, none of the mAbs were agonists on their own, and the bin 3 epitope region appears to contain functional diversity (e.g., all 3 mAbs listed below exhibited different activity). Also of note is that mAb 20A slightly increased/enhanced the M-CSF signal (Figures 4A and 4B).

Table 7. Summary of anti-CD115 mAb Activity

mAb	Bin	Blocks M-CSF	Blocks IL-34
24A	1	+++	+++
29A	1	+++	+++
32B	1	+++	+++
20A	2	Slight agonist	NO
10B	3	+++	NO
35A	3	+++	+++

47B	3	NO	NO
40A	4	+	NO
27A	4	+	NO
19B	5	+++	NO
7B	6	NO	NO
44B	6	NO	NO

EXAMPLE 4

AFFINITY DETERMINATION

[0112] Affinity was determined for 24 selected monoclonal hybridoma supernatants (Biosensor Tools, Salt Lake City, Utah). Binding kinetics were measured using a BioRad ProteOn XPR36 optical biosensor equipped with an anti-mouse IgG-Coated GLC sensor chip. Hybridoma supernatants were diluted 10-fold into running buffer and captured for 4 minutes on the anti-mouse IgG surface. Hu CD115 (Sino Biological, China #10161-H08H) was tested in duplicate using a 3-fold dilution series starting at 150 nM. The processed data were fit using a 1:1 interaction model that includes a mass-transport parameter (Scrubber2, Canberra Australia). Within the panel of AlivaMab Mouse anti-CD115 antibodies, there are antibodies with KD values below a nanomolar and KD values in the low nanomolar range, and with fast k_{on} and slow k_{off} rates (Tables 8 and 9).

Table 8. Binding Kinetics of Anti-CD115 IgG λ mAbs

mAb	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})	K_D (pM)
329	1.18×10^5	4.46×10^{-6}	38
310	5.06×10^4	4.35×10^{-6}	79
331	1.86×10^5	2.20×10^{-5}	118
215	8.53×10^4	1.83×10^{-5}	215
225	5.00×10^4	1.38×10^{-5}	277
340	7.41×10^4	2.72×10^{-5}	367
312	1.39×10^5	6.03×10^{-5}	435
206	5.90×10^4	2.92×10^{-5}	495
231	8.10×10^4	4.71×10^{-5}	578
249	1.49×10^5	8.91×10^{-5}	599
217	9.81×10^4	1.20×10^{-4}	1,220
313	7.36×10^4	9.09×10^{-5}	1,240
327	1.33×10^5	1.89×10^{-4}	1,410

Table 9. Binding Kinetics of Anti-CD115 IgGκ mAbs

mAb	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})	K_D (pM)
418	1.33×10^5	2.11×10^{-5}	158
533	1.24×10^5	3.92×10^{-5}	317
412	1.58×10^5	5.94×10^{-4}	376
460	8.92×10^5	5.51×10^{-4}	618
467	2.30×10^4	1.48×10^{-5}	650
459	5.00×10^4	3.38×10^{-5}	680
519	9.00×10^4	7.71×10^{-5}	860
407	6.90×10^4	6.54×10^{-5}	950
541	2.67×10^5	3.74×10^{-4}	1,400
465	5.31×10^5	7.49×10^{-4}	1,410
456	2.70×10^4	5.30×10^{-5}	1,970
539	3.20×10^4	8.50×10^{-5}	2,700

EXAMPLE 5

INTERNALIZATION OF ANTI-CD115 ANTIBODIES

[0113] Anti-CD115 antibodies were tested for their ability to internalize upon binding to native CD115 on the surface of OCI-AML5 cells (DSMZ #ACC-247). OCI-AML5 cells were treated with individual AlivaMab Mouse anti-CD115 supernatants for 1 hour at 37°C. The cells were then transferred to ice and stained with a fluorescently labeled anti-CD115 mAb known to be able to bind CD115 in the presence of bound test antibody (either Biologend Rat x Hu-CD115-PE #6393 or CCK533A conjugated with Dylight488 Pierce #46403). Detection of fluorescent signal was then measured using a BD FACScalibur instrument. Cells that gave a strong fluorescent signal are considered to be non-internalizers for that individual test anti-CD115 mAb. Cells that are measured to have weak or no fluorescent signal are considered to be strong internalizers for that individual test anti-CD115 mAb. This procedure was repeated with several purified AlivaMab Mouse anti-CD115 mAbs that showed internalization as a supernatant at 20 ug/ml. Other AlivaMab Mouse anti-CD115 mAbs are also shown to exhibit various levels of internalization of CD115 induced by mAb binding (Figure 12 and Table 10).

Table 10. Internalization of Anti-CD115 mAbs

mAb	Internalization
-----	-----------------

CCK-423A	-
CCK-543A	+
CCK-416A	++
CCL-252A	+++
CCL-331A	++++

(- = no internalization, 1-4 + = strength of internalization)

EXAMPLE 6

NEUTRALIZATION OF CSF-1 BINDING TO CD115

[0114] Anti-CD115 antibodies were tested for their ability to block binding of recombinant CSF-1 to recombinant CD115. ELISA plates were coated with recombinant hu-CD115 (Sino Biological, China #10161-H08H) at 0.5 ug/ml and blocked with Superblock (Thermo Scientific #37518). Wells were incubated for 15 min with anti-CD115 mAbs, then biotinylated Hu-CSF-1 (R&D Systems, #216-MC-005) (biotinylation using NHS-Peg4-biotin, Life Technologies, #21330) was added to a final concentration of 0.25 ug/ml for an additional 15'. After a 4X wash, CSF-1-biotin was detected using 1:10,000 SAV-poly HRP (Life Technologies, #N200). Other AlivaMab Mouse anti-CD115 mAbs are also shown to exhibit various abilities and IC50 values in blocking CSF-1 binding to CD115 (Figure 10 and Table 11).

Table 11. AlivaMab Mouse anti-CD115 mAbs block CSF-1 binding to CD115

mAb	IC ₅₀ (pM)
CCK-415A	355
CCK-416A	811
CCK-423A	135

EXAMPLE 7

INHIBITION OF P-TYR FORMATION ON CD115 INDUCED BY CSF-1

[0115] Anti-CD115 antibodies were tested for their ability to block hu-CSF1 induced phosphorylation of native CD115. OCI-AML5 cells (DSMZ, # ACC-247) were serum starved (1% FBS) overnight, then harvested and washed twice in PBS with 0.1% BSA. 250,000 cells were plated per well into a 96-well v-bottom polypropylene plate. Anti-CD115 supernatants were added neat for 15 min while incubating the plate on ice. Hu-CSF-1 (R&D Systems, #216-MC-005) was added to each well at a final concentration of 100 ng/ml and incubated for 30' on ice. Cells were then spun down at 1500 RPM for 5' at 4°C and supernatant was removed. Cells were then resuspended in lysis buffer (Cell Signaling, #9803 with 1X HALT protease inhibitors, Pierce, #78430) and incubated on ice for 15 min. Lysates

were then measured for tyrosine phosphorylated CD115 using a p-MCSFR validated DUOSET assay (R&D Systems #CYC3268E) and detected using Supersignal Pico ELISA Substrate (Pierce, #37069) (Figures 6 and 8).

[0116] Unpurified anti-CD115 IgGs (as identified by ELISA as described above) secreted from hybridomas into the tissue culture supernatant was assessed for neutralization of P-TYR formation induced by CSF-1. Neutralization using these unpurified, non-quantified antibodies was rank ordered. From this assessment, sets of the better neutralizing mAbs were identified, one set of IgG κ mAbs and one set of IgG λ mAbs. The hybridomas making these mAbs were grown and mAb purified using a commercially-available kit. The P-TYR neutralization assay was repeated with several purified anti-CD115 mAbs using a dilution series enabling an IC₅₀ calculation, first in an eight-point dilution curve (Figure 9) and then with a further subset of best neutralizing mAbs in a twelve-point dilution curve to better calculate IC₅₀ values. Of the antibodies tested, CCK423 was identified as having the best IC₅₀ for neutralizing CSF-1 P-TYR formation on CD115 (Figures 11 and 16; Table 12).

Table 12. Anti-CD115 mAbs inhibit CSF-1 induced phosphorylation of CD115

mAb	IC ₅₀ (pM)
CCK-415A	570
CCK-416A	1350
CCK-423A	45

EXAMPLE 8

CONVERSION OF ALIVAMAB MOUSE ANTI-CD115 MABS TO FULLY HUMAN

[0117] The AlivaMab Mouse anti-CD115 mAbs are easily converted, expressed recombinantly and purified as fully-human antibodies of any isotype. The recombinant fully-human antibody retains all of the characteristics of the parental AlivaMab Mouse antibody. For example, the nucleotide sequences of the heavy and light chain variable region of CCK423A were transmitted to and synthesized into DNA by Lake Pharma (Belmont CA) and then, using vectors for recombinant expression in mammalian cells, the V_H cloned in-frame with coding sequences for human IgG1, IgG2, or IgG4 constant regions and the V_κ cloned in-frame with coding sequences for the human C_κ region. Vectors were then transformed into HEK293 cells for expression of recombinant fully human antibody. Fully human IgG1 κ , IgG2 κ and IgG4 κ mAb versions of CCK423A were purified from tissue culture supernatants using protein A (Figure 13).

EXAMPLE 9

AFFINITY OF FULLY HUMAN MABS

[0118] Affinity was determined for AlivaMab CCK423A as well as for the 3 human variants CCK423A-IgG1 κ , CCK423A-IgG2 κ , and CCK423A-IgG4 κ (Biosensor Tools, Salt Lake City, Utah). Binding kinetics were measured using a BioRad ProteOn XPR36 optical biosensor equipped with a GLM sensor chip. Purified mAbs were amine coupled to the GLM sensor chip. Hu CD115 (Sino Biological, China #10161-H08H) was tested in triplicate using a 3 fold dilution series starting at 10 nM. The processed data were fit using a 1:1 interaction model that includes a mass-transport parameter (Scrubber2, Canberra Australia). All 4 constructs were found to bind hu-CD115 with the same kinetics and affinity (Table 13).

Table 13.

mAb	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})	KD (nM)
AlivaMab CCK423A	1.3×10^7	1.5×10^{-2}	1.2
Human IgG1 κ	1.2×10^7	1.7×10^{-2}	1.2
Human IgG2 κ	1.3×10^7	1.4×10^{-2}	1.3
Human IgG4 κ	1.1×10^7	1.5×10^{-2}	1.3

EXAMPLE 10

CSF-1 BINDING NEUTRALIZATION WITH FULLY HUMAN MABS

[0119] AlivaMab Mouse CCK423A as well as the 3 human variants CCK423A-IgG1, CCK423A-IgG2, and CCK423A-IgG4 antibodies were tested for their ability to block binding of recombinant CSF-1 to recombinant CD115. ELISA plates were coated with recombinant hu-CD115 (Sino Biological, China #10161-H08H) at 0.5 ug/ml and blocked with Superblock (Thermo Scientific #37518). Wells were incubated for 15 min with anti-CD115 mAbs, then biotinylated Hu-CSF-1 (R&D Systems, #216-MC-005) (biotinylation using NHS-Peg4-biotin, Life Technologies, #21330) was added to a final concentration of 0.25 ug/ml for an additional 15'. After a 4X wash, CSF-1-biotin was detected using 1:10,000 SAV-poly HRP (Life Technologies, #N200). The fully human variants exhibited identical potency as the parental AlivaMab antibody (Figure 14 and Table 14).

Table 14.

mAb	IC ₅₀ (pM)
-----	-----------------------

AlivaMab CCK423A	191
Human IgG1 κ	146
Human IgG2 κ	253
Human IgG4 κ	133

EXAMPLE 11

INHIBITION OF CSF-1 INDUCED P-TYR WITH FULLY HUMAN MABS

[0120] AlivaMab Mouse CCK423A as well as the 3 human variants CCK423A-IgG1, CCK423A-IgG2, and CCK423A-IgG4 were tested for their ability to block hu-CSF1 induced phosphorylation of native CD115. OCI-AML5 cells (DSMZ, # ACC-247) were serum starved (1% FBS) overnight, then harvested and washed twice in PBS with 0.1% BSA. 250,000 cells were plated per well into a 96-well v-bottom polypropylene plate. Anti-CD115 mAbs were added in a dilution series for 15 min while incubating the plate on ice. Hu-CSF-1 (R&D Systems, #216-MC-005) was added to each well at a final concentration of 100 ng/ml and incubated for 30 min on ice. Cells were then spun down at 1500 RPM for 5 min at 4°C and supernatant was removed. Cells were then resuspended in lysis buffer (Cell Signaling, #9803 with 1X HALT protease inhibitors, Pierce, #78430) and incubated on ice for 15 min. Lysates were then measured for tyrosine phosphorylated-CD115 using a p-MCSFR validated duoset assay (R&D Systems #CYC3268E) and detected using Supersignal Pico ELISA Substrate (Pierce, #37069). The fully human variants exhibited identical potency as the parental AlivaMab antibody (Figure 15 and Table 15).

Table 15.

mAb	IC ₅₀ (pM)
AlivaMab CCK423A	58
Human IgG1 κ	68
Human IgG2 κ	59
Human IgG4 κ	60

EXAMPLE 12

INHIBITION OF P-TYR FORMATION ON CD115 INDUCED BY IL-34

[0121] Anti-CD115 antibodies of the invention were found to neutralize P-TYR formation on CD115 induced by interleukin-34 (IL-34). Some antibodies that block CSF-1 induced P-TYR formation on CD115 are found to also block IL-34 induced P-TYR formation on CD115. Other antibodies are found that block only CSF-1 induced P-TYR formation on

CD115 and do not block IL-34 induced P-TYR formation on CD115. Antibodies are tested for their ability to block IL-34 induced phosphorylation of native CD115.

[0122] In an example assay, SR cells or other CD115+ IL-34 responsive cell line(s) are serum starved (1% FBS) overnight, then harvested and washed twice in PBS with 0.1% BSA. Cells are plated into a 96-well v-bottom polypropylene plate. Anti-CD115 antibodies, either in hybridoma supernatants, purified antibody, or in purified fully-human recombinant antibody format, are added for 15 min while incubating the plate on ice. Human IL-34 is added to each well at a final concentration sufficient and necessary to trigger P-TYR formation on CD115 and incubated for 30' on ice. Cells were then spun down at 1500 RPM for 5' at 4°C and supernatant was removed. Cells were then resuspended in lysis buffer (Cell Signaling, #9803 with 1X HALT protease inhibitors, Pierce, #78430) and incubated on ice for 15 min. Lysates were then measured for tyrosine phosphorylated CD115 using a p-MCSFR validated DUOSET assay (R&D Systems #CYC3268E) and detected using Supersignal Pico ELISA Substrate (Pierce, #37069).

EXAMPLE 13

NEUTRALIZATION OF IL-34 BINDING TO CD115

[0123] AlivaMab Mouse anti-CD115 antibodies block binding of IL-34 to CD115. For example, anti-CD115 antibodies were tested for their ability to block binding of recombinant human IL-34 to recombinant CD115. ELISA plates were coated with recombinant hu-CD115 (Sino Biological, China #10161-H08H) at 0.5 ug/ml and blocked with Superblock (Thermo Scientific #37518). Wells were incubated for 15 min with anti-CD115 mAbs, then biotinylated Hu-IL-34 (biotinylation using NHS-Peg4-biotin, Life Technologies, #21330) is added for an additional 15 minutes. After a 4X wash, HU-IL-34-biotin was detected using 1:10,000 SAV-poly HRP (Life Technologies, #N200). Other AlivaMab Mouse anti-CD115 mAbs are also shown to exhibit various abilities and IC50 values in blocking HU-IL-34 binding to CD115.

EXAMPLE 14

NEUTRALIZATION OF P-TYR FORMATION ON CD115 WITHOUT NEUTRALIZATION OF BINDING OF CSF-1 AND IL-34

[0124] AlivaMab Mouse anti-CD115 mAbs were found that neutralize p-Tyr formation in cells exposed to either CSF-1 or IL-34. However, these mAbs still allow

binding of CSF-1 and IL-34 to CD115. This set of mAbs block p-Tyr formation through inhibition of dimerization of CD115.

EXAMPLE 15

AlivaMab Mouse ANTI-CD115 MABS AND THEIR FULLY HUMAN DERVATIVES BIND TO AND NEUTRALIZE CD115 FROM CYNOMOLGUS MONKEY

[0125] CD115 is cloned and expressed from cynomolgus monkey using standard molecular biological techniques. The recombinant CD115 may be tagged (histidine, Fc) to support efficient purification. The recombinant cynomolgus CD115 may also be transiently or stably expressed on cell lines. The AlivaMab Mouse anti-CD115 mAbs and their human variants are shown to bind to cynomolgus monkey CD115. The AlivaMab Mouse anti-CD115 mAbs and their human variants are shown to neutralize cynomolgus monkey CD115 in assays as described above for human CD115.

[0126] The various embodiments described above can be combined to provide further embodiments. All of the U.S. patents, U.S. patent application publications, U.S. patent application, foreign patents, foreign patent application and non-patent publications referred to in this specification and/or listed in the Application Data Sheet are incorporated herein by reference, in their entirety. Aspects of the embodiments can be modified, if necessary to employ concepts of the various patents, application and publications to provide yet further embodiments.

[0127] These and other changes can be made to the embodiments in light of the above-detailed description. In general, in the following claims, the terms used should not be construed to limit the claims to the specific embodiments disclosed in the specification and the claims, but should be construed to include all possible embodiments along with the full scope of equivalents to which such claims are entitled. Accordingly, the claims are not limited by the disclosure.

[0128] Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integers or steps.

[0129] The reference in this specification to any prior publication (or information derived from it), or to any matter which is known, is not, and should not be taken as an acknowledgment or admission or any form of suggestion that that prior publication (or information derived from it) or known matter forms part of the common general knowledge in the field of endeavour to which this specification relates.

CLAIMS

1. An isolated anti-CD115 antibody, or an antigen-binding fragment thereof, comprising i) a heavy chain variable region (VH) comprising a heavy chain variable complementarity determining region 1 (VHCDR1) comprising SEQ ID NO: 450, a VHCDR2 comprising SEQ ID NO: 882, and a VHCDR3 comprising SEQ ID NO: 1314; and ii) a light chain variable region (VL) comprising a VLCDR1 comprising SEQ ID NO: 666, a VLCDR2 comprising SEQ ID NO: 1098, and a VLCDR3 comprising SEQ ID NO: 1530.
2. The antibody, or antigen-binding fragment thereof, of claim 1, wherein the VH comprises SEQ ID NO:123.
3. The antibody, or antigen-binding fragment thereof, of claim 1 or claim 2, wherein the VL comprises SEQ ID NO:339.
4. The antibody, or antigen-binding fragment thereof, of claim 1, wherein the antibody is human.
5. The antibody, or antigen-binding fragment thereof, of claim 1, wherein the antibody is chimeric.
6. The antibody, or antigen-binding fragment thereof, of any one of claims 1-5, wherein the antibody is selected from a single-variable domain antibody, single chain antibody, a scFv, a bispecific antibody, a multi-specific antibody, a Fab, a F(ab')₂, and a whole antibody.
7. A recombinant polynucleotide encoding the antibody, or antigen-binding fragment thereof, of any one of claims 1-6.
8. An expression vector comprising the recombinant polynucleotide of claim 7.

9. An isolated host cell comprising the expression vector of claim 8.
10. A composition comprising the antibody, or antigen-binding fragment thereof, of any one of claims 1-6 and a physiologically acceptable carrier.
11. A method of treating cancer in a subject in need thereof, the method comprising administering to the subject a therapeutically effective amount of the antibody, or antigen-binding fragment thereof, of any one of claims 1-6 or the composition of claim 10, wherein the cancer is associated with overexpression of CD115, CSF-1 or IL-34.
12. Use of the antibody, or an antigen-binding fragment thereof, of any one of claims 1-6 or the composition of claim 10 in the manufacture of a medicament for treating cancer, wherein the cancer is associated with overexpression of CD115, CSF-1 or IL-34.

Pilot p-MCSFR ELISA - SR cells

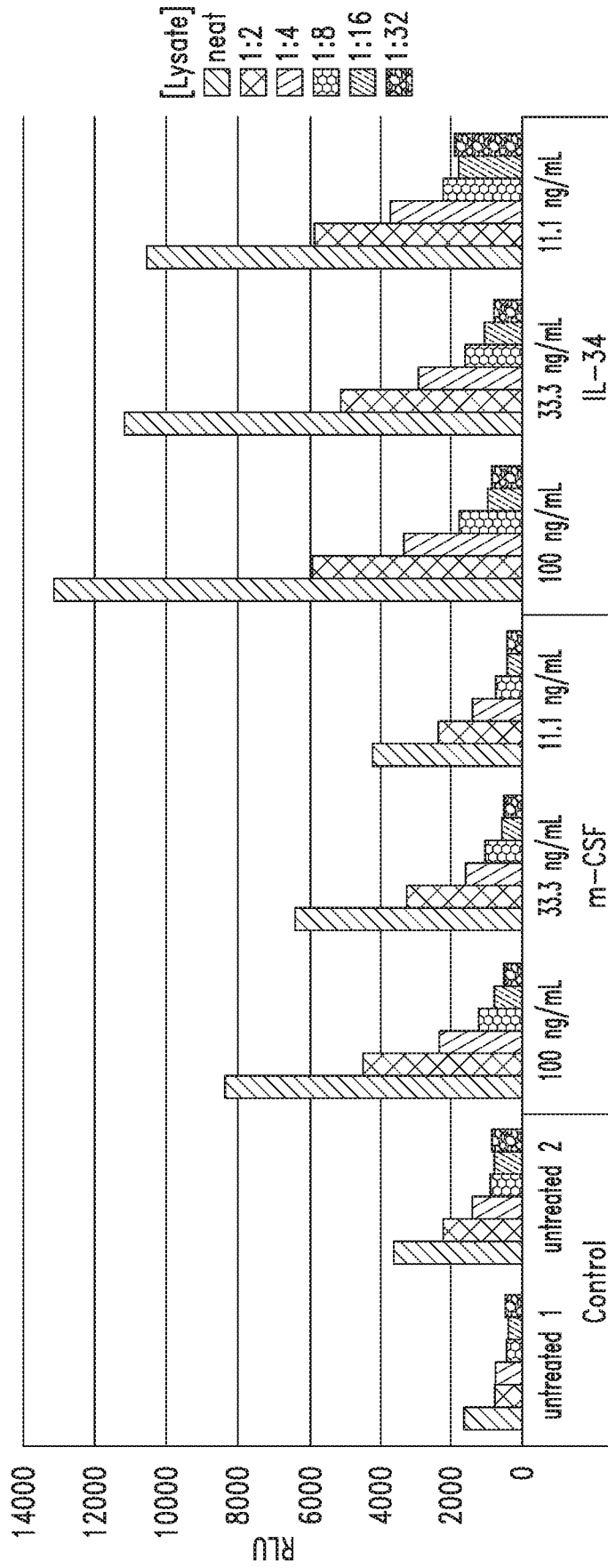


FIG. 1

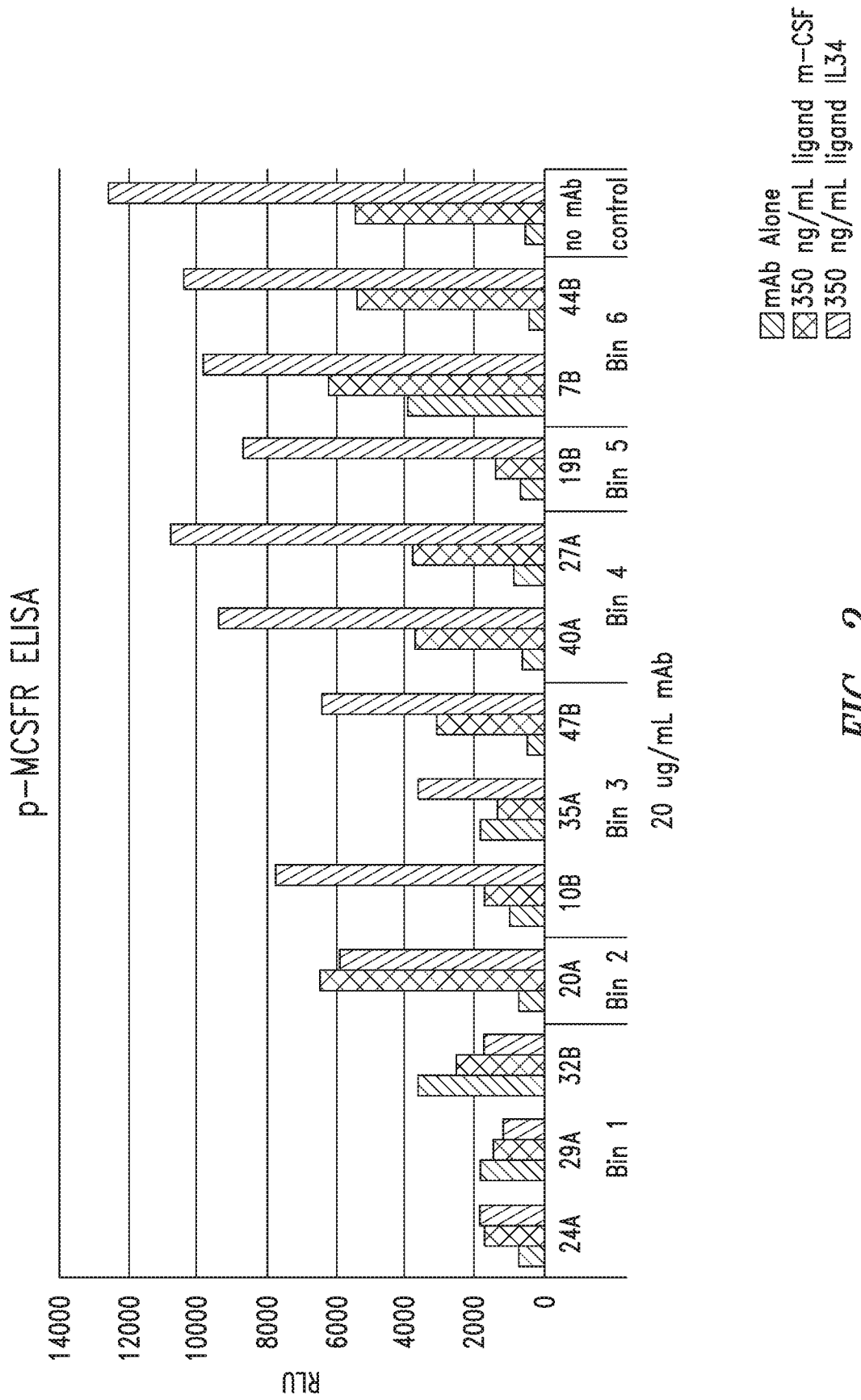
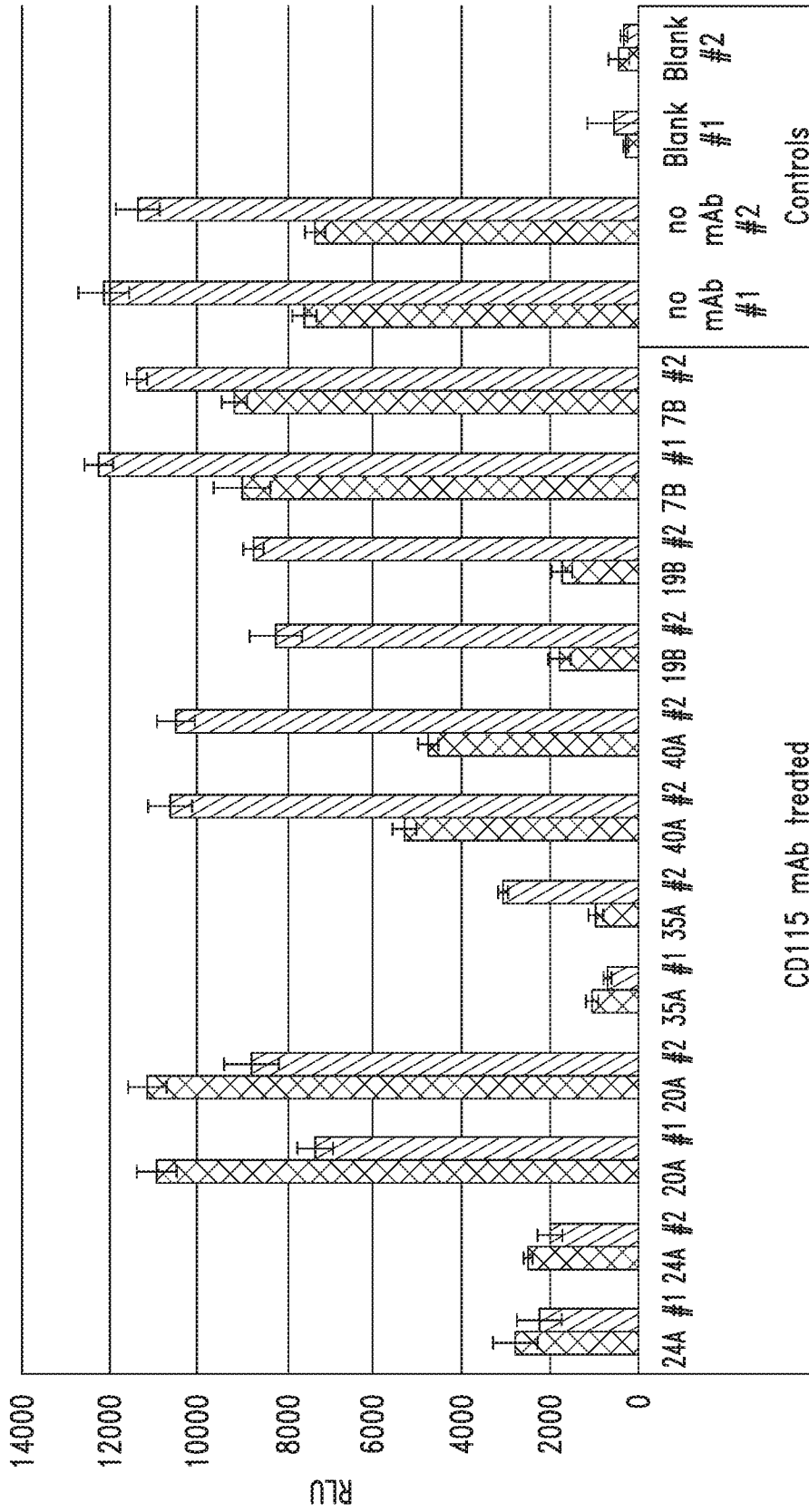


FIG. 2

p-MCSFR ELISA - mAb inhibition of CD115 phosphorylation



▨ Treated with m-CSF
▧ Treated with IL34

CD115 mAb treated

FIG. 3

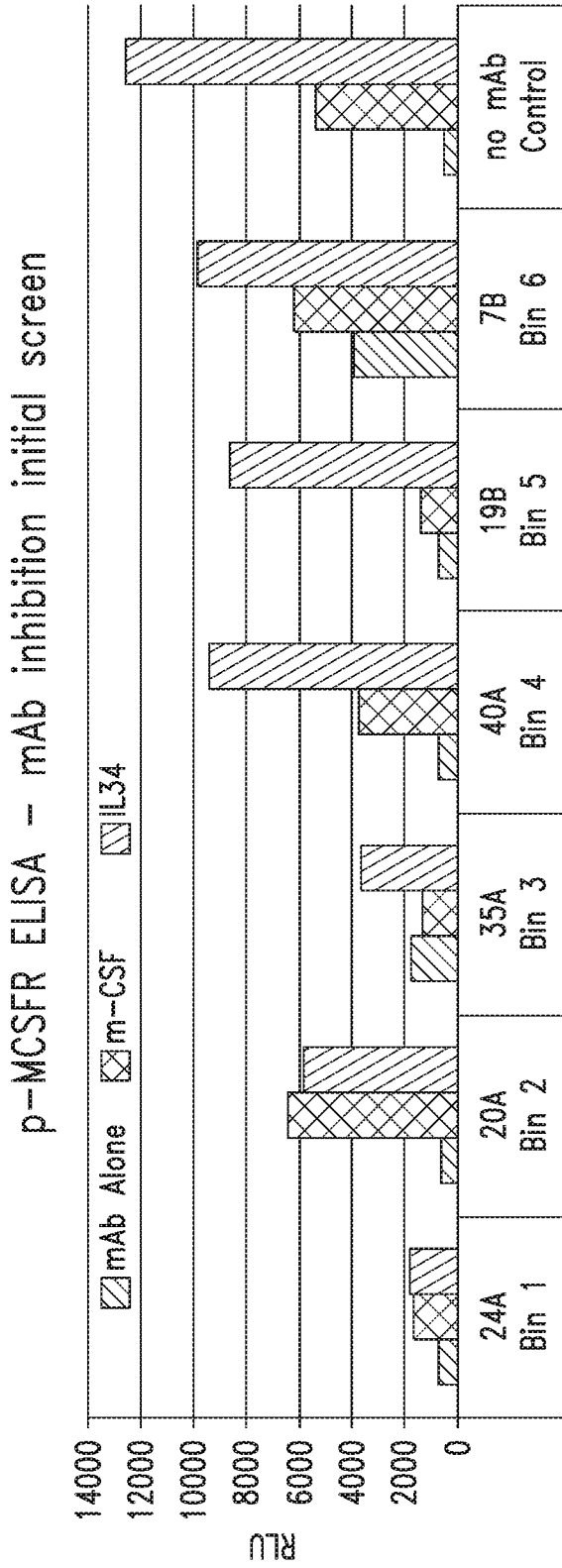


FIG. 4A p-MCSFR ELISA - mAb inhibition confirmation screen

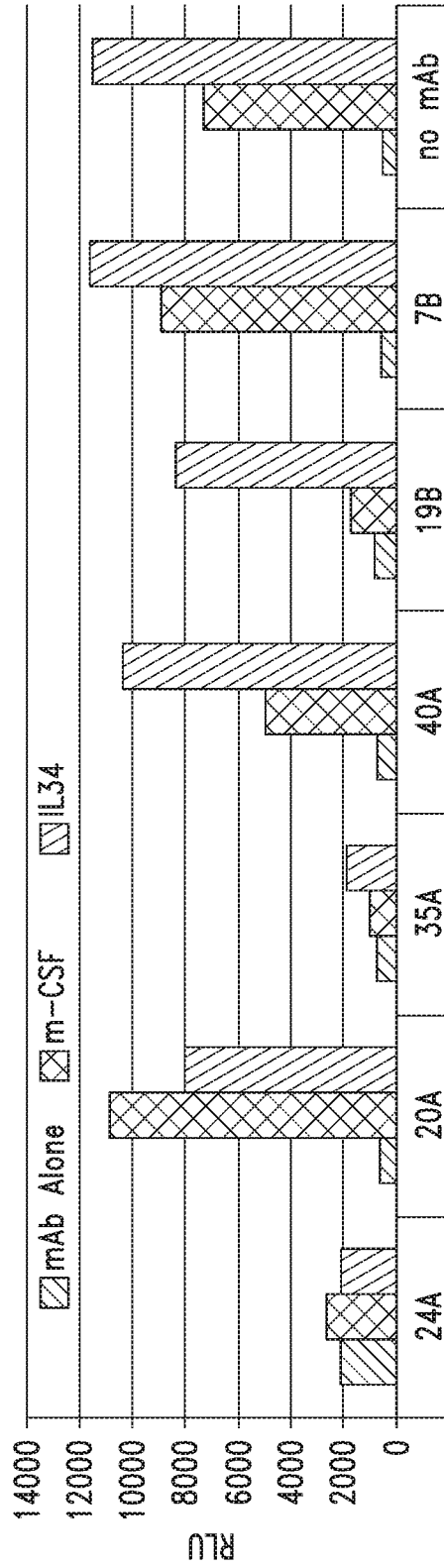


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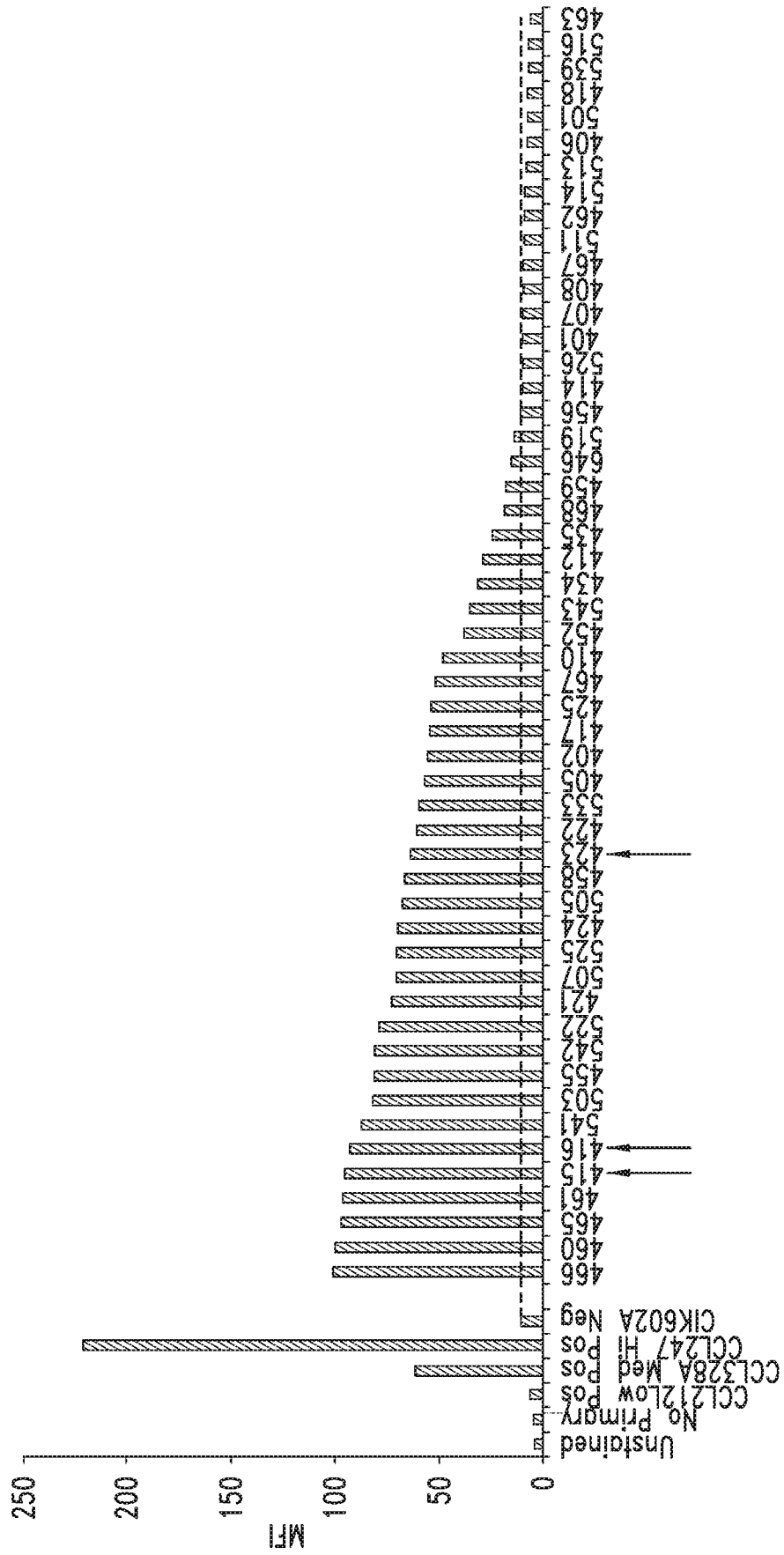


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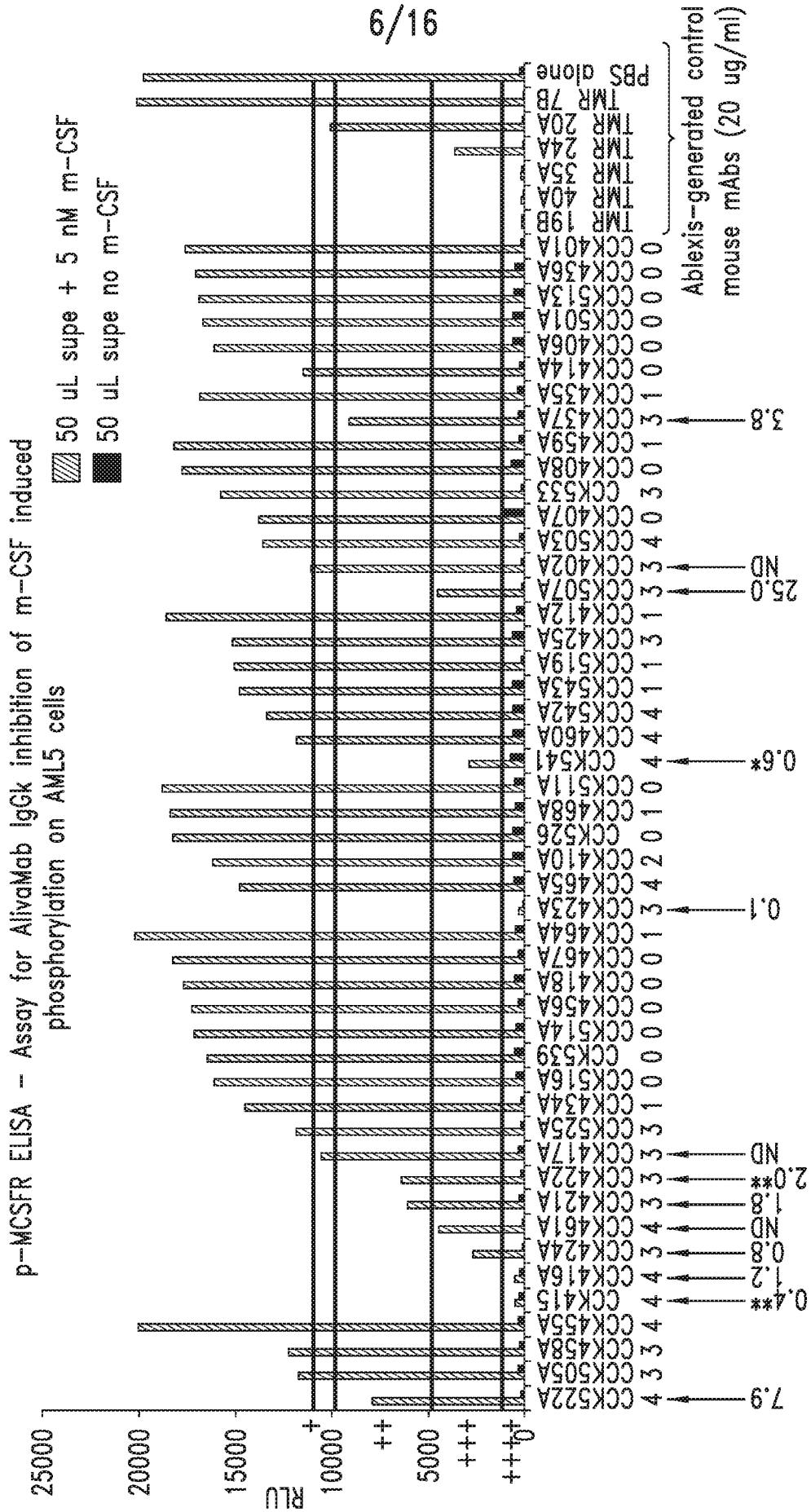


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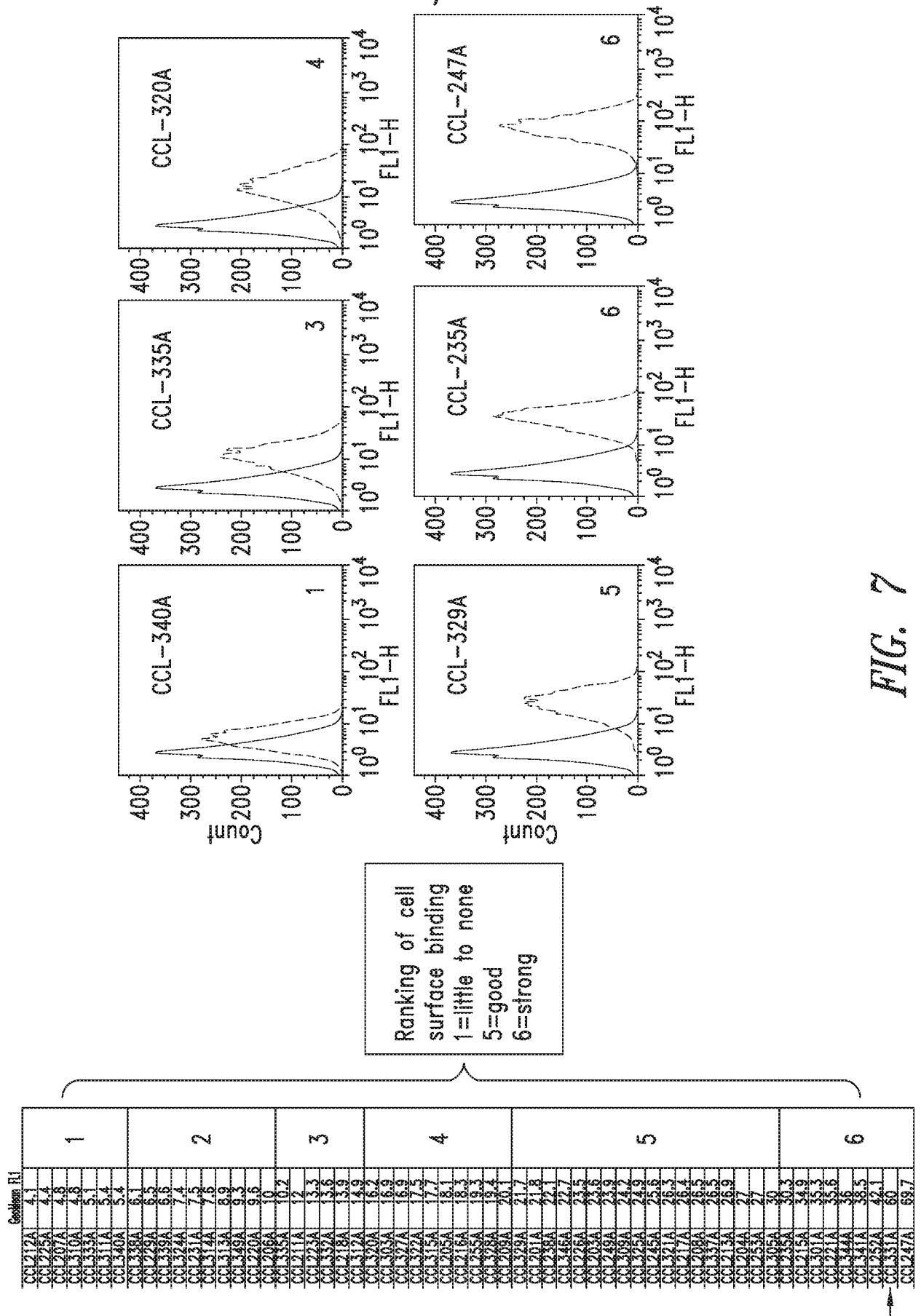


FIG. 7

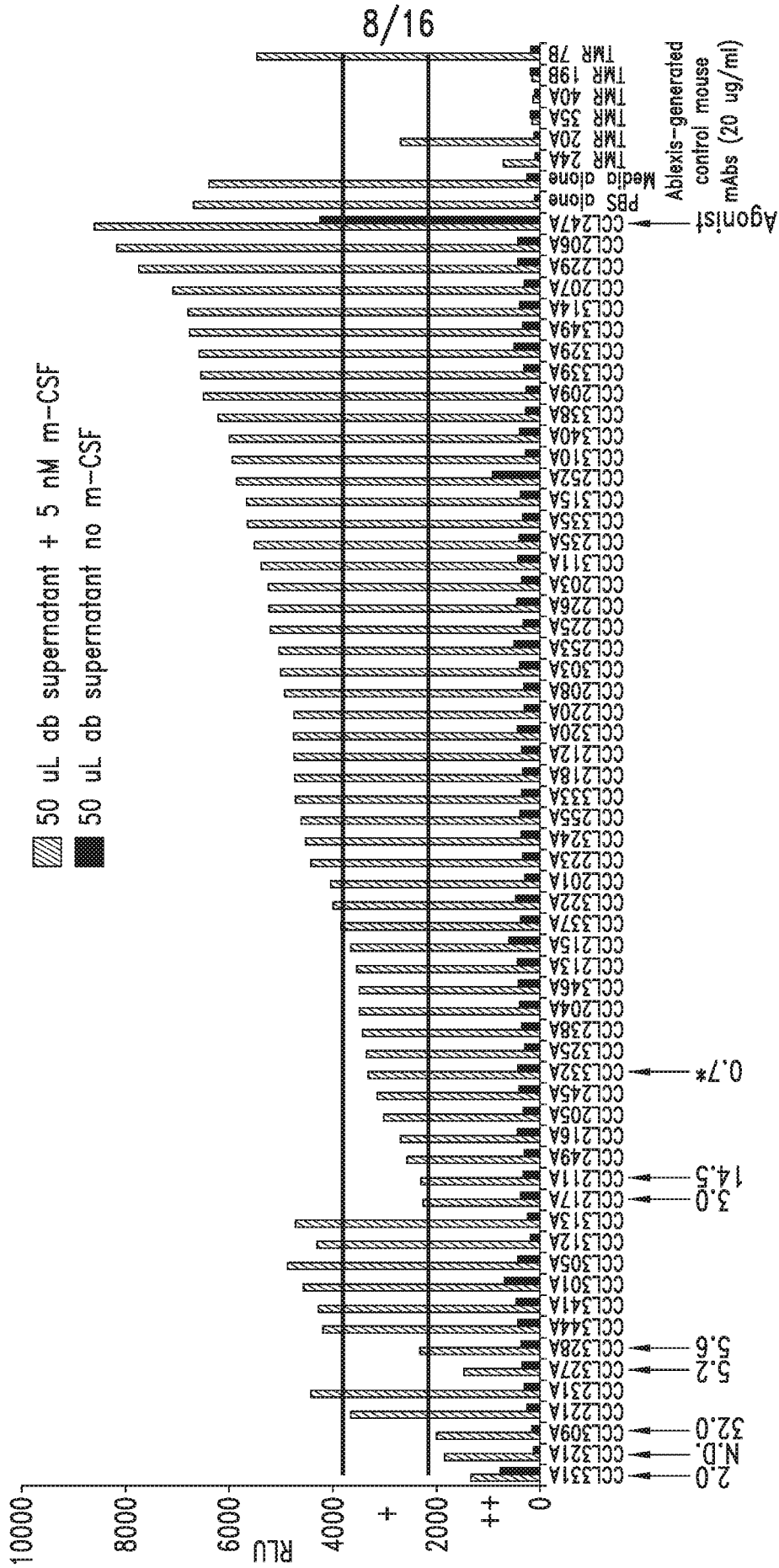


FIG. 8

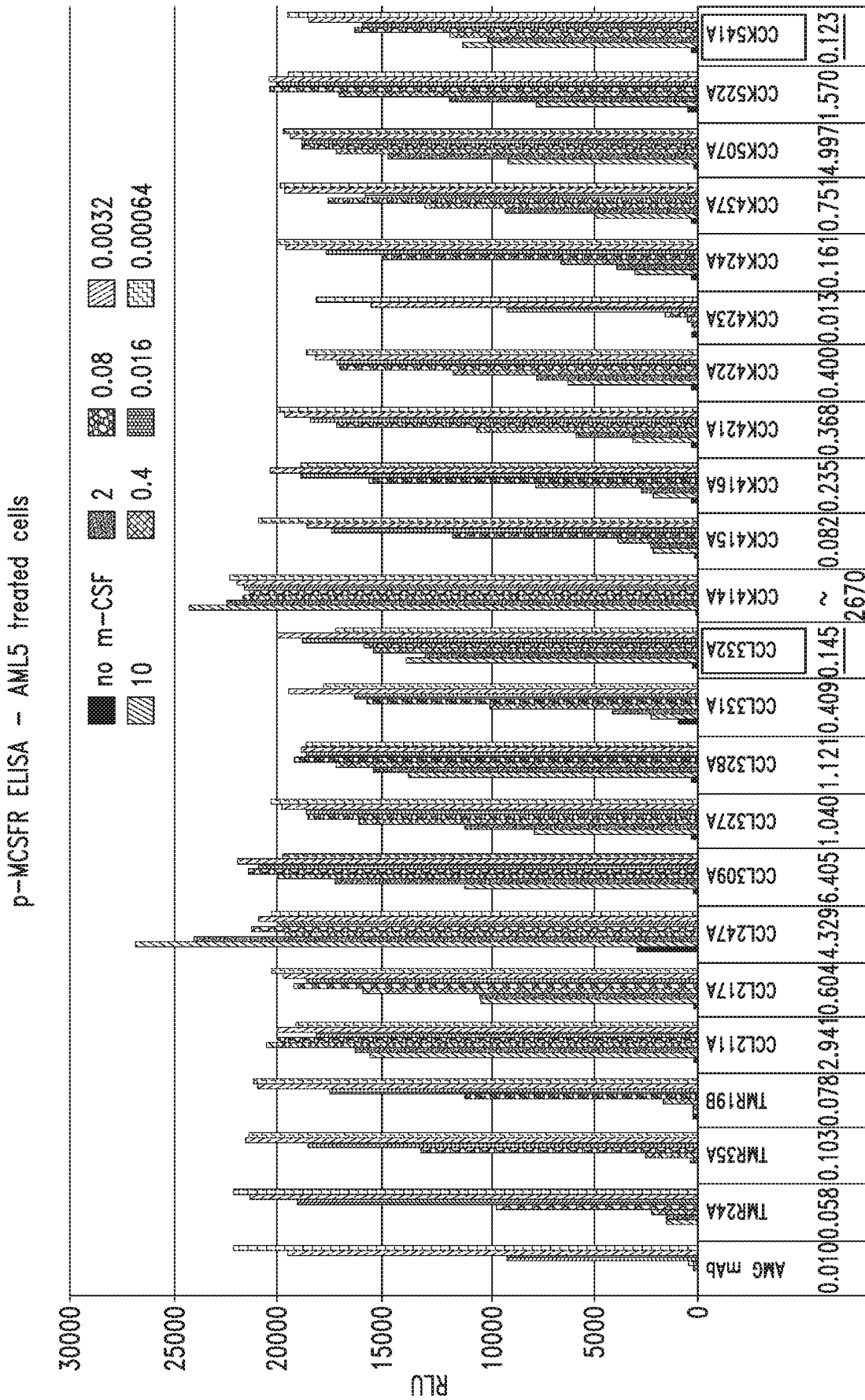


FIG. 9

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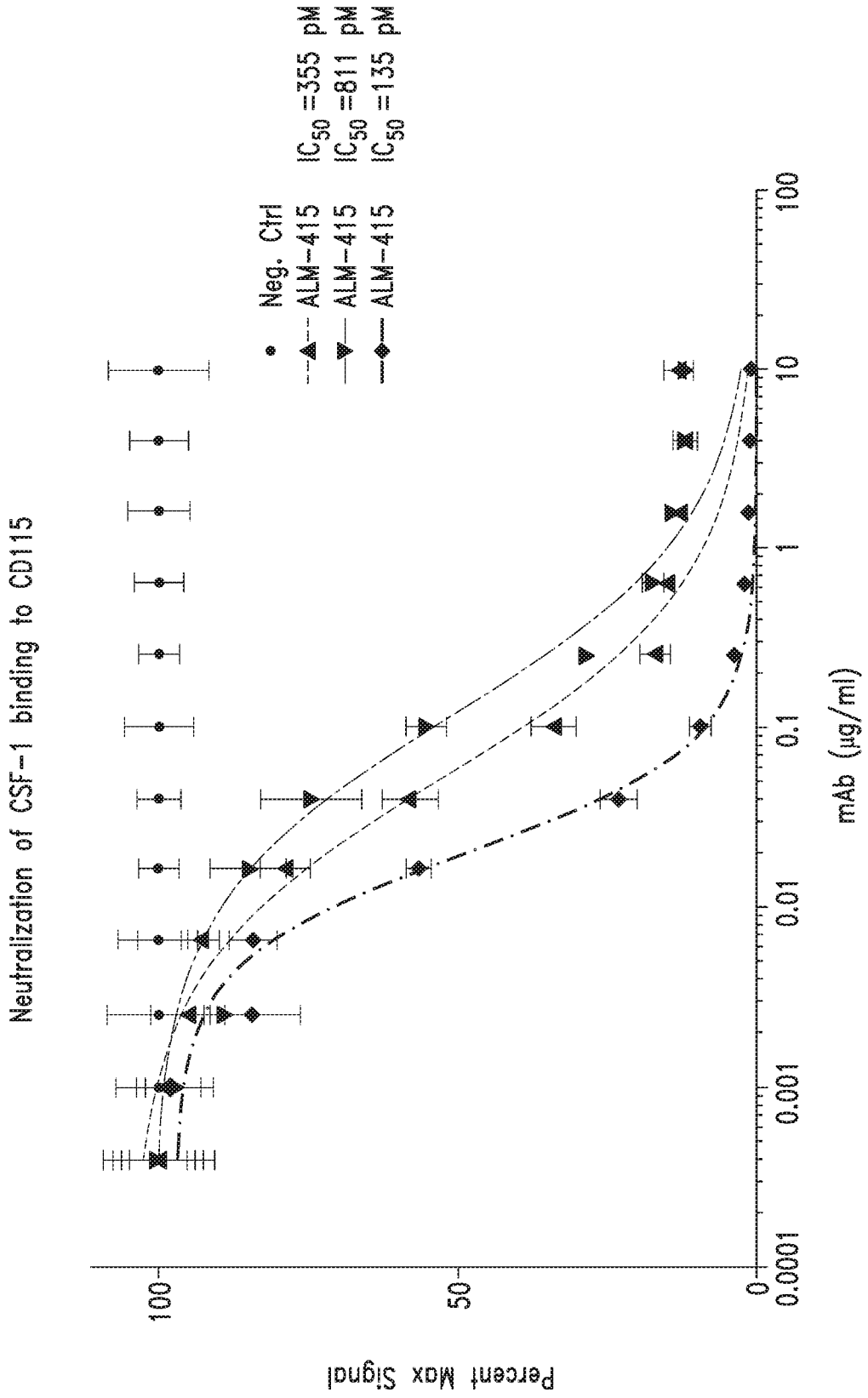


FIG. 10

11/16

Neutralization of CSF-1 induced phosphorylation of CD115

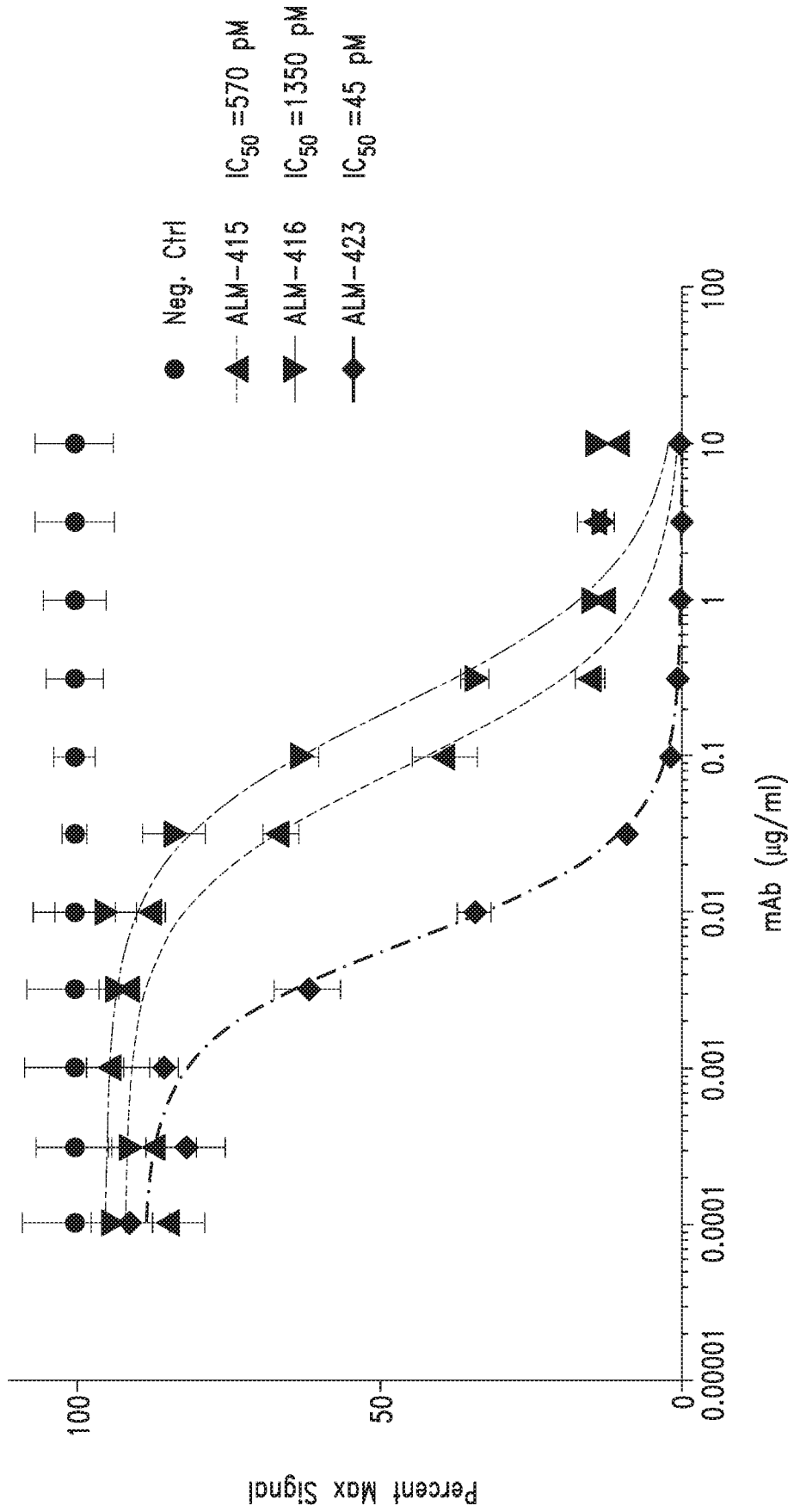


FIG. 11

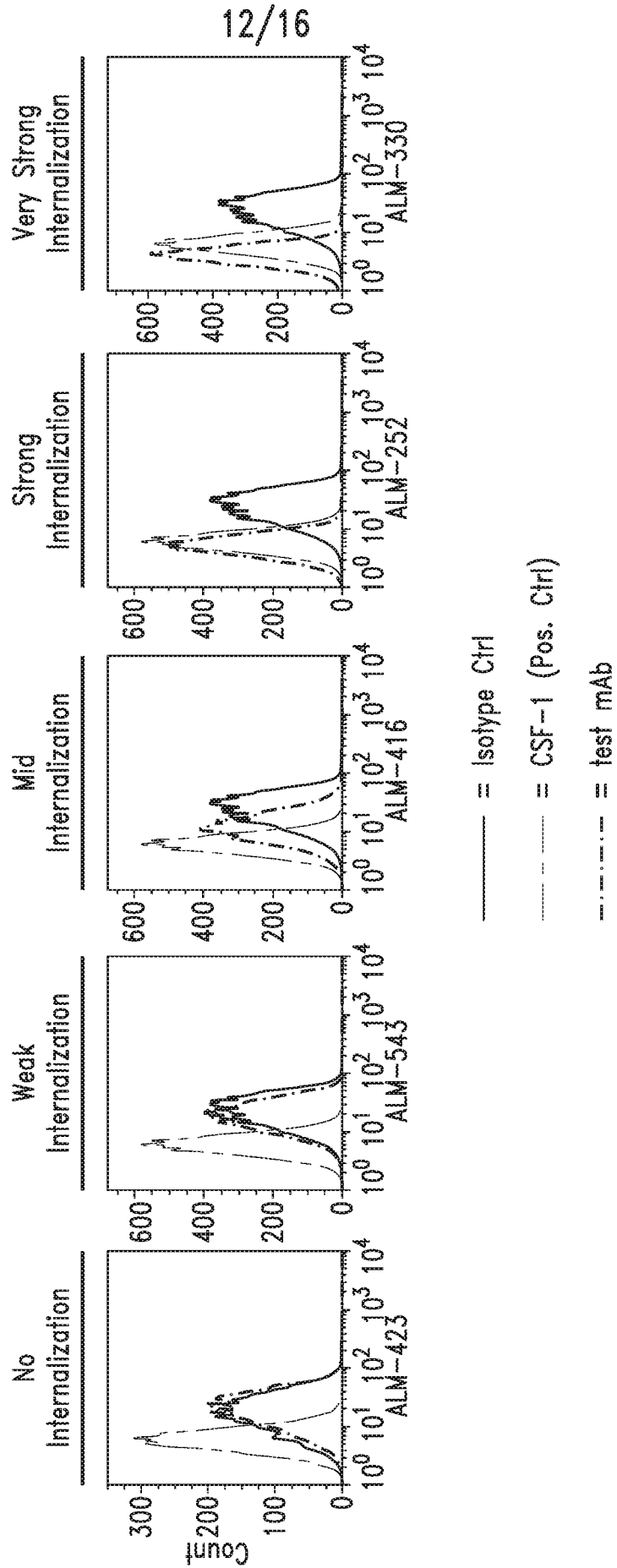


FIG. 12

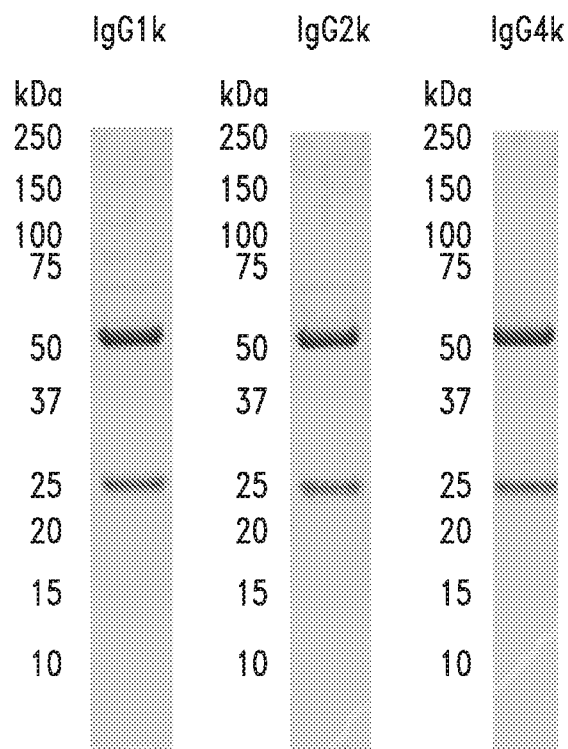


FIG. 13

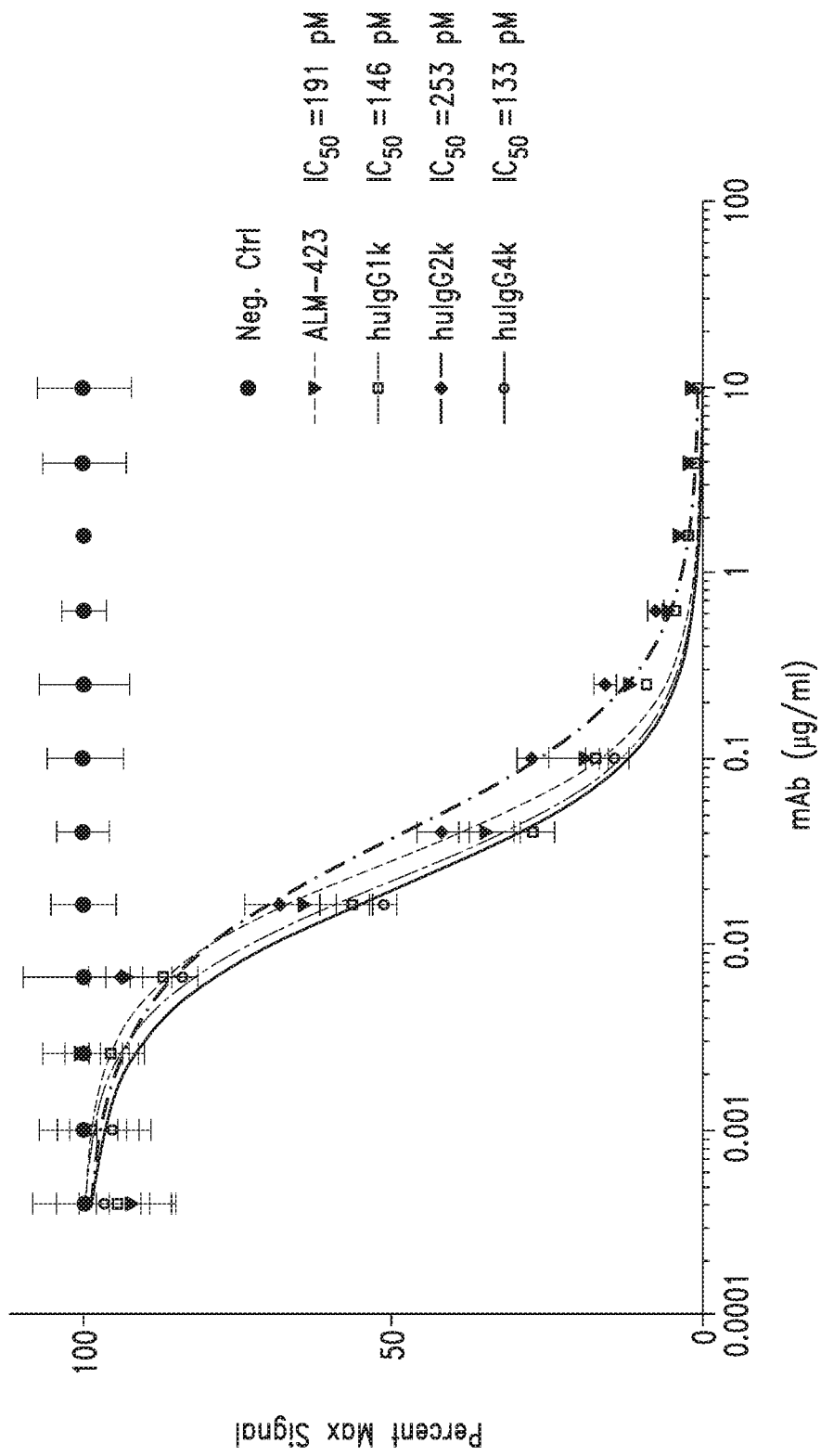


FIG. 14

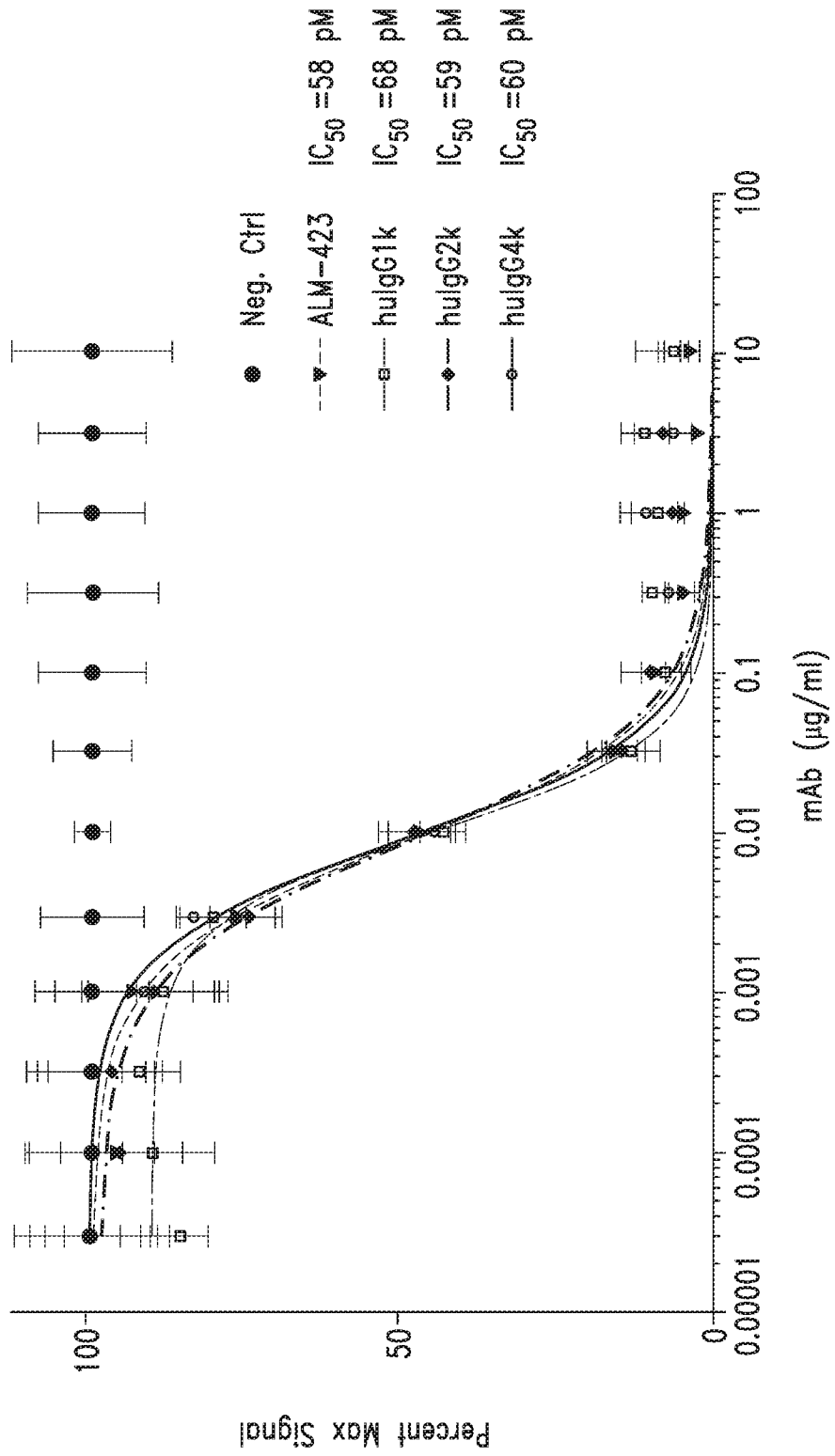


FIG. 15

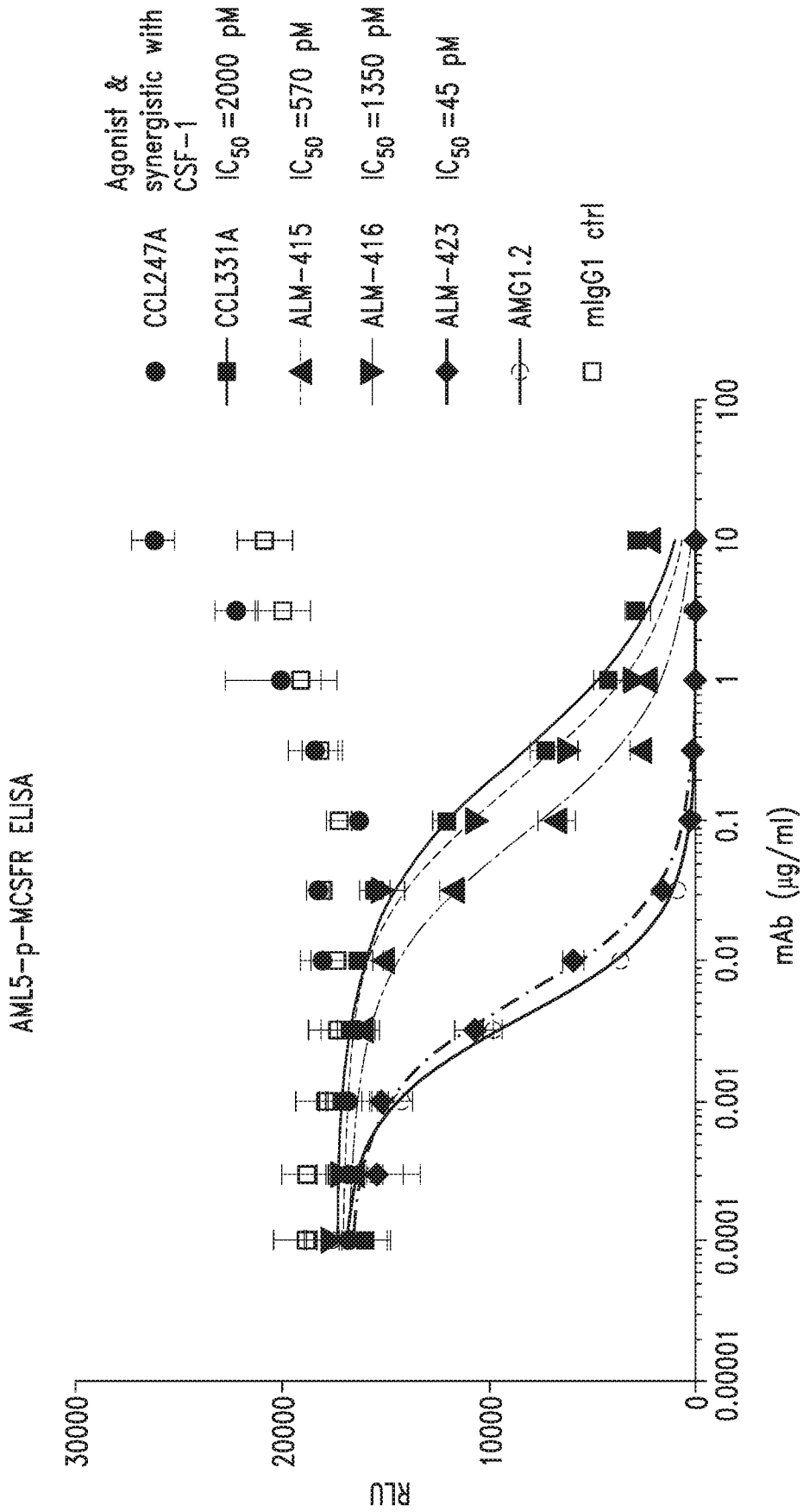


FIG. 16

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<211> 379
<212> DNA
<213> Homo sapiens

<400> 82
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 <213> Homo sapiens

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<212> DNA
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 aagctgagtt ctgtgaccgc cgccgacacg gccgtgtatt actgtgagag agagaggctg 300
 gggatgttct ttgactactg gggccagga accctggtca ccgtctcctc ag 352

<210> 106
 <211> 352
 <212> DNA
 <213> Homo sapiens

<400> 106
 caggtgcagc tgcaggagtc gggcccagga ctggtgaagc cttcggagac cctgtccctc 60
 acctgactg tctctggtgg ctccatcagt agttactact ggagctggat ccggcagccc 120
 gccgggaagg gactggagtg gattgggcgt atctatacca gtgggagcac caactacaac 180
 ccctccctca agagtcgagt caccatgtca gtagacacgt ccaagaacca gttctccctg 240
 aagctgagtt ctgtgaccgc cgccgacacg gccgtgtatt actgtgagag agagaggctg 300
 gggatgttct ttgactactg gggccagga accctggtca ccgtctcctc ag 352

<210> 107
 <211> 352
 <212> DNA
 <213> Homo sapiens

<400> 107
 caggtgcagc tacaggagtc gggcccagga ctggtgaagc cttcggagac cctgtccctc 60
 acctgactg tctctggtgg ctccatcagt agttactact ggagctggat ccggcagccc 120
 gccgggaagg gactggagtg gattggacgt ttctatacca gtgggagcac cagctgcaac 180
 ccctccctca agagtcgagt caccatgtca gtagacacgt ccaagaacca gttctccctg 240
 aagctgagct ctgtgaccgc cgccgacacg gccgtgtatt actgtgagag agaggggata 300
 ctggggatatt ttgactattg gggccagga accctggtca ccgtctcctc ag 352

<210> 108
 <211> 352
 <212> DNA
 <213> Homo sapiens

<400> 108
 caggtgcagc tgcaggagtc gggcccagga ctggtgaagc cttcggagac cctgtccctc 60
 acctgactg tctctggtgg ctccatcagt atttactact ggaactggat ccggcagccc 120

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ccaggggaagg gactggagtg gattgggtat atctattaca ttgggagcac caactacaac 180
 ccctccctca agagtcgagt caccatatca gtagacacgt ccaagaacca gttctccctg 240
 aagctgagtt ctgtgaccgc tgcggacacg gccgtgtatt actgtgcgcg aggaaggcta 300
 acttatgctt ttgatatctg gggccaaggg acaatggtca ccgtctcttc ag 352

<210> 109
 <211> 122
 <212> PRT
 <213> Homo sapiens

<400> 109

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Ala
 20 25 30

Trp Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

Gly Arg Ile Lys Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Ala Ala
 50 55 60

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

Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120

<210> 110
 <211> 118
 <212> PRT
 <213> Homo sapiens

<400> 110

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

Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
 35 40 45

Gly Trp Ile Asn Pro Ile Ser Gly Gly Thr Asn Tyr Ala Gln Lys Phe
 50 55 60

Gln Gly 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 Thr Ala Val Tyr Tyr Cys
 85 90 95

Ala Arg Gln Asp Asn Trp Asn Tyr Phe Asp Tyr Trp Gly Gln Gly Thr
 100 105 110

Leu Val Thr Val Ser Ser
 115

<210> 111
 <211> 123
 <212> PRT
 <213> Homo sapiens

<400> 111

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Ala
 20 25 30

Trp Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Asp Trp Val
 35 40 45

Gly Arg Val Lys Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Ala Ala
 50 55 60

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

Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120

<210> 112
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 112

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly His Ile Phe Ser Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Ser Gln Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Leu Cys Ala
85 90 95

Arg Gly Gly Val Thr Trp Phe Asp Pro Trp Gly Gln Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 113
<211> 116
<212> PRT
<213> Homo sapiens

<400> 113

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr His Pro Ser Phe Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Thr Gln Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Leu Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gly Gly Ile Thr Trp Phe Asp Pro Trp Gly Gln Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 114
 <211> 120
 <212> PRT
 <213> Homo sapiens

<400> 114

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

Ser Leu Arg Leu Ser Cys Thr Ala Ser Gly Phe Thr Phe Gly Asp Tyr
 20 25 30

Ala Met Ser Trp Phe Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

Gly Phe Ile Arg Ser Lys Ala Tyr Gly Gly Thr Thr Glu Tyr Ala Ala
 50 55 60

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

Ala Tyr Leu Gln Met Asn Ser Leu Lys Ile Glu Asp Thr Ala Val His
 85 90 95

Tyr Cys Thr Arg Glu Gly Ser Phe Gly Ala Leu Asp Ile Trp Gly Gln
 100 105 110

Gly Ile Met Val Thr Val Ser Ser
 115 120

<210> 115
 <211> 119
 <212> PRT
 <213> Homo sapiens

<400> 115

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 Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
 35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Asn Ala Gln Lys Phe
 50 55 60

Gln Gly 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

Al a Arg Gly Gly Tyr Ser Gly Pro Tyr Phe Asp Tyr Trp Gly Gl n Gly
 100 105 110

Thr Leu Val Thr Val Ser Ser
 115

<210> 116
 <211> 122
 <212> PRT
 <213> Homo sapi ens

<400> 116

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gly Leu Val Lys Pro Gly Gly
 1 5 10 15

Ser Leu Arg Leu Ser Cys Al a Al a Ser Gly Phe Thr Phe Ser Asn Al a
 20 25 30

Trp Met Ser Trp Val Arg Gl n Al a Pro Gly Lys Gly Leu Gl u Trp Val
 35 40 45

Gly Arg Ile Lys Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Al a Al a
 50 55 60

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

Leu Tyr Leu Gl n Met Asn Ser Leu Lys Thr Gl u Asp Thr Al a Val Hi s
 85 90 95

Tyr Cys Thr Thr Gly Asp Tyr Ser Tyr Thr Asp Al a Phe Asp Ile Trp
 100 105 110

Gly Gl n Gly Thr Met Val Thr Val Ser Ser
 115 120

<210> 117
 <211> 112
 <212> PRT
 <213> Homo sapi ens

<400> 117

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
 1 5 10 15

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Ile Ser Ser Ser
 20 25 30

Tyr Trp Thr Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
 35 40 45

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Gly Tyr Ile His Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gly Phe Asp Tyr Trp Gly Gl n Gly Thr Leu Val Thr Val Ser Ser
100 105 110

<210> 118
<211> 124
<212> PRT
<213> Homo sapi ens

<400> 118

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Asp Leu Val Gl n Pro Gly Gly
1 5 10 15

Ser Leu Arg Leu Ser Cys Ala Asp Ser Arg Phe Thr Phe Ser Ser Tyr
20 25 30

Ser Met Asn Trp Val Arg Gl n Ala Pro Gly Lys Gly Leu Gl u Trp Val
35 40 45

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

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

Leu Gl n Met Asn Ser Leu Arg Asn Gl u Asp Thr Ala Val Tyr Tyr Cys
85 90 95

Ala Arg Asp Phe Pro His Asp Tyr Gly Tyr Tyr Pro His Tyr Phe Asp
100 105 110

Tyr Trp Gly Gl n Gly Thr Leu Val Thr Val Ser Ser
115 120

<210> 119
<211> 118
<212> PRT
<213> Homo sapi ens

<400> 119

Gl n Val Thr Leu Arg Gl u Ser Gly Pro Ala Leu Val Lys Pro Thr Gl n
1 5 10 15

Thr Leu Thr Leu Thr Cys Thr Phe Ser Gly Phe Ser Leu Ser Thr Ser

Gly Met Cys Val Ser Trp Ile Arg Gln Pro Pro Gly Lys Ala Leu Glu
 35 40 45

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

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

Val Leu Thr Met Thr Asn Met Asp Pro Val Asp Thr Ala Thr Tyr Tyr
 85 90 95

Cys Ala Arg Thr Gln Leu Gly Ile Ala Asp Tyr Trp Gly Gln Gly Thr
 100 105 110

Leu Val Thr Val Ser Ser
 115

<210> 120
 <211> 120
 <212> PRT
 <213> Homo sapiens

<400> 120

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Thr Thr Asn Phe Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Thr Leu Ser Gly Thr Asn Trp Gly Ser Pro Phe Asp Tyr Trp Gly Gln
 100 105 110

Gly Thr Leu Val Thr Val Ser Ser
 115 120

<210> 121
 <211> 125

<212> PRT
 <213> Homo sapiens

<400> 121

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

<210> 122
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 122

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

Arg Gly Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
 100 105 110

<210> 123
 <211> 119
 <212> PRT
 <213> Homo sapiens

<400> 123

Gln Val Gln Leu Glu Gln Ser Gly Ala Glu Val Lys Arg Pro Gly Ala
 1 5 10 15

Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
 35 40 45

Gly Trp Ile Asn Pro Asn Arg Gly Asp Thr Asn Tyr Ala Gln Lys Phe
 50 55 60

Gln Gly 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 Asp Leu Glu Gly Gly Gly Pro Phe Asp Tyr Trp Gly Gln Gly
 100 105 110

Thr Leu Val Thr Val Ser Ser
 115

<210> 124
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 124

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Val Leu Glu Trp Ile
 35 40 45

Gly Tyr Phe Ser Tyr Asn Gly Asn Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

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Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Thr
85 90 95

Arg Gly Met Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser
100 105 110

<210> 125
<211> 117
<212> PRT
<213> Homo sapiens
<400> 125

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Asp Arg Thr Phe Tyr Ala Phe Asp Met Trp Gly Gln Gly Thr Met
100 105 110

Val Thr Val Ser Ser
115

<210> 126
<211> 119
<212> PRT
<213> Homo sapiens
<400> 126

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 Ser Ser Tyr
20 25 30

Asp Met His Trp Val Arg Gln Ala Thr Gly Lys Gly Leu Glu Trp Val
35 40 45

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

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

Gln Met Asn Ser Leu Arg Ala Gly Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Gly His Ile Val Val Val Thr Ala Met Asp Tyr Trp Gly Gln Gly
 100 105 110

Thr Leu Val Thr Val Ser Ser
 115

<210> 127
 <211> 115
 <212> PRT
 <213> Homo sapiens

<400> 127

Gln Val Gln Leu Gln Gln Trp Gly Ala Gly Leu Leu Lys Pro Ser Glu
 1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Glu Ile Asn His Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Arg Asn Gln Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Ala Asp Ala Phe Asp Ile Trp Gly Gln Gly Thr Met Val Thr
 100 105 110

Val Ser Ser
 115

<210> 128
 <211> 122
 <212> PRT
 <213> Homo sapiens

<400> 128

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

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Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Ala
20 25 30

Trp Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

Gly Arg Ile Lys Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Ala Ala
50 55 60

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

Gly Gln Gly Thr Met Val Thr Val Ser Ser
115 120

<210> 129
<211> 115
<212> PRT
<213> Homo sapiens
<400> 129

Gln Val Gln Leu Gln Gln Trp Gly Ala Gly Leu Leu Lys Pro Ser Glu
1 5 10 15

Thr Leu Ser Leu Thr Cys Ala Val Tyr Gly Gly Ser Phe Ser Ala Tyr
20 25 30

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Glu Ile Asn His Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Ala Asp Ala Phe Asp Ile Trp Gly Gln Gly Thr Met Val Thr
100 105 110

Val Ser Ser
115

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<210> 130
 <211> 119
 <212> PRT
 <213> Homo sapiens

<400> 130

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 Ser Ser Tyr
 20 25 30

Trp Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Val Trp Val
 35 40 45

Ser Arg Ile Asn Ser Asp Gly Ser Ser Thr Ser Tyr Ala Asp Ser Val
 50 55 60

Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala 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 Gly Thr Thr Gly Ala Phe Asp Ile Trp Gly Gln Gly
 100 105 110

Thr Met Val Thr Val Ser Ser
 115

<210> 131
 <211> 122
 <212> PRT
 <213> Homo sapiens

<400> 131

Glu Val His Leu Val Glu Ser Gly Gly Gly Leu Val Lys 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

Ser Met Asn Trp Phe Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

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

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

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys

Al a Arg Al a Gly Ala Val Ala Ala Leu Tyr Asn Trp Phe Asp Pro Trp
 100 105 110

Gly Gl n Gly Thr Leu Val Thr Val Ser Ser
 115 120

<210> 132
 <211> 120
 <212> PRT
 <213> Homo sapi ens

<400> 132

Gl n Ile Thr Leu Lys Gl u Ser Gly Pro Pro Leu Val Lys Pro Thr Gl n
 1 5 10 15

Thr Leu Thr Leu Thr Cys Thr Phe Ser Gly Phe Ser Leu Ser Thr Ser
 20 25 30

Gly Val Gly Val Gly Trp Ile Arg Gl n Pro Pro Gly Lys Ala Leu Gl u
 35 40 45

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

Leu Lys Ser Arg Leu Thr Ile Thr Lys Asp Thr Ser Lys Asn Gl n Val
 65 70 75 80

Val Leu Thr Met Thr Asn Met Asp Pro Val Asp Thr Ala Thr Tyr Tyr
 85 90 95

Cys Ala His Ser Pro Arg Tyr Ser Gly Tyr Phe Asp Tyr Trp Gly Gl n
 100 105 110

Gly Thr Leu Val Thr Val Ser Ser
 115 120

<210> 133
 <211> 118
 <212> PRT
 <213> Homo sapi ens

<400> 133

Gl n Val Gl n Leu Val Gl n Ser Gly Ala Gl u Val Lys Lys Pro Gly Ala
 1 5 10 15

Ser Val Lys Val Ser Cys Lys Ala Ser Gly Tyr Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gl n Ala Pro Gly Gl n Gly Leu Gl u Trp Met
 35 40 45

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Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Ala Gln Lys Phe
50 55 60

Gln Gly Lys Phe 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 Glu Ser Pro Tyr Trp Tyr Phe Asp Leu Trp Gly Arg Gly Thr
100 105 110

Leu Val Thr Val Ser Ser
115

<210> 134
<211> 116
<212> PRT
<213> Homo sapiens

<400> 134

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Lys Asn Pro Ser Leu Lys Ser
50 55 60

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

Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala Arg
85 90 95

Trp Arg Thr Phe Tyr Gly Met Asp Val Trp Gly Gln Gly Thr Thr Val
100 105 110

Thr Val Ser Ser
115

<210> 135
<211> 113
<212> PRT
<213> Homo sapiens

<400> 135

Gln Val Gln Leu Gln Glu Ser Gly Pro Gly Leu Val Lys Pro Ser Glu

1 5 10 15

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

Tyr Trp Thr Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
 35 40 45

Gly Tyr Ile Ser Tyr Ser Gly Gly Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Gl u Leu Gly Asn Tyr Trp Gly Gl n Gly Thr Leu Val Thr Val Ser
 100 105 110

Ser

<210> 136
 <211> 123
 <212> PRT
 <213> Homo sapiens

<400> 136

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gly Leu Val Lys 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

Asn Met Asn Trp Val Arg Gl n Ala Pro Gly Lys Gly Leu Gl u Trp Val
 35 40 45

Ser Ser Ile Ser Ser Ser Ser Thr Tyr Ile Tyr Tyr Ala Asp Ser Leu
 50 55 60

Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala Met Asn Ser Leu Tyr
 65 70 75 80

Leu Gl n Met Ser Ser Leu Arg Ala Gl u Asp Thr Ala Val Tyr Tyr Cys
 85 90 95

Ala Arg Asp Ser Gly Thr Tyr Pro Tyr Tyr Tyr Phe Gly Met Asp Val
 100 105 110

Trp Gly Gl n Gly Thr Thr Val Thr Val Ser Ser
 115 120

<210> 137
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 137

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

<210> 138
 <211> 120
 <212> PRT
 <213> Homo sapiens

<400> 138

Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Lys 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
 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 Ser Tyr Ile Tyr Tyr Ala Asp Ser Val
 50 55 60
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala Lys Asn Ser Leu Phe
 65 70 75 80

Leu Gl n Met Asn Ser Leu Arg Ala Gl u Asp Thr Ala Val Tyr Tyr Cys
85 90 95

Ala Arg Gly Gly Ser Phe Pro Tyr Asn Trp Phe Asp Pro Trp Gly Gl n
100 105 110

Gly Thr Leu Val Thr Val Ser Ser
115 120

<210> 139
<211> 120
<212> PRT
<213> Homo sapiens

<400> 139

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gl y Leu Val Gl n Pro Gly Arg
1 5 10 15

Ser Leu Arg Leu Ser Cys Thr Ala Ser Gly Phe Thr Phe Gly Asp Tyr
20 25 30

Ala Met Thr Trp Phe Arg Gl n Ala Pro Gly Lys Gly Leu Gl u Trp Val
35 40 45

Gly Phe Ile Arg Ser Lys Ala Tyr Gly Gly Thr Thr Gl u Tyr Ala Ala
50 55 60

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

Ala Tyr Leu Gl n Met Asn Ser Leu Lys Thr Gl u Asp Thr Ala Val Tyr
85 90 95

Tyr Cys Thr Arg Gl u Gly Ser Phe Gly Ala Leu Asp Ile Trp Gly Gl n
100 105 110

Gly Thr Met Val Thr Val Ser Ser
115 120

<210> 140
<211> 121
<212> PRT
<213> Homo sapiens

<400> 140

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gl y Leu Val Lys Pro Gly Gly
1 5 10 15

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Ala
20 25 30

Trp Met Asn Trp Val Arg Gl n Ala Pro Gly Lys Gly Leu Gl u Trp Val
35 40 45

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Gly Arg Ile Asn Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Ala Ala
50 55 60

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

Gln Gly Thr Met Val Thr Val Ser Ser
115 120

<210> 141
<211> 118
<212> PRT
<213> Homo sapiens

<400> 141

Gln Val Gln Val 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 Thr Phe Thr Gly Tyr
20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Ala Gln Arg Phe
50 55 60

Gln Gly 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 Leu Ser Pro Tyr Trp Tyr Phe Asp Leu Trp Gly Arg Gly Thr
100 105 110

Leu Val Thr Val Ser Ser
115

<210> 142
<211> 126
<212> PRT
<213> Homo sapiens

<400> 142

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 Ser Ser Tyr
 20 25 30

Ser Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

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

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

Leu Gln Met Asn Ser Leu Arg Asp Glu Asp Thr Ala Val Tyr Tyr Cys
 85 90 95

Ala Arg Glu Gly Glu Arg Phe Cys Gly Ala Asp Cys Tyr Pro His Trp
 100 105 110

Phe Asp Pro Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
 115 120 125

<210> 143
 <211> 126
 <212> PRT
 <213> Homo sapiens
 <400> 143

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Arg Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Asp Asn Trp Asn Tyr Gly Gly Pro Thr Tyr Tyr Tyr Tyr Tyr Gly
 100 105 110

Met Asp Val Trp Gly Gln Gly Thr Thr Val Thr Val Ser Ser

115

<210> 144
<211> 116
<212> PRT
<213> Homo sapiens

<400> 144

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Pro Phe Ser Val Tyr
20 25 30

Trp Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

Ala Asn Ile Lys Gln Asp Gly Ser Glu Lys Tyr Tyr Val Asp Ser Val
50 55 60

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

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Cys Cys
85 90 95

Ala Ser Gly Tyr His Leu Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 145
<211> 122
<212> PRT
<213> Homo sapiens

<400> 145

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Asn Ala
20 25 30

Trp Met Ile Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

Gly Arg Ile Lys Ser Lys Thr Asp Gly Gly Thr Thr Asp Tyr Ala Ala
50 55 60

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

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Leu Tyr Leu Gl n Met Asn Ser Leu Lys Thr Gl u Asp Thr Al a Val Tyr
85 90 95

Tyr Cys Thr Ser Thr Asp Tyr Gly Asp Tyr Asp Al a Phe Asp Ile Trp
100 105 110

Gly Gl n Gly Thr Met Val Thr Val Ser Ser
115 120

<210> 146
<211> 120
<212> PRT
<213> Homo sapi ens
<400> 146

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gly Leu Val Lys Pro Gly Gly
1 5 10 15

Ser Leu Arg Leu Ser Cys Al a Al a Ser Gly Phe Thr Phe Ser Ser Tyr
20 25 30

Ser Met Asn Trp Val Arg Gl n Al a Pro Gly Lys Gly Leu Gl u Trp Val
35 40 45

Ser Ser Ile Ser Ser Ser Ser Ser Tyr Ile Tyr Tyr Al a Asp Ser Val
50 55 60

Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Al a Lys Asn Ser Leu Tyr
65 70 75 80

Leu Gl n Met Asn Ser Leu Arg Al a Gl u Asp Thr Al a Val Tyr Tyr Cys
85 90 95

Al a Arg Gly Gly Ser Tyr Pro Tyr Asn Trp Phe Asp Pro Trp Gly Gl n
100 105 110

Gly Thr Leu Val Thr Val Ser Ser
115 120

<210> 147
<211> 118
<212> PRT
<213> Homo sapi ens
<400> 147

Gl n Val Gl n Leu Val Gl n Ser Gly Al a Gl u Val Lys Lys Pro Gly Al a
1 5 10 15

Ser Val Lys Val Ser Cys Lys Al a Ser Gly Tyr Thr Phe Thr Gly Tyr
20 25 30

Tyr Met His Trp Val Arg Gl n Al a Pro Gly Gl n Gly Leu Gl u Trp Met

35

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Ala Gln Lys Phe
50 55 60

Gln Gly 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 Asp Asn His Asp Gly Ala Phe Asp Ile Trp Gly Gln Gly Thr
100 105 110

Met Val Thr Val Ser Ser
115

<210> 148
<211> 119
<212> PRT
<213> Homo sapiens

<400> 148

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 Thr Phe Thr Gly Tyr
20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Ala Gln Lys Phe
50 55 60

Gln Gly 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 Asp Glu Asp Ser Gly Ser Tyr Phe Asp Tyr Trp Gly Gln Gly
100 105 110

Thr Leu Val Thr Val Ser Ser
115

<210> 149
<211> 121
<212> PRT
<213> Homo sapiens

<400> 149

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Gln Ile Thr Leu Lys Glu Ser Gly Pro Thr Leu Val Lys Pro Thr Gln
1 5 10 15

Thr Leu Thr Leu Thr Cys Thr Phe Ser Gly Phe Ser Leu Ser Thr Ser
20 25 30

Gly Val Gly Val Gly Trp Ile Arg Gln Pro Pro Gly Lys Ala Leu Gln
35 40 45

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

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

Val Leu Thr Met Thr Asn Met Asp Pro Val Asp Thr Ala Thr Tyr Tyr
85 90 95

Cys Ala His Gln Glu Arg Arg Ser Gly Trp Ser Phe Asp Tyr Trp Gly
100 105 110

Gln Gly Thr Leu Val Ser Val Ser Ser
115 120

<210> 150
<211> 126
<212> PRT
<213> Homo sapiens

<400> 150

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
20 25 30

Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

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

Phe Asp Pro Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
 115 120 125

<210> 151
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 151

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Thr Tyr
 20 25 30

Trp Met Thr Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

Ala Asn Ile Lys Gln Asp Gly Ser Glu Lys Tyr Tyr Val Asp Ser Val
 50 55 60

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

Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Cys Cys
 85 90 95

Ala Ser Gly Tyr His Leu Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val
 100 105 110

Thr Val Ser Ser
 115

<210> 152
 <211> 118
 <212> PRT
 <213> Homo sapiens

<400> 152

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 Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
 35 40 45

Gly Arg Ile Asn Pro Asn Ser Gly Gly Thr Tyr Tyr Ala Gln Lys Phe
 50 55 60

Gln Gly Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Thr Ala Tyr
 65 70 75 80

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Val Glu Leu Ser Arg Leu Arg Ser Asp Asp Thr Ala Val Tyr Tyr Cys
85 90 95

Ala Arg Lys Asn Pro Trp Gly His Phe Asp Tyr Trp Gly Gln Gly Thr
100 105 110

Leu Val Thr Val Ser Ser
115

<210> 153
<211> 118
<212> PRT
<213> Homo sapiens
<400> 153

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

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Thr Phe
20 25 30

Ala Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
35 40 45

Ala Val Ile Ser Tyr Asp Gly Ser Asn Lys Tyr 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 Met Tyr Tyr Cys
85 90 95

Ala Arg Asp Gly Thr Gly Asp Ala Leu Asp Ile Trp Gly Gln Gly Thr
100 105 110

Met Val Thr Val Ser Ser
115

<210> 154
<211> 117
<212> PRT
<213> Homo sapiens
<400> 154

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

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

Phe Trp Ser Trp Leu Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

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

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

Glu Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Asp Arg Thr Phe Tyr Gly Met Asp Val Trp Gly Gln Gly Thr Thr
 100 105 110

Val Thr Val Ser Ser
 115

<210> 155
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 155

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Ser Ser Gly Ser Ile Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Phe Tyr Cys Ala
 85 90 95

Arg Asp Arg Ala Phe Tyr Gly Met Asp Val Trp Gly Gln Gly Thr Thr
 100 105 110

Val Thr Val Ser Ser
 115

<210> 156
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 156

Gln Val Gln Phe Val Glu Ser Gly Gly Gly Val Val Gln Pro Gly Arg
 1 5 10 15
 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Leu Ser Ser Tyr
 20 25 30
 Gly Met His Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45
 Ala Val Ile Trp Tyr Asp Gly Ser Asn Glu 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 Asp Arg Ala Trp Gly Ser Gly Asp Trp Gly Gln Gly Thr Leu
 100 105 110
 Val Thr Val Ser Ser
 115

<210> 157
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 157

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

Val Thr Val Ser Ser
115

<210> 158
<211> 117
<212> PRT
<213> Homo sapiens

<400> 158

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

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

Tyr Trp Asn Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr His Cys Ala
85 90 95

Arg Gly Arg Leu Asn Gly Ala Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 159
<211> 117
<212> PRT
<213> Homo sapiens

<400> 159

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

Thr Leu Ser Leu Thr Cys Thr Ile Ser Ser Gly Ser Ile Ser Ser Tyr
20 25 30

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr His Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn His Phe Ser Leu

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Tyr Trp Thr Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ile Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Arg Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Gly Arg Ile Ile Gly Ala Phe Asp Ile Trp Gly Gln Gly Thr Met
 100 105 110

Val Thr Val Ser Ser
 115

<210> 162
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 162

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Thr Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Asn Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Arg Val Ile Trp Phe Asp Pro Trp Gly Gln Gly Thr Leu Val
 100 105 110

Thr Val Ser Ser
 115

<210> 163
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 163

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 Thr Phe Thr Asp Tyr
20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Ala Asn Tyr Ala Gln Lys Phe
50 55 60

Gln Gly Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Ile Ala Tyr
65 70 75 80

Met Glu Leu Ser Arg Leu Arg Ser Asp Asp Thr Ala Ile Tyr Tyr Cys
85 90 95

Ala Arg Gly Arg Leu Thr Phe Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 164
<211> 117
<212> PRT
<213> Homo sapiens

<400> 164

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

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Ile Ser Ser Phe
20 25 30

Tyr Trp Ser Trp Ile Arg Gln Ser Pro Gly Arg Gly Leu Glu Trp Ile
35 40 45

Gly Tyr Ile Phe Tyr Ser Gly Ser Thr Lys Tyr Asn Pro Ser Leu Met
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gly Arg Leu Gly Asn Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 165
<211> 117
<212> PRT
<213> Homo sapiens

<400> 165

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Ile Ser Gly Thr Thr Asn Tyr Asn Pro Ser Leu Met
50 55 60

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

Lys Leu Thr Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Leu Val Arg Ala Phe Asp Ile Trp Gly Gln Gly Thr Met
100 105 110

Val Thr Val Ser Ser
115

<210> 166
<211> 117
<212> PRT
<213> Homo sapiens

<400> 166

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Lys Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Gly Val Thr Gly Gly Phe Asp Pro Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 167
<211> 116
<212> PRT
<213> Homo sapiens

<400> 167

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Lys Asn Pro Ser Leu Lys Ser
50 55 60

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

Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala Arg
85 90 95

Leu Thr Val Val Gly Ala Leu Asp Tyr Trp Gly Gln Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 168
<211> 118
<212> PRT
<213> Homo sapiens

<400> 168

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 Ser Ser Tyr
20 25 30

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Ser Met Asn Trp Val Arg Gl n Al a Pro Gly Lys Gly Leu Gl u Trp Val
35 40 45

Ser Tyr Ile Ser Ser Ser Ser Thr Ile Tyr Tyr Al a Asp Ser Val
50 55 60

Lys Gly Arg Phe Thr Ile Ser Ser Asp Asn Al a Lys Asn Ser Leu Tyr
65 70 75 80

Leu Gl n Met Asp Ser Leu Arg Asp Gl u Asp Thr Al a Val Tyr Tyr Cys
85 90 95

Al a Arg Asp Arg Leu Gly Ile Pro Phe Asp Tyr Trp Gly Gl n Gly Thr
100 105 110

Leu Val Thr Val Ser Ser
115

<210> 169
<211> 116
<212> PRT
<213> Homo sapi ens

<400> 169

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Asp Ser Ile Ser Asn Tyr
20 25 30

Tyr Trp Ser Trp Ile Arg Gl n Pro Al a Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Asn Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Tyr Tyr Cys Al a
85 90 95

Arg Gl u Arg Ile Asn Trp Phe Asp Pro Trp Gly Gl n Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 170
<211> 117
<212> PRT

<213> Homo sapiens

<400> 170

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Ser Ser Gly Thr Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Asn Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Ile Thr Gly His Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 171

<211> 117

<212> PRT

<213> Homo sapiens

<400> 171

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Val Leu Gly Tyr Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu

100

105

110

Val Thr Val Ser Ser
115

<210> 172
<211> 116
<212> PRT
<213> Homo sapiens

<400> 172

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Gly Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Leu Tyr Gly Met Asp Val Trp Gly Gln Gly Thr Thr Val
100 105 110

Thr Val Ser Ser
115

<210> 173
<211> 119
<212> PRT
<213> Homo sapiens

<400> 173

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Asn Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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Ser Arg Val Thr Met Ser Leu Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Val Gly Ala Ile Gly Val Asp Ala Phe Asp Ile Trp Gly Gl n Gly
100 105 110

Thr Met Val Thr Val Ser Ser
115

<210> 174
<211> 116
<212> PRT
<213> Homo sapi ens

<400> 174

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Ala Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Ser Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gl u Arg Ile Ile Trp Phe Asp Pro Trp Gly Gl n Gly Thr Leu Val
100 105 110

Thr Val Ser Ser
115

<210> 175
<211> 118
<212> PRT
<213> Homo sapi ens

<400> 175

Gl u Val Gl n Leu Val Gl u Ser Gly Gly Gly Leu Val Lys Pro Gly Gly
1 5 10 15

Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr

Asn Met Asn Trp Val Arg Gl n Al a Pro Gl y Lys Gl y Leu Gl u Trp Val
 35 40 45

Ser Ser Ile Ser Ser Ser Ser Asn Tyr Ile Tyr Tyr Al a Asp Ser Val
 50 55 60

Lys Gl y Arg Phe Thr Ile Ser Gl y Asp Thr Al a Lys Asn Ser Leu Phe
 65 70 75 80

Leu Gl n Met Ile Ser Leu Arg Val Gl u Asp Thr Al a Val Tyr Tyr Cys
 85 90 95

Al a Arg Asp Arg Leu Gl y Ile Pro Phe Asp Tyr Trp Gl y Gl n Gl y Ser
 100 105 110

Leu Val Thr Val Ser Ser
 115

<210> 176
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 176

Gl n Val Gl n Leu Gl n Gl u Ser Gl y Pro Gl y Leu Val Lys Pro Ser Gl u
 1 5 10 15

Thr Leu Ser Leu Thr Cys Thr Val Ser Gl y Al a Ser Ile Ser Ser Tyr
 20 25 30

Tyr Trp Ser Trp Ile Arg Gl n Pro Pro Gl y Lys Gl y Leu Gl u Trp Ile
 35 40 45

Gl y Tyr Phe Tyr Tyr Ser Gl y Ser Thr Asn Tyr Ser Pro Ser Leu Lys
 50 55 60

Ser Arg Val Thr Phe Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Tyr Phe Cys Al a
 85 90 95

Arg Gl y Lys Val Gl y Val Pro Phe Asp Tyr Trp Gl y Gl n Gl y Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 177
 <211> 117

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<212> PRT
 <213> Homo sapiens

<400> 177

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

<210> 178
 <211> 119
 <212> PRT
 <213> Homo sapiens

<400> 178

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

Arg Val Gly Ala Ile Gly Val Asp Ala Phe Asp Ile Trp Gly Gln Gly
 100 105 110

Thr Met Val Thr Val Ser Ser
 115

<210> 179
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 179

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

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

Tyr Trp Ser Trp Ile Trp Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Pro Lys Asn Gln Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Arg Leu Thr Gly Ala Phe Asp Ile Trp Gly Gln Gly Thr Met
 100 105 110

Val Thr Val Ser Ser
 115

<210> 180
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 180

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 Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
 35 40 45

Val Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Ala Gln Lys Phe
 50 55 60

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Gln Gly 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 Gly Arg Leu Thr Phe Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 181
<211> 117
<212> PRT
<213> Homo sapiens

<400> 181

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 Thr Phe Thr Gly Tyr
20 25 30

Tyr Met His Trp Val Arg Gln Ala Pro Gly Gln Gly Leu Glu Trp Met
35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Tyr Cys Ala Gln Lys Phe
50 55 60

Gln Gly 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 Gly Arg Leu Thr Phe Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 182
<211> 117
<212> PRT
<213> Homo sapiens

<400> 182

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

Thr Leu Ser Leu Thr Cys Thr Val Ser Asp Gly Ser Ile Ser Ser Tyr
 20 25 30

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Thr Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

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

Val Thr Val Ser Ser
 115

<210> 183
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 183

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Val
 35 40 45

Gly Arg Leu Phe Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Ile Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Arg Leu Thr Gly Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 184

<211> 117
 <212> PRT
 <213> Homo sapiens

<400> 184

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

<210> 185
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 185

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

Arg Asp Arg Leu Gly Arg Ala Phe Asp Ile Trp Gly Gln Gly Thr Met
 100 105 110

Val Thr Val Ser Ser
 115

<210> 186
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 186

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Gly Arg Leu Thr Gly His Phe Asp Tyr Trp Gly Gln Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 187
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 187

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Tyr Thr Asn Tyr Asn Pro Ser Leu Lys

50

55

60

Ser Arg Val Ser Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Tyr Tyr Cys Al a
85 90 95

Arg Gl u Arg Leu Gly Al a Phe Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 188
<211> 117
<212> PRT
<213> Homo sapi ens

<400> 188

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Phe Tyr Cys Al a
85 90 95

Arg Ile Thr Val Thr Ser Al a Phe Asp Ile Trp Gly Gl n Gly Thr Met
100 105 110

Val Thr Val Ser Ser
115

<210> 189
<211> 116
<212> PRT
<213> Homo sapi ens

<400> 189

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Lys Lys Pro Ser Leu Lys Ser
 50 55 60

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

Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala Arg
 85 90 95

Ser Thr Val Val Asn Ala Phe Asp Ile Trp Gly Gln Gly Thr Met Val
 100 105 110

Thr Val Ser Ser
 115

<210> 190
 <211> 126
 <212> PRT
 <213> Homo sapiens

<400> 190

Gl u Val Gln Leu Val Gl u 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

Ser Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

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

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

Leu Gln Met Asn Ser Leu Arg Asp Glu Asp Thr Ala Val Tyr Tyr Cys
 85 90 95

Ala Arg Asp Leu Pro Ile Thr Met Ile Val Val Val Met Pro Asp Ala
 100 105 110

Phe Asp Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120 125

<210> 191
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 191

Glu Val Gln Leu Val₅ Glu Ser Gly Gly₁₀ Gly Leu Val Gln Pro Gly₁₅ Gly

Ser Leu Arg Leu₂₀ Ser Cys Ala Ala Ser₂₅ Gly Phe Thr Phe Ser₃₀ Asn Tyr

Trp Met Arg₃₅ Trp Val Arg Gln Ala₄₀ Pro Gly Lys Gly Leu₄₅ Glu Trp Val

Ala Asn Ile Lys Gln Asp Gly₅₅ Ser Glu Lys Tyr Tyr₆₀ Val Asp Ser Val

Lys Gly Arg Phe Thr Ile₇₀ Ser Arg Asp Asn Ala₇₅ Lys Tyr Ser Leu Tyr₈₀

Leu Gln Met Asn₈₅ Ser Leu Arg Ala Glu Asp₉₀ Thr Ala Val Tyr Tyr₉₅ Cys

Ala Arg Asp Arg₁₀₀ Leu Gly Ile Phe Asp₁₀₅ Tyr Trp Gly Gln Gly₁₁₀ Thr Leu

Val Thr Val₁₁₅ Ser Ser

<210> 192
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 192

Glu Val Gln Leu Val₅ Gln Ser Gly Ala Glu₁₀ Val Lys Lys Pro Gly₁₅ Glu

Ser Leu Lys Ile₂₀ Ser Cys Lys Gly Ser₂₅ Gly Tyr Ser Phe Thr Asn Tyr

Trp Ile Gly₃₅ Trp Val Arg Gln Met₄₀ Pro Gly Lys Gly Leu₄₅ Glu Trp Met

Gly Ile Ile Tyr Pro Gly Asp₅₅ Ser Asp Thr Arg Tyr₆₀ Ser Pro Ser Phe

Gln Gly Gln Val Thr Ile₇₀ Ser Ala Asp Ile Ser₇₅ Ile Thr Thr Ala Tyr₈₀

Leu Gln Trp Ser Ser₈₅ Leu Lys Ala Ser Asp₉₀ Thr Ala Met Tyr Tyr₉₅ Cys

Al a Arg His Arg Leu Gly Phe Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 193
 <211> 118
 <212> PRT
 <213> Homo sapiens
 <400> 193

Gl n Val Gl n Leu Val Gl n Ser Gly Al a Gl u Val Lys Lys Pro Gly Al a
 1 5 10 15

Ser Val Lys Val Ser Cys Lys Al a Ser Gly Tyr Thr Phe Thr Gly Tyr
 20 25 30

Tyr Met His Trp Val Arg Gl n Al a Pro Gly Gl n Gly Leu Gl u Trp Met
 35 40 45

Gly Trp Ile Asn Pro Asn Ser Gly Gly Thr Asn Tyr Al a Gl n Lys Phe
 50 55 60

Gl n Gly Arg Val Thr Met Thr Arg Asp Thr Ser Ile Ser Thr Al a Tyr
 65 70 75 80

Met Gl u Leu Ser Arg Leu Arg Ser Asp Asp Thr Al a Val Tyr Tyr Cys
 85 90 95

Al a Arg Leu Tyr Tyr Tyr Tyr Asn Met Asp Val Trp Gly Gl n Gly Thr
 100 105 110

Thr Val Thr Val Ser Ser
 115

<210> 194
 <211> 118
 <212> PRT
 <213> Homo sapiens
 <400> 194

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
 1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Thr Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Gly Trp Trp Glu Leu Thr Phe Phe Asp Tyr Trp Gly Gln Gly Thr
 100 105 110

Leu Val Thr Val Ser Ser
 115

<210> 195
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 195

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Ala Ser Gly Gly Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Val Ser Ser Val Ile Ala Ala Asp Thr Ala Ile Tyr Tyr Cys Ala
 85 90 95

Arg Asn Arg Leu Gly Ile Tyr Asp Tyr Trp Gly Gln Gly Ser Leu Val
 100 105 110

Thr Val Ser Ser
 115

<210> 196
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 196

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

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

Tyr Trp Ser Trp Ile Arg Gln Ser Pro Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Phe Cys Ala
85 90 95

Arg Ala Leu Leu Thr Gly Gly Phe Ala Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 197
<211> 117
<212> PRT
<213> Homo sapiens

<400> 197

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
35 40 45

Gly Arg Ile Tyr Ala Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Thr Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Val Gly Ile Tyr Phe Asp Tyr Trp Gly Gln Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

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<210> 198
 <211> 125
 <212> PRT
 <213> Homo sapiens

<400> 198

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 Ser Ser Tyr
 20 25 30

Ser Met Asn Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
 35 40 45

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

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

Leu Gln Met Asp Ser Leu Arg Asp Glu Asp Thr Ala Val Tyr Tyr Cys
 85 90 95

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

Asp Ile Trp Gly Gln Gly Thr Met Val Thr Val Ser Ser
 115 120 125

<210> 199
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 199

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Ile Lys Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Val Ala Met Ser Val Asp Thr Ser Lys Asn Gln Phe Ser Leu
 65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala

Arg Gl u Gly Ile Leu Gly Tyr Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 200
 <211> 117
 <212> PRT
 <213> Homo sapi ens
 <400> 200

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
 1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
 35 40 45

Gly Tyr Val Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

Ser Arg Val Thr Thr Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
 65 70 75 80

Lys Leu Arg Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Asp Arg Leu Gly Ile Ala Phe Asp Ile Trp Gly Gl n Gly Thr Met
 100 105 110

Val Thr Val Ser Ser
 115

<210> 201
 <211> 118
 <212> PRT
 <213> Homo sapi ens
 <400> 201

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
 1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Ser Ala Gly Lys Gly Leu Gl u Trp Ile
 35 40 45

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Gly His Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gl u Arg Thr Val Val Thr Tyr Phe Asp Tyr Trp Gly Gl n Gly Thr
100 105 110

Leu Val Thr Val Ser Ser
115

<210> 202
<211> 117
<212> PRT
<213> Homo sapiens

<400> 202

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

His Trp Ser Trp Ile Arg Gl n Pro Ala Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Gly Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gl u Arg Leu Thr Gly Tyr Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 203
<211> 116
<212> PRT
<213> Homo sapiens

<400> 203

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u

1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Ser Arg Leu Gly Ile Phe Asp Tyr Trp Gly Gln Gly Thr Leu Val
 100 105 110

Thr Val Ser Ser
 115

<210> 204
 <211> 117
 <212> PRT
 <213> Homo sapiens
 <400> 204

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Thr Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Ser Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Arg Met Ala Thr Ile Phe Asp Tyr Trp Gly Gln Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 205
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 205

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

<210> 206
 <211> 125
 <212> PRT
 <213> Homo sapiens

<400> 206

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

Leu Gl n Met Asn Ser 85 Leu Arg Asp Gl u Asp Thr Ala Val Tyr Tyr Cys 95

Ala Arg Asp Tyr 100 Tyr Gly Ser Ser Gly 105 Tyr Tyr Tyr Pro His Ala Phe 110

Asp Ile Trp 115 Gly Gl n Gly Thr Met 120 Val Thr Val Ser Ser 125

<210> 207
 <211> 116
 <212> PRT
 <213> Homo sapiens

<400> 207

Gl u Val Gl n Leu Val 5 Gl u Ser Gly Gly 10 Gl y Leu Val Lys Pro Gly Gly 15

Ser Leu Arg Leu 20 Ser Cys Ala Ala Ser 25 Gl y Phe Thr Phe Ser 30 Tyr Tyr

Ser Met Asn 35 Trp Val Arg Gl n Ala Pro Gly Lys Gly Leu Gl u Trp Val 45

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

Lys Gly Arg Phe Thr 70 Ile Ser Arg Asp Asn Ala 75 Lys Asn Ser Leu Tyr 80

Leu Gl n Met Asn Ser 85 Leu Arg Ala Gl u Asp Thr Ala Val Tyr Tyr Cys 95

Ala Arg Gly Gly 100 Leu Arg Phe Asp Tyr 105 Trp Gly Gl n Gly Thr Leu Val 110

Thr Val Ser Ser 115

<210> 208
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 208

Gl n Val Gl n Leu Gl n 5 Gl u Ser Gly Pro Gly 10 Leu Val Lys Pro Ser Gl u 15

Thr Leu Ser Leu 20 Thr Cys Thr Val Ser 25 Gl y Gly Ser Ile Ser Ser Tyr 30

Tyr Trp Ser 35 Trp Ile Arg Gl n Ser 40 Pro Gly Lys Gly Leu Gl u Trp Ile 45

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Gly Tyr Ile Tyr Tyr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Ala Leu Leu Thr Gly Gly Phe Ala Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 209
<211> 117
<212> PRT
<213> Homo sapiens

<400> 209

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Ile Asn Ser Tyr
20 25 30

Tyr Trp Ser Trp Ile Arg Gl n Pro Ala Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Asn Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Gl u Arg Leu Thr Gly Tyr Phe Asp Tyr Trp Ala Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 210
<211> 117
<212> PRT
<213> Homo sapiens

<400> 210

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

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

Tyr Trp Ser Trp Ile Arg Gln Pro Pro Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Tyr Ile Tyr Tyr Ser Gly Thr Thr Asn Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Ala Leu Ile Val Gly Ala Phe Ala Tyr Trp Gly Gln Gly Thr Leu
 100 105 110

Val Thr Val Ser Ser
 115

<210> 211
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 211

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

Thr Leu Ser Leu Thr Cys Thr Val Ser Gly Gly Ser Ile Ser Ser Asn
 20 25 30

Tyr Trp Ser Trp Ile Arg Gln Pro Ala Gly Lys Gly Leu Glu Trp Ile
 35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Lys Tyr Asn Pro Ser Leu Lys
 50 55 60

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

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
 85 90 95

Arg Glu Gly Ile Val Gly Ala Phe His Tyr Trp Gly Gln Gly Ala Leu
 100 105 110

Val Thr Val Ser Ser

115

<210> 212
 <211> 118
 <212> PRT
 <213> Homo sapiens

<400> 212

Glu Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Lys Pro Gly Gly
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 Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Ser Ser Tyr
 20 25 30
 Ser Met Lys Trp Val Arg Gln Ala Pro Val Lys Gly Leu Glu Trp Val
 35 40 45
 Ser Ser Ile Ser Ser Ser Ser Tyr Ile Tyr Tyr Gly Asp Ser Val
 50 55 60
 Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ala Lys Asn Ser 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 Leu Gly Arg Ala Phe Asp Ile Trp Gly Gln Gly Thr
 100 105 110
 Met Val Thr Val Ser Ser
 115

<210> 213
 <211> 117
 <212> PRT
 <213> Homo sapiens

<400> 213

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

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Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Leu Gly Met Phe Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 214
<211> 117
<212> PRT
<213> Homo sapiens

<400> 214

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Ala Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Arg Ile Tyr Thr Ser Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Ala Ala Asp Thr Ala Val Tyr Tyr Cys Ala
85 90 95

Arg Glu Arg Leu Gly Met Phe Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 215
<211> 117
<212> PRT
<213> Homo sapiens

<400> 215

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Ser Trp Ile Arg Gl n Pro Ala Gly Lys Gly Leu Gl u Trp Ile

35

Gly Arg Phe Tyr Thr Ser Gly Ser Thr Ser Cys Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Met Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Tyr Tyr Cys Al a
85 90 95

Arg Gl u Gly Ile Leu Gly Tyr Phe Asp Tyr Trp Gly Gl n Gly Thr Leu
100 105 110

Val Thr Val Ser Ser
115

<210> 216
<211> 117
<212> PRT
<213> Homo sapi ens

<400> 216

Gl n Val Gl n Leu Gl n Gl u Ser Gly Pro Gly Leu Val Lys Pro Ser Gl u
1 5 10 15

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

Tyr Trp Asn Trp Ile Arg Gl n Pro Pro Gly Lys Gly Leu Gl u Trp Ile
35 40 45

Gly Tyr Ile Tyr Tyr Ile Gly Ser Thr Asn Tyr Asn Pro Ser Leu Lys
50 55 60

Ser Arg Val Thr Ile Ser Val Asp Thr Ser Lys Asn Gl n Phe Ser Leu
65 70 75 80

Lys Leu Ser Ser Val Thr Al a Al a Asp Thr Al a Val Tyr Tyr Cys Al a
85 90 95

Arg Gly Arg Leu Thr Tyr Al a Phe Asp Ile Trp Gly Gl n Gly Thr Met
100 105 110

Val Thr Val Ser Ser
115

<210> 217
<211> 319
<212> DNA
<213> Homo sapi ens

<400> 217

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 gggaaagccc ctaagctcct gatctacgat gaatccaatt tggaaacagg ggtcccatca 180
 aggttcagtg gaagtggatc tgggacagat tttactttca ccatcagcag cctgcagcct 240
 gaagatattg gaacatatta ctgtcaacag tatgataatc ttctcacttt cggccctggg 300
 accaaagtgg atatcaaac 319

<210> 218
 <211> 340
 <212> DNA
 <213> Homo sapi ens

<400> 218
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 tggttcctgc agaagccagg gcagtctcca cagctcctga tctatacgct ttcctatcgg 180
 gcctctggag tcccagacag gttcagtggc agtgggtcag gcactgattt cacttgaaa 240
 atcagcaggg tggaggctga cgatgttggg gtttattact gcatgcaacg tatagagttt 300
 ccgttcactt ttggccaggg gaccaagctg gagatcaaac 340

<210> 219
 <211> 340
 <212> DNA
 <213> Homo sapi ens

<400> 219
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 tggtagcagc agaaagtagg acagcctccg aagctgctca tttactgggc atctacccgg 180
 gaatccgggg tccctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
 atcagcagcc tgcaggctga agatgtggca gtttattact gtcagcaata ttatagtact 300
 ccgttcactt tcggcggagg gaccaagtg gagatcaaac 340

<210> 220
 <211> 322
 <212> DNA
 <213> Homo sapi ens

<400> 220
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 cctggccagg ctcccaggct cctcatctat ggtgcatcca gcagggccac tggcatccca 180
 gacaggttca gtggcagtgg gtctgggaca gacttcactc tcaccatcag cagactggag 240
 cctgaagatt ttgcagtgta ttactgtcag cagtatggta gctcgtcac tttcggcgga 300
 gggaccaagg tggagatcaa ac 322

<210> 221
 <211> 322
 <212> DNA
 <213> Homo sapiens

<400> 221
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 cctggccagg ctcccaggct cctcatctat ggtgcatcca gcagggccac tggcatccca 180
 gacaggttca gtggcagtgg gtctgggaca gacttcactc tcaccatcag cagactggag 240
 cctgaagatt ttgcaagtga ttactgtcag cagtatggta gctcgctcac tttcggcgga 300
 gggaccaagg tggagatcaa ac 322

<210> 222
 <211> 337
 <212> DNA
 <213> Homo sapiens

<400> 222
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 atctcctgca ggtctagtc gagcctcctg catagtaatg gatacaacta tttggaatgg 120
 tactgcaga agccaggga gtctccacag ctctgatct atttgggttc taatcgggcc 180
 tccgggtcc ctgacaggtt cagtggcagt ggatcaggca cagattttac actgaaaatc 240
 agcagagtgg aggctgagga tgttgggggt tattactgca tgcaagctct acaaactccg 300
 ctactttcg gcggaggga caaggtggag atcaaac 337

<210> 223
 <211> 322
 <212> DNA
 <213> Homo sapiens

<400> 223
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 gagaaagccc ctaagctct gatctatgct gcatccagtt tgcaaagtgg ggtccatca 180
 aggttcagcg gcagtggatc tgggacagat ttactctca ccatcagcag cctgcagcct 240
 gaagatttg caactacta ttgtcaacag gcttacagtt tcccgtacac ttttggccag 300
 gggaccaagc tggagatcaa ac 322

<210> 224
 <211> 340
 <212> DNA
 <213> Homo sapiens

<400> 224
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 atcaactgca agtccagcca gagtgttta tacagctcca acaataagaa ctacttagct 120

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tgttaccagc agaaaccagg acagcctcct aagctgctca ttttctgggc atctaccgg 180
gattccgggg tccctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
atcagcagcc tgcaggctga agatgtggca gtttattact gtcagcaata ttatagtact 300
ccgctcactt tcggcggagg gaccaaggtg gagatcaaac 340

<210> 225
<211> 319
<212> DNA
<213> Homo sapi ens

<400> 225
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gggaaagccc ctaagctcct gatctatatg gcgtctagtt tagaaagtgg ggtcccatca 180
aggttcagcg gcagtggatc tgggacagaa ttcactctca ccatcagcag cctgcagcct 240
gatgattttg caacttatta ctgccaacag tataatagtt attggacgtt cggccaaggg 300
accaaggtgg aatcaaac 319

<210> 226
<211> 322
<212> DNA
<213> Homo sapi ens

<400> 226
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atcacttgcc aggcgagtca ggacattagc gactatntaa attggtatca gcagaaacca 120
gggaaagccc ctaagctcct gatctacgat gcatccaatt tggaaacagg ggtcccatca 180
aggttcagtg gaagtggatc tgggacagat tttactttca ccatcagtag cctgcagcct 240
gaagatattg caacatatta ctgtcaacag tatgataatc tccctccac tttcggcgga 300
gggaccaagg tggatatac ac 322

<210> 227
<211> 322
<212> DNA
<213> Homo sapi ens

<400> 227
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atcacttgct gggcgagtca gggatttagc agttatttag cctggtatca gcaaaaacca 120
gggaaagccc ctaagctcct gatctatgct gcatccactt tgcaaagtgg ggtcccatca 180
aggttcagcg gcagtggatc tgggacagat ttcactctca ccatcagctg cctgcagtct 240
gaagattttg caacttatta ctgtcaacag tattatagtt acccattcac tttcggccct 300
gggaccaag tggatatac ac 322

<210> 228

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<211> 337
 <212> DNA
 <213> Homo sapiens

<400> 228
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 tacctgcaga agccagggca gtctccacag ctctgatct atttgggttc taatcgggcc 180
 tccgggggtcc ctgacaggtt cagtggcagt ggatcaggca cagattttac actgaaaatc 240
 agcagagtgg aggctgagga tgttgggggtt tattactgca tgcaagctct acaaactccg 300
 ctcactttcg gcggagggac caaggtggag atcaaac 337

<210> 229
 <211> 319
 <212> DNA
 <213> Homo sapiens

<400> 229
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 atcacttgcc gggccagtca gagtattagt agctggttgg cctggtatca gcggaaacca 120
 gggaaagccc ctaagctcct gatctataag gcgtctactt tagaaagtgg ggtcccatca 180
 aggttcagcg gcagtggatc tgggacagaa ttcactctca ccatcagcag cctgcagcct 240
 gatgattttg caacttatta ctgccaacag tttaatagtt attggacgtt cggccaaggg 300
 accaaggtgg aatcaaac 319

<210> 230
 <211> 319
 <212> DNA
 <213> Homo sapiens

<400> 230
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 atcacttgcc gggccagtca gagtattagt agctggttgg cctggtatca gcagaaacca 120
 gggaaagccc ctaagctcct gatctataag gcgtctagtt tagaaagtgg ggtcccatca 180
 aggttcagcg gcagtggatc tgggacagaa ttcactctca ccatcagcag cctgcagcct 240
 gatgattttg caacttatta ctgccaacag tataatagtt attggacgtt cggccaaggg 300
 accaaggtgg aatcaaac 319

<210> 231
 <211> 340
 <212> DNA
 <213> Homo sapiens

<400> 231
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 atctcctgca ggtctagtca ggcctcttg gatagtgatg atggaaacac ctatttggac 120
 tggtagctgc agaagccagg gcagtctcca cagctcctga tctatacgct ttcctatcgg 180

gcctctggag tcccagacag gttcagtggc agtgggtcag gcactgattt cacactgaaa 240
 atcagcaggg tggaggctga ggatgttga gtttattact gcatgcaacg tatagagttt 300
 ccttcgacct tcggccaagg gacacgactg gagattaaac 340

<210> 232
 <211> 319
 <212> DNA
 <213> Homo sapiens

<400> 232
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 gggaaagccc ctaagctcct gatctataag gcgtctagtt tagaaagtgg ggtcccatca 180
 aggttcagcg gcagtggatc tgggacagca ttactctca ccatcagcag cctgcagcct 240
 gatgattttg caacttatta ctgccaacag tataatagtc attggacggt cggccaaggg 300
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 <211> 325
 <212> DNA
 <213> Homo sapiens

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 cctggccagg ctccaggct cctcatctat ggtgcatcca gcagggccac tggcatcca 180
 gacaggttca gtggcagtg gtctgggaca gacttcactc tcaccatcag cagactggag 240
 cctgaagatt ttgagtgta ttactgtcag cagtatggta gctcaccgta cacttttggc 300
 caggggacca agctggagat caaac 325

<210> 234
 <211> 322
 <212> DNA
 <213> Homo sapiens

<400> 234
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 gggaaagccc ctaagctcct gatctacgat gcatccaatt tggaaacagg ggtcccatca 180
 aggttcagtg gaagtggatc tgggacagat ttactttca ccatcagcag cctgcagcct 240
 gaagatattg caacatatta ctgtcaacag tatgataatc tccctctcac tttcggcgga 300
 gggaccaagg tggagatcaa ac 322

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 cctggccagg ctcccaggct cctcatctat ggtgcatcca gcagggccac tggcatccca 180
 gacaggttca gtggcagtgg gtctgggaca gacttcactc tcaccatcag cagactggag 240
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 caggggacca agctggagat caaac 325

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 <211> 340
 <212> DNA
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 tggatcagc agaaccagg acagcctcct aaactgctca ttactgggc atctaccgg 180
 gaatccgggg tccctgaccg attcagcggc agcgggtctg ggacagattt cactctcacc 240
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<210> 237
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 <212> DNA
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<400> 237
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 cctggccagg ctcccaggct cctcatctat ggtgcatcca gcagggccac tggcatccca 180
 gacaggttca gtggcagtgg gtctgggaca gacttcactc tcaccatcag cagactggag 240
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 caggggacca agctggagat caaac 325

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 <211> 340
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 tggtagcagc agaaccagg acagcctcct aagctgctca ttactgggc atctaccgg 180
 gaatccgggg tccctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
 atcagcagcc tgcaggctga agatgtggca gtttattact gtcagcaata ttatagtact 300

ccgctcactt tcggcggagg gaccaaggtg gagatcaaac 340

<210> 239
 <211> 325
 <212> DNA
 <213> Homo sapiens

<400> 239
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 cctggccagg ctcccaggct cctcatctat ggtgcatcca gcagggccac tggcatccca 180
 gacaggttca gtggcagtggt gtctgggaca gacttcactc tcaccatcag cagactggag 240
 cctgaagatt ttgcagtgta ttactgtcag cagtatggta gctcactatt cactttcggc 300
 cctgggacca aagtggatat caaac 325

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 <211> 340
 <212> DNA
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 tggtagcaac agaaaccagg acagcctcct aagggtctca ttactgggc atctatccgg 180
 gaatccgggg tctctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
 atcagcagcc tgcaggctga agatgtggca gtttattact gtcagcaata ttatagtact 300
 ccgtacactt ttggccaggg gaccaagctg gagatcaaac 340

<210> 241
 <211> 340
 <212> DNA
 <213> Homo sapiens

<400> 241
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 tggtagcagc agaaaccagg acagcctcct aagctgctca ttactgggc atctacccgg 180
 gaatccgggg tccctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
 atcagcagcc tgcaggctga agatgtggca gtttattact gtcagcaata ttatagtact 300
 cctcggacgt tcggccaagg gaccaaggtg gaaatcaaac 340

<210> 242
 <211> 325
 <212> DNA
 <213> Homo sapiens

<400> 242
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 aggttcagcg gcagtggatc tgggacagaa ttactctca caatcagcag cctgcagcct 240
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 tccgggtcc ctgacaggtt cagtggcagt ggatcaggca cagatnttac actgaaaatc 240
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 tccgggtcc ctgacaggtt cagtggcagt ggatcaggca cagatnttac actgaagatc 240
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 gaatccgggg tccctgaccg attcagtggc agcgggtctg ggacagattt cactctcacc 240
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 tatacttttg gccaggggac caagctggag atcaaac 337

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 aggttcagcg gcagtggatc tgggacagat ttactctca ccatcagcag cctgcagcct 240
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ctcattctcg gcggagggac caaggtggag atcaaac 337

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 tggacgttcg gcccaaggac caaggtgga atcaaac 337

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 ttcggcggag ggaccaagct gaccgtccta g 331

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 <211> 331
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 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta tttctgtgca gcatgggatg acagcctgaa tgggtccggtg 300
 ttcggcggag ggaccagggt gaccgtccta g 331

<210> 312
 <211> 331
 <212> DNA
 <213> Homo sapiens

<400> 312
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 tcttgttctg gaagcagctc caacatcgga agtaatactg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcaggctccc tggccatcag tgggctcaag 240
 tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tgggtgtggtg 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 313
 <211> 331
 <212> DNA
 <213> Homo sapiens

<400> 313
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 tcttgttctg gaagcagctc caacatcgga agtaattatg tatactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccgg 240
 tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgag tgggtgtggtg 300
 ttcggcggag ggaccaagtt gaccgtccta g 331

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<210> 314
 <211> 331
 <212> DNA
 <213> Homo sapi ens

<400> 314
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 tcttgttctg gaagcagctc caacatcgga agtaactctg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
 gaccgattct ctggctcaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta tgactgtgca gcatgggatg acagcctgaa tgggtccggta 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 315
 <211> 331
 <212> DNA
 <213> Homo sapi ens

<400> 315
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 tcttgttctg gaagcagctc caacatcgga agtaactctg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
 gaccgattct ctggctcaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tggttgggtg 300
 ttcggcggag ggaccaagct gaccgtcctg g 331

<210> 316
 <211> 331
 <212> DNA
 <213> Homo sapi ens

<400> 316
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 tcttgttctg gaagcagctc caacatcgga agtaactctg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
 gaccgattct ctggctcaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
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 ttcggcggag ggaccaaact gaccgtccta g 331

<210> 317
 <211> 331
 <212> DNA
 <213> Homo sapi ens

<400> 317
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 tcttgttctg gaagcagctc caacatcgga agtaactctg taaactggta ccagcagctc 120

ABLX_007_02W0_ST25. txt

ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tgggtccgga 300
ttcggcggag ggaccaagct gaccgtccta g 331

<210> 318
<211> 325
<212> DNA
<213> Homo sapiens

<400> 318
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acctgtgggg gaaacaacat tggaagtaa gctgtgact ggtaccagca aaagccaggc 120
caggaccctg tgctggatc ctatagcag agcaaccggc cctcaggat ccctgagcga 180
ttctctggct ccaaccagg gaacaccgcc accctaacca tcagcaggat cgaggctggg 240
gatgaggctg actattactg tcaggtgtgg gacagtagta gtgatcatcg ggtgttcggc 300
ggagggacca agctgaccgt cctag 325

<210> 319
<211> 331
<212> DNA
<213> Homo sapiens

<400> 319
cagtctgtgc tgactcagcc accctcagcg tctgggaccc ccgggcagag ggtcaccatc 60
tcttgttctg gaagcagctc caacatcgga agtaactg taaactggta ccagcagctc 120
ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
tctgaggatg aggctgatta ttactgtgca ccatgggatg acagcctgaa tgggtgtgga 300
ttcggcggag ggaccaagct gaccgtccta g 331

<210> 320
<211> 331
<212> DNA
<213> Homo sapiens

<400> 320
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tcttgttctg gaagcagctc caacatcgga agtaactg taaactggta ccagcagctc 120
ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtccct 180
gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tagttgggtg 300
ttcggcggag ggaccatgct gaccgtccta g 331

<210> 321
<211> 331

ABLX_007_02WO_ST25. txt

<212> DNA
 <213> Homo sapiens

<400> 321
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 tcttgttctg gaagcagctc caacatcgga agtaatactg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta ttactgtgca gcacgggatg acagcctgaa tgggtgtggta 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 322
 <211> 331
 <212> DNA
 <213> Homo sapiens

<400> 322
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 tcttgttctg gaagcagctc caacatcgga agtaatactg taaactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta ttactgtgca gcacgggatg acagcctgaa tgggtgtggta 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 323
 <211> 331
 <212> DNA
 <213> Homo sapiens

<400> 323
 cagtctgtgc tgactcagcc accctcagcg tctgggaccc ccgggcagag ggtcaccatc 60
 tcttgttctg gaaccagctc caacatcgga agtaatactg tacactggta ccagcagctc 120
 ccaggaacgg cccccaaact cctcatctat agtaataatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240
 tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tgggtccggta 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 324
 <211> 331
 <212> DNA
 <213> Homo sapiens

<400> 324
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 tcttgttctg gaagcagctc caacatcgga agtaatactg taaactggta ccagcgggtc 120
 ccaggaacgg cccccaaact cctcatctat agtaatgatc agcggccctc aggggtcctc 180
 gaccgattct ctggctccaa gtctggcacc tcagcctccc tggccatcag tgggctccag 240

tctgaggatg aggctgatta ttactgtgca gcatgggatg acagcctgaa tgggtgtgta 300
 ttcggcggag ggaccaagct gaccgtccta g 331

<210> 325
 <211> 106
 <212> PRT
 <213> Homo sapiens

<400> 325

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

<210> 326
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 326

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

Arg Ile Glu Phe Pro Phe Thr Phe Gly Glu Gly Thr Lys Leu Glu Ile
100 105 110

Lys

<210> 327
<211> 113
<212> PRT
<213> Homo sapiens

<400> 327

Asp Ile Val Met Thr Glu Ser Pro Asp Ser Leu Ala Val Ser Leu Gly
1 5 10 15

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

Ser Asn Lys Lys Asn Tyr Leu Thr Trp Tyr Glu Glu Lys Val Gly Glu
35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
65 70 75 80

Ile Ser Ser Leu Glu Ala Glu Asp Val Ala Val Tyr Tyr Cys Glu Glu
85 90 95

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

Lys

<210> 328
<211> 107
<212> PRT
<213> Homo sapiens

<400> 328

Glu Ile Val Leu Thr Glu Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
1 5 10 15

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Glu Ser Val Ser Ser Ser
20 25 30

Tyr Leu Ala Trp Tyr Glu Glu Lys Pro Gly Glu Ala Pro Arg Leu Leu
35 40 45

I l e T y r G l y A l a S e r S e r A r g A l a T h r G l y I l e P r o A s p A r g P h e S e r
 50 55 60

G l y S e r G l y S e r G l y T h r A s p P h e T h r L e u T h r I l e S e r A r g L e u G l u
 65 70 75 80

P r o G l u A s p P h e A l a V a l T y r T y r C y s G l n G l n T y r G l y S e r S e r L e u
 85 90 95

T h r P h e G l y G l y G l y T h r L y s V a l G l u I l e L y s
 100 105

<210> 329
 <211> 107
 <212> PRT
 <213> Homo sapiens

<400> 329

G l u I l e V a l L e u T h r G l n S e r P r o G l y T h r L e u S e r L e u S e r P r o G l y
 1 5 10 15

G l u A r g A l a T h r L e u S e r C y s A r g A l a S e r G l n S e r V a l S e r S e r S e r
 20 25 30

T y r L e u A l a T r p T y r G l n G l n L y s P r o G l y G l n A l a P r o A r g L e u L e u
 35 40 45

I l e T y r G l y A l a S e r S e r A r g A l a T h r G l y I l e P r o A s p A r g P h e S e r
 50 55 60

G l y S e r G l y S e r G l y T h r A s p P h e T h r L e u T h r I l e S e r A r g L e u G l u
 65 70 75 80

P r o G l u A s p P h e A l a V a l T y r T y r C y s G l n G l n T y r G l y S e r S e r L e u
 85 90 95

T h r P h e G l y G l y G l y T h r L y s V a l G l u I l e L y s
 100 105

<210> 330
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 330

A s p I l e V a l M e t T h r G l n S e r P r o L e u S e r L e u P r o V a l T h r P r o G l y
 1 5 10 15

G l u P r o A l a S e r I l e S e r C y s A r g S e r S e r G l n S e r L e u L e u H i s S e r
 20 25 30

Asn Gly Tyr Asn Tyr Leu Glu Trp Tyr Leu Gl n Lys Pro Gly Gl n Ser
 35 40 45

Pro Gl n Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gl n Ala
 85 90 95

Leu Gl n Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
 100 105 110

<210> 331
 <211> 107
 <212> PRT
 <213> Homo sapiens

<400> 331

Asp Ile Gl n Met Thr Gl n Ser Pro Ser Ser Val Ser Ala Ser Val Gly
 1 5 10 15

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Pro Gly Ile Ser Ser Trp
 20 25 30

Leu Ala Trp Tyr Gl n Gl n Lys Pro Glu Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Ala Ala Ser Ser Leu Gl n Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Gl n Pro
 65 70 75 80

Gl u Asp Phe Ala Thr Tyr Tyr Cys Gl n Gl n Ala Tyr Ser Phe Pro Tyr
 85 90 95

Thr Phe Gly Gl n Gly Thr Lys Leu Glu Ile Lys
 100 105

<210> 332
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 332

Asp Ile Val Met Thr Gl n Ser Pro Asp Ser Leu Ala Val Ser Leu Gly
 1 5 10 15

Gl u Arg Ala Thr Ile Asn Cys Lys Ser Ser Gl n Ser Val Leu Tyr Ser
 20 25 30

Ser Asn Asn Lys Asn Tyr Leu Ala Trp Tyr Gl n Gl n Lys Pro Gly Gl n
 35 40 45

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

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 65 70 75 80

Ile Ser Ser Leu Gl n Ala Gl u Asp Val Ala Val Tyr Tyr Cys Gl n Gl n
 85 90 95

Tyr Tyr Ser Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Gl u Ile
 100 105 110

Lys

<210> 333
 <211> 106
 <212> PRT
 <213> Homo sapi ens

<400> 333

Asp Ile Gl n Met Thr Gl n Ser Pro Ser Thr Leu Ser Ala Ser Val Gly
 1 5 10 15

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gl n Ser Ile Ser Ser Trp
 20 25 30

Leu Ala Trp Tyr Gl n Gl n Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Met Ala Ser Ser Leu Gl u Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

Ser Gly Ser Gly Thr Gl u Phe Thr Leu Thr Ile Ser Ser Leu Gl n Pro
 65 70 75 80

Asp Asp Phe Ala Thr Tyr Tyr Cys Gl n Gl n Tyr Asn Ser Tyr Trp Thr
 85 90 95

Phe Gly Gl n Gly Thr Lys Val Gl u Ile Lys
 100 105

<210> 334
 <211> 107
 <212> PRT
 <213> Homo sapi ens

<400> 334

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 Gln Ala Ser Gln Asp Ile Ser Asp Tyr
 20 25 30

Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

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

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

Glu Asp Ile Ala Thr Tyr Tyr Cys Gln Gln Tyr Asp Asn Leu Pro Pro
 85 90 95

Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
 100 105

<210> 335
 <211> 107
 <212> PRT
 <213> Homo sapiens
 <400> 335

Ala Ile Arg Met Thr Gln Ser Pro Ser Ser Phe Ser Ala Ser Thr Gly
 1 5 10 15

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser Tyr
 20 25 30

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Ala Ala Ser Thr Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

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

Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Tyr Ser Tyr Pro Phe
 85 90 95

Thr Phe Gly Pro Gly Thr Lys Val Asp Ile Lys
 100 105

<210> 336
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 336

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu Tyr Ser
20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln Ser
35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
85 90 95

Leu Gln Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
100 105 110

<210> 337

<211> 106

<212> PRT

<213> Homo sapiens

<400> 337

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser Trp
20 25 30

Leu Ala Trp Tyr Gln Arg Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
35 40 45

Tyr Lys Ala Ser Thr Leu Glu Ser Gly Val Pro Ser Arg Phe Ser Gly
50 55 60

Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
65 70 75 80

Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Phe Asn Ser Tyr Trp Thr
85 90 95

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
100 105

<210> 338

<211> 106

<212> PRT

<213> Homo sapiens

<400> 338

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser Trp
20 25 30

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
35 40 45

Tyr Lys Ala Ser Ser Leu Glu Ser Gly Val Pro Ser Arg Phe Ser Gly
50 55 60

Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
65 70 75 80

Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ser Tyr Trp Thr
85 90 95

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
100 105

<210> 339

<211> 113

<212> PRT

<213> Homo sapiens

<400> 339

Asp Ile Val Met Thr Gln Thr Pro Leu Ser Leu Pro Val Thr Pro Gly
1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu Asp Ser
20 25 30

Asp Asp Gly Asn Thr Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln
35 40 45

Ser Pro Gln Leu Leu Ile Tyr Thr Leu Ser Tyr 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

Arg Ile Glu Phe Pro Ser Thr Phe Gly Gln Gly Thr Arg Leu Glu Ile
100 105 110

Lys

<210> 340
 <211> 106
 <212> PRT
 <213> Homo sapiens

<400> 340

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Asp Trp
 20 25 30

Leu Ala Trp Tyr Leu Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Lys Ala Ser Ser Leu Glu Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

Ser Gly Ser Gly Thr Ala Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
 65 70 75 80

Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ser His Trp Thr
 85 90 95

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 100 105

<210> 341
 <211> 108
 <212> PRT
 <213> Homo sapiens

<400> 341

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Ser 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 Gly Ala Ser Ser 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 Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 85 90 95

Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile Lys
 100 105

<210> 342
 <211> 107
 <212> PRT
 <213> Homo sapiens
 <400> 342

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

Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

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

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

Glu Asp Ile Ala Thr Tyr Tyr Cys Gln Gln Tyr Asp Asn Leu Pro Leu
 85 90 95

Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
 100 105

<210> 343
 <211> 108
 <212> PRT
 <213> Homo sapiens
 <400> 343

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser 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 Gly Ala Ser Ser 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 Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro

Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile Lys
 100 105

<210> 344
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 344

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

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

Ser Asn Asn Lys Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
 50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 65 70 75 80

Ile Ser Ser Leu Gln Ala Glu Asp Glu Ala Val Tyr Tyr Cys Gln Gln
 85 90 95

Tyr Tyr Tyr Thr Pro Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile
 100 105 110

Lys

<210> 345
 <211> 108
 <212> PRT
 <213> Homo sapiens

<400> 345

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser 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 Gly Ala Ser Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser
 50 55 60

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Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu
65 70 75 80

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Glu Glu Tyr Gly Ser Ser Pro
85 90 95

Tyr Thr Phe Gly Glu Gly Thr Lys Leu Glu Ile Lys
100 105

<210> 346
<211> 113
<212> PRT
<213> Homo sapiens
<400> 346

Asp Ile Val Met Thr Glu Ser Pro Asp Ser Leu Ala Val Ser Leu Gly
1 5 10 15

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

Ser Asn Asn Lys Asn Tyr Leu Ser Trp Tyr Glu Glu Lys Pro Gly Glu
35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
65 70 75 80

Ile Ser Ser Leu Glu Ala Glu Asp Val Ala Val Tyr Tyr Cys Glu Glu
85 90 95

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

Lys

<210> 347
<211> 108
<212> PRT
<213> Homo sapiens
<400> 347

Glu Ile Val Leu Thr Glu Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
1 5 10 15

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Glu Ser Val Ser Ser Ser
20 25 30

Tyr Leu Ala Trp Tyr Glu Glu Lys Pro Gly Glu Ala Pro Arg Leu Leu

35

I l e T y r G l y A l a S e r S e r A r g A l a T h r G l y I l e P r o A s p A r g P h e S e r
50 55 60

G l y S e r G l y S e r G l y T h r A s p P h e T h r L e u T h r I l e S e r A r g L e u G l u
65 70 75 80

P r o G l u A s p P h e A l a V a l T y r T y r C y s G l n G l n T y r G l y S e r S e r L e u
85 90 95

P h e T h r P h e G l y P r o G l y T h r L y s V a l A s p I l e L y s
100 105

<210> 348
<211> 113
<212> PRT
<213> Homo sapiens

<400> 348

A s p I l e V a l M e t T h r G l n S e r P r o A s p S e r L e u A l a V a l S e r L e u G l y
1 5 10 15

G l u A r g A l a T h r I l e A s n C y s L y s S e r S e r G l n S e r V a l L e u T y r S e r
20 25 30

S e r A s n A s n L y s A s n T y r L e u A l a T r p T y r G l n G l n L y s P r o G l y G l n
35 40 45

P r o P r o L y s V a l L e u I l e T y r T r p A l a S e r I l e A r g G l u S e r G l y V a l
50 55 60

S e r A s p A r g P h e S e r G l y S e r G l y S e r G l y T h r A s p P h e T h r L e u T h r
65 70 75 80

I l e S e r S e r L e u G l n A l a G l u A s p V a l A l a V a l T y r T y r C y s G l n G l n
85 90 95

T y r T y r S e r T h r P r o T y r T h r P h e G l y G l n G l y T h r L y s L e u G l u I l e
100 105 110

Lys

<210> 349
<211> 113
<212> PRT
<213> Homo sapiens

<400> 349

A s p I l e V a l M e t T h r G l n S e r P r o A s p S e r L e u A l a V a l S e r L e u G l y
1 5 10 15

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Glu Arg Ala Thr Ile Asn Cys Lys Ser Ser Gln Thr Val Leu Tyr Ser
 20 25 30

Ser Asp Asn Lys Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
 50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 65 70 75 80

Ile Ser Ser Leu Gln Ala Glu Asp Val Ala Val Tyr Tyr Cys Gln Gln
 85 90 95

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

Lys

<210> 350
 <211> 108
 <212> PRT
 <213> Homo sapiens
 <400> 350

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Ile Ser 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 Gly Ala Ser Ser 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 Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 85 90 95

Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile Lys
 100 105

<210> 351
 <211> 106
 <212> PRT
 <213> Homo sapiens

<400> 351

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Ser Ile Ser Ser Trp
20 25 30

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
35 40 45

Tyr Lys Ala Ser Ser Leu Glu Ser Gly Val Pro Ser Arg Phe Ser Gly
50 55 60

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

Asp Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ser Tyr Trp Thr
85 90 95

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
100 105

<210> 352

<211> 107

<212> PRT

<213> Homo sapiens

<400> 352

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser Tyr
20 25 30

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
35 40 45

Tyr Ala Ala Ser Thr Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
50 55 60

Ser Gly Ser Gly Thr Glu Phe Thr Leu Thr Ile Ser Ser Leu Gln Pro
65 70 75 80

Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Leu Asn Thr Tyr Pro Arg
85 90 95

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
100 105

<210> 353

<211> 107

<212> PRT
 <213> Homo sapiens
 <400> 353

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

<210> 354
 <211> 112
 <212> PRT
 <213> Homo sapiens
 <400> 354

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

<210> 355
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 355

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Glu Trp Tyr Leu Gln Lys Pro Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
 85 90 95

Leu Gln Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
 100 105 110

<210> 356
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 356

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Ser Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
 85 90 95

Leu Gln Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
 100 105 110

<210> 357
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 357

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

Glu Arg Ala Thr Ile Asn Cys Lys Ser Ser Gln Ser Val Leu Tyr Ser
 20 25 30

Ser Asn Asn Lys Asn Tyr Leu Thr Trp Tyr Gln Gln Lys Pro Gly Gln
 35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
 50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 65 70 75 80

Ile Ser Ser Leu Gln Ala Glu Asp Val Ala Val Tyr Tyr Cys Gln Gln
 85 90 95

Tyr Tyr Ser Thr Pro Pro Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 100 105 110

Lys

<210> 358
 <211> 107
 <212> PRT
 <213> Homo sapiens

<400> 358

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

Leu Asn Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

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

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

Glu Asp Ile Ala Thr Tyr Tyr Cys Gln Gln Tyr Asp Asn Leu Pro Ile
 85 90 95

Thr Phe Gly Gln Gly Thr Arg Leu Glu Ile Lys
 100 105

<210> 359
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 359

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
 85 90 95

Leu Gln Thr Pro Pro Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile
 100 105 110

Lys

<210> 360
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 360

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
85 90 95

Leu Gln Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
100 105 110

<210> 361
<211> 113
<212> PRT
<213> Homo sapiens
<400> 361

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

Glu Arg Ala Thr Ile Asn Cys Lys Ser Ser Gln Ser Val Leu Tyr Ser
20 25 30

Ser Asn Asn Lys Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
65 70 75 80

Ile Ser Ser Leu Gln Ala Glu Asp Val Ala Val Tyr Tyr Cys Gln Gln
85 90 95

Tyr Tyr Arg Thr Met Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile
100 105 110

Lys

<210> 362
<211> 112
<212> PRT
<213> Homo sapiens
<400> 362

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
20 25 30

Asn Gly Tyr 35 Asn Tyr Leu Asp 40 Trp Ser Leu Gln Lys 45 Pro Gly Gln Ser

Pro Gln Leu Leu Ile Tyr 55 Leu Gly Ser Asn Arg Ala 60 Ser Gly Val Pro

Asp Arg Phe Ser Gly 70 Ser Gly Ser Gly Thr 75 Asp Phe Thr Leu Lys 80 Ile

Ser Arg Val Glu 85 Ala Glu Asp Val Gly 90 Val Tyr Tyr Cys Met 95 Gln Thr

Leu Gln Ile 100 Pro Tyr Thr Phe Gly 105 Gln Gly Thr Lys Leu Glu 110 Ile Lys

<210> 363
<211> 113
<212> PRT
<213> Homo sapiens
<400> 363

Asp Ile Val 1 Met Thr 5 Gln Thr Pro Leu Ser 10 Leu Pro Val Thr 15 Pro Gly

Glu Pro Ala 20 Ser Ile Ser Cys Arg 25 Ser Ser Gln Ser Leu 30 Leu Asp Ser

Asp Asp Gly 35 Asn Thr Tyr Leu 40 Asp Trp Tyr Leu Gln 45 Lys Pro Gly Gln

Ser Pro Gln Leu Leu Ile Tyr 55 Thr Leu Ser Tyr Arg 60 Ala Ser Gly Val

Pro Asp Arg Phe Ser 70 Gly Ser Gly Ser Gly Thr 75 Asp Phe Thr Leu Lys 80

Ile Ser Arg Val 85 Glu Ala Glu Asp Val 90 Gly Val Tyr Tyr Cys 95 Met Gln

Arg Ile Glu 100 Phe Pro Leu Thr Phe 105 Gly Gly Gly Thr Lys 110 Val Glu Ile

Lys

<210> 364
<211> 113
<212> PRT
<213> Homo sapiens
<400> 364

Asp Ile Val 1 Met Thr 5 Gln Thr Pro Leu Ser 10 Leu Pro Val Thr 15 Pro Gly

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu Asp Ser
 20 25 30

Asp Asp Gly Asn Thr Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln
 35 40 45

Ser Pro Gln Leu Leu Ile Tyr Thr Leu Ser Tyr 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

Arg Ile Glu Phe Pro Ile Thr Phe Gly Gln Gly Thr Arg Leu Glu Ile
 100 105 110

Lys

<210> 365
 <211> 113
 <212> PRT
 <213> Homo sapiens
 <400> 365

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

Glu Arg Ala Thr Ile Asn Cys Lys Ser Ser Gln Ser Val Leu Tyr Ser
 20 25 30

Ser Asn Asn Lys Asn Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 35 40 45

Pro Pro Lys Leu Leu Ile Tyr Trp Ala Ser Thr Arg Glu Ser Gly Val
 50 55 60

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 65 70 75 80

Ile Ser Ser Leu Gln Ala Glu Asp Val Ala Val Tyr Tyr Cys Gln Gln
 85 90 95

Tyr Tyr Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Glu Leu Trp
 100 105 110

Leu

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<210> 366
 <211> 107
 <212> PRT
 <213> Homo sapiens

<400> 366

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

Asp Arg Val Thr Ile Thr Cys Arg Ala Ser Gln Gly Ile Ser Ser Trp
 20 25 30

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Lys Ala Pro Lys Leu Leu Ile
 35 40 45

Tyr Ala Ala Ser Ser Leu Gln Ser Gly Val Pro Ser Arg Phe Ser Gly
 50 55 60

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

Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Ala Asn Thr Phe Pro Phe
 85 90 95

Thr Val Gly Pro Gly Thr Lys Val Asp Ile Lys
 100 105

<210> 367
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 367

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Leu Gln Ala
 85 90 95

Leu Gln Thr Pro Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys

100

105

110

<210> 368
 <211> 113
 <212> PRT
 <213> Homo sapiens

<400> 368

Asp Ile Val Met Thr Gln Thr Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu Asp Ser
 20 25 30

Asp Asp Gly Asn Thr Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln
 35 40 45

Ser Pro Gln Leu Leu Ile Tyr Thr Leu Ser Tyr 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

Arg Ile Glu Phe Pro Tyr Thr Phe Gly Gln Gly Thr Lys Leu Glu Ile
 100 105 110

Lys

<210> 369
 <211> 112
 <212> PRT
 <213> Homo sapiens

<400> 369

Asp Ile Val Met Thr Gln Ser Pro Leu Ser Leu Pro Val Thr Pro Gly
 1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gln Ser Leu Leu His Ser
 20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gln Lys Pro Gly Gln Ser
 35 40 45

Pro Gln Leu Leu Ile Tyr Leu Ala Ser Asn Arg Ala Ser Gly Val Pro
 50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
 65 70 75 80

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Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gln Ala
 85 90 95

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

<210> 370
 <211> 108
 <212> PRT
 <213> Homo sapiens

<400> 370

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Asn Val Ser Ser Thr
 20 25 30

Tyr Leu Ala Trp Tyr His Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
 35 40 45

Ile Tyr Gly Ala Ser Ser 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 Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 85 90 95

Arg Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 100 105

<210> 371
 <211> 108
 <212> PRT
 <213> Homo sapiens

<400> 371

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser 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 Gly Ala Ser Ser 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 Ala Val Tyr Tyr Cys Gl n Gl n Tyr Gly Asn Ser Pro
85 90 95

Leu Thr Phe Gly Gly Gly Thr Lys Val Glu Ile Lys
100 105

<210> 372
<211> 112
<212> PRT
<213> Homo sapiens

<400> 372

Asp Ile Val Met Thr Gl n Ser Pro Ile Ser Leu Pro Val Thr Pro Gly
1 5 10 15

Glu Pro Ala Ser Ile Ser Cys Arg Ser Ser Gl n Ser Leu Leu Tyr Ser
20 25 30

Asn Gly Tyr Asn Tyr Leu Asp Trp Tyr Leu Gl n Lys Pro Gly Gl n Ser
35 40 45

Pro Gl n Leu Leu Ile Tyr Leu Gly Ser Asn Arg Ala Ser Gly Val Pro
50 55 60

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Lys Ile
65 70 75 80

Ser Arg Val Glu Ala Glu Asp Val Gly Val Tyr Tyr Cys Met Gl n Ala
85 90 95

Leu Gl n Thr Pro Trp Thr Phe Gly Gl n Gly Thr Lys Val Glu Ile Lys
100 105 110

<210> 373
<211> 110
<212> PRT
<213> Homo sapiens

<400> 373

Gl n Ser Val Leu Thr Gl n Pro Pro Ser Ala Ser Gly Thr Pro Gly Gl n
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 Gl n Gl n Leu Pro Gly Thr Ala Pro Lys Leu Leu
35 40 45

Ile Tyr Ser Asn Asn Gl n Arg Pro Ser Gly Val Pro Asp Arg Phe Ser
50 55 60

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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

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

<210> 374
<211> 110
<212> PRT
<213> Homo sapiens
<400> 374

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 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 80

Ser Glu Asp Glu Ala Asp Tyr Asp Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

<210> 375
<211> 110
<212> PRT
<213> Homo sapiens
<400> 375

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 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 Gly Arg Phe Ser
Page 150

50

55

60

Gly Ser Lys Ser Gly Thr Ser Ala Ser Leu Ala Ile Ser Gly Leu Gln
65 70 75 80

Ser Asp Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

<210> 376
<211> 110
<212> PRT
<213> Homo sapiens

<400> 376

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 Asn
20 25 30

Ser Val Asn Trp Phe 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

<210> 377
<211> 110
<212> PRT
<213> Homo sapiens

<400> 377

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 Asn
20 25 30

Thr Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu
35 40 45

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I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Pro Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

<210> 378
<211> 110
<212> PRT
<213> Homo sapi ens

<400> 378

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Ser Asn I l e G l y Ser Asn
20 25 30

Al a Val Asn Trp Tyr G l n G l n Leu Pro G l y Thr Al a Pro Lys Leu Leu
35 40 45

I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Pro Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

<210> 379
<211> 110
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<213> Homo sapi ens

<400> 379

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Ser Asn I l e G l y Ser Asn
20 25 30

Thr Val Asn Trp Tyr G l n G l n Phe Pro G l y Thr Al a Pro Lys Leu Leu
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I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Val Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

<210> 380
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<212> PRT
<213> Homo sapi ens

<400> 380

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Ser Asn I l e G l y Asn Asn
20 25 30

Al a Val Asn Trp Tyr G l n G l n Leu Pro G l y Thr Al a Pro Lys Leu Leu
35 40 45

I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Pro Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

<210> 381
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<213> Homo sapi ens

<400> 381

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Ser Asn I l e G l y Ser Asn
20 25 30

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Thr Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Asn 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 80

Ser Glu Asp Glu Ala Asp Tyr Phe Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 382
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<400> 382

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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Pro Val Phe Gly Gly Gly Thr Asn Leu Thr Val Leu
 100 105 110

<210> 383
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<400> 383

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 Gly Ser Ser Asn Ile Gly Ser Asn
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20

25

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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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

<210> 384
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<400> 384

Gln Ser Val Leu Thr Gln Ser 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 80

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 Thr Gly Thr Lys Val Thr Val Leu
 100 105 110

<210> 385
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<400> 385

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

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp His Asp Ser Leu
85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

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<400> 386

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 Asn
20 25 30

Thr Val Asn Trp Tyr Gln Gln Phe 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

Asn Gly Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105

<210> 387
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<400> 387

Gln Ser Val Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln
Page 156

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 Phe Gl n Gl n Leu Pro Gly Thr Ala Pro Lys Leu Leu
 35 40 45

Ile Tyr Ser Asn Asn Gl n 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 Gl n
65 70 75 80

Ser Gl u Asp Gl u Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 388
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<400> 388

Gl n Ser Val Leu Thr Gl n Pro Pro Ser Ala Ser Gly Thr Pro Gly Gl n
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 Gl n Gl n Leu Pro Gly Thr Ala Pro Lys Leu Leu
 35 40 45

Ile Tyr Ser Asn Asn Gl n 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 Gl n
65 70 75 80

Ser Gl u Asp Gl u Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 389
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<400> 389

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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 Leu
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 Ser
85 90 95

Ser Thr Phe Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

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<400> 390

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 Asn
20 25 30

Thr Val Asn Trp Tyr Gln Gln Val 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

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<213> Homo sapiens

<400> 391

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 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 80
 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 Thr Gly Thr Lys Val Thr Val Leu
 100 105 110

<210> 392
 <211> 110
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<400> 392

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Gly Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

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<212> PRT
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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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Gly Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 394
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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 Leu
 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 Ser
 85 90 95
 Ser Thr Phe Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

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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 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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 397
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 <212> PRT
 <213> Homo sapiens

<400> 397

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 398
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 <213> Homo sapiens

<400> 398

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Phe Cys Ala Ala Trp His Asp Ser Leu
 85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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Gln Ser Val Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln
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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 80

Ser Glu Asp Glu Ala Asp Tyr Ser Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 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 Phe Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

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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 Asn
20 25 30

Tyr Val Tyr 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 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 Leu Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

<210> 407
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<400> 407

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 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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

Ser Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

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<213> Homo sapiens

<400> 408

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 Asn
20 25 30

Thr Val Asn Trp Tyr Gln Leu 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

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Gly Ser Gln 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 Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

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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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

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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 Asn
20 25 30

Tyr Val Tyr Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Lys Leu Leu
35 40 45

I l e Ser Ser Asn Asn G l n Arg Pro Ser Gly Val Pro Asp Arg Phe Ser
 50 55 60

Gly Ser Lys Ser Gly Thr Ser Al a Ser Leu Al a I l e Ser Gly Leu Arg
 65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
 85 90 95

Ser Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

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G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser Gly Thr Pro Gly G l n
 1 5 10 15

Arg Phe Thr I l e Ser Cys Ser Gly Ser Ser Ser Asn I l e Gly Ser Asn
 20 25 30

Thr Val Asn Trp Tyr G l n G l n Leu Pro Gly Thr Al a Pro Lys Leu Leu
 35 40 45

I l e Tyr Ser Asn Asn G l n Arg Pro Ser Gly Val Pro Asp Arg Phe Ser
 50 55 60

Gly Ser Lys Ser Gly Thr Ser Al a Ser Leu Al a Ser Ser Gly Leu G l n
 65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Arg Leu
 85 90 95

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

<210> 412
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 <400> 412

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser Gly Thr Pro Gly G l n
 1 5 10 15

Arg Val Thr I l e Ser Cys Ser Gly Ser Ser Ser Asn I l e Gly Ser Asn
 20 25 30

Thr Val Asn Trp Tyr G l n G l n Phe Pro Gly Thr Al a Pro Lys Leu Leu
 35 40 45

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I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Pro Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

<210> 413
<211> 110
<212> PRT
<213> Homo sapiens

<400> 413

G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Ser Asn I l e G l y Ser Asn
20 25 30

Thr Val Asn Trp Tyr G l n G l n Leu Pro G l y Thr Al a Pro Lys Leu Leu
35 40 45

I l e Tyr Ser Asn Asn G l n Arg Pro Ser G l y Val Pro Asp Arg Phe Ser
50 55 60

G l y Ser Lys Ser G l y Thr Ser Al a Ser Leu Al a I l e Ser G l y Leu G l n
65 70 75 80

Ser Gl u Asp Gl u Al a Asp Tyr Tyr Cys Al a Al a Trp Asp Asp Ser Leu
85 90 95

Asn G l y Pro Val Phe G l y G l y G l y Thr Lys Leu Thr Val Leu
100 105 110

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G l n Ser Val Leu Thr G l n Pro Pro Ser Al a Ser G l y Thr Pro G l y G l n
1 5 10 15

Arg Val Thr I l e Ser Cys Ser G l y Ser Ser Phe Asn I l e G l y G l y 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

<210> 415
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<400> 415

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 Asn
 20 25 30

Thr Val Thr 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

<210> 416
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<400> 416

Ser Ser Glu Leu Thr Gln Asp Pro Ala Val Ser Val Ala Leu Gly Gln
 1 5 10 15

Thr Val Arg Ile Thr Cys Gln Gly Asp Ser Leu Arg Ser Tyr Tyr Ala
 20 25 30

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Ser Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Ile 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

Ser Ser Gly Asn 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 Gln Asp Asn Ser Gly Asn His
85 90 95

Leu Val Val Phe Gly Arg Gly Thr Lys Leu Thr Val Leu
100 105

<210> 417
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<400> 417

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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
85 90 95

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

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<400> 418

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 Asn Ser Asn Ile Gly Ser Asn
 20 25 30

Thr Val Asn Trp Tyr Gln Gln Leu Pro Gly Thr Ala Pro Arg 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 80

Ser Glu Asp Glu Ala Asp Ser Tyr Cys Ala Ala Trp Tyr Asp Ser Leu
 85 90 95

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

<210> 419
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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 Asn
 20 25 30

Thr Val Asn Trp Tyr Gln His 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 80

Ser Glu Asp Glu Ala Asp Tyr Phe Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Pro Val Phe Gly Gly Gly Thr Arg Val Thr Val Leu
 100 105 110

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<400> 420

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

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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 Asn
20 25 30

Tyr Val Tyr 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 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 Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

<210> 422
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<213> Homo sapiens
<400> 422

Gln Ser Val Leu Thr Gln Pro Pro Ser Val 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 80

Ser Glu Asp Glu Ala Asp Tyr Asp Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

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

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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 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 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> 424
 <211> 110
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<400> 424

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Gly Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 425
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<400> 425

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Gly Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 426
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<213> Homo sapiens

<400> 426

Ser Tyr Glu Leu Thr Gln Pro His Ser Val Ser Val Ala Thr Ala Gln
1 5 10 15

Met Ala Arg Ile Thr Cys Gly Gly Asn Asn Ile Gly Ser Lys Ala Val
20 25 30

His Trp Tyr Gln Gln Lys Pro Gly Gln Asp Pro Val Leu Val Ile Tyr
35 40 45

Ser Asp Ser Asn Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
50 55 60

Asn Pro Gly Asn Thr Ala Thr Leu Thr Ile Ser Arg Ile Glu Ala Gly
65 70 75 80

Asp Glu Ala Asp Tyr Tyr Cys Gln Val Trp Asp Ser Ser Ser Asp His
85 90 95

Arg Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105

<210> 427

<211> 110

<212> PRT

<213> Homo sapiens

<400> 427

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 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Pro Trp Asp Asp Ser Leu
85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
100 105 110

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<400> 428

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 Asn
 20 25 30
 Thr Val Asn Trp Tyr Gln Gln Val 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95
 Asn Ser Trp Val Phe Gly Gly Gly Thr Met Leu Thr Val Leu
 100 105 110

<210> 429
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 <213> Homo sapiens

<400> 429

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 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 80
 Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Arg Asp Asp Ser Leu
 85 90 95
 Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

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<210> 430
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Gln Ser Val Leu Thr Gln Pro Pro Ser Ala Ser Gly Thr Pro Gly Gln
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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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Arg Asp Asp Ser Leu
 85 90 95

Asn Gly Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

<210> 431
 <211> 110
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 <213> Homo sapiens

<400> 431

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 Thr Ser Ser Asn Ile Gly Ser Asn
 20 25 30

Thr Val His 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 80

Ser Glu Asp Glu Ala Asp Tyr Tyr Cys Ala Ala Trp Asp Asp Ser Leu
 85 90 95

Asn Gly Pro Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
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105

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<400> 432

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 Asn
 20 25 30

Thr Val Asn Trp Tyr Gln Arg Val Pro Gly Thr Ala Pro Lys Leu Leu
 35 40 45

Ile Tyr Ser 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 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 Val Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu
 100 105 110

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<400> 433
 ggttcgggga agtagtcctt gacc

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<220>
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29

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<223> Synthesi zed pri mer

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Gly Phe Thr Phe Ser Asn Ala Trp
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Gly Tyr Thr Phe Thr Gly Tyr Tyr
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Gly Phe Thr Phe Ser Asn Ala Trp
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Thr Phe Gly Asp Tyr Ala
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Gly Tyr Thr Phe Thr Gly Tyr Tyr
 1 5

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Gly Phe Thr Phe Ser Asn Ala Trp
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Gly Gly Ser Ile Ser Ser Ser Tyr
 1 5

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Arg Phe Thr Phe Ser Ser Tyr Ser
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Gly Phe Ser Leu Ser Thr Ser Gly Met Cys
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Tyr Thr Phe Thr Gly Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Thr Phe Ser Ser Tyr Asp
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Gly Gly Ser Cys Ser Ala Tyr Tyr
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Gly Phe Thr Phe Ser Asn Ala Trp
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Gly Gly Ser Phe Ser Ala Tyr Tyr
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Gly Phe Thr Phe Ser Ser Tyr Trp
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Gly Phe Thr Phe Ser Ser Tyr Ser
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Gly Phe Ser Leu Ser Thr Ser Gly Val Gly
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Gly Tyr Thr Phe Thr Gly Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Thr Phe Ser Ser Tyr Asn
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Thr Phe Ser Ser Tyr Ser
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Gly Phe Thr Phe Gly Asp Tyr Ala
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Gly Phe Thr Phe Ser Asn Ala Trp
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Gly Tyr Thr Phe Thr Gly Tyr Tyr
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Gly Phe Thr Phe Ser Ser Tyr Ser
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Pro Phe Ser Val Tyr Trp
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Gly Phe Thr Phe Ser Asn Ala Trp
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Gly Phe Thr Phe Ser Ser Tyr Ser
1 5

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Gly Tyr Thr Phe Thr Gly Tyr Tyr
1 5

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Gly Tyr Thr Phe Thr Gly Tyr Tyr
1 5

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Gly Phe Ser Leu Ser Thr Ser Gly Val Gly
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Gly Phe Thr Phe Ser Ser Tyr Gly
1 5

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Gly Phe Thr Phe Ser Thr Tyr Trp

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Gly Tyr Thr Phe Thr Gly Tyr Tyr
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Gly Phe Thr Phe Ser Thr Phe Ala
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Gly Ala Ser Ile Ser Thr Tyr Phe
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Phe Thr Leu Ser Ser Tyr Gly
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Ser Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Tyr Thr Phe Thr Asp Tyr Tyr
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Gly Gly Ser Ile Ser Ser Phe Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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Gly Phe Thr Phe Ser Ser Tyr Ser
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Gly Asp Ser Ile Ser Asn Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
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Gly Gly Ser Ile Ser Ser Tyr Tyr
1 5

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