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(54) **Title**: ANTI-MST1R ANTIBODIES AND USES THEREOF

(57) **Abstract**: The present disclosure provides recombinant antigen-binding regions and antibodies and functional fragments containing such antigen-binding regions that are specific for MST1R, which plays an integral role in various disorders or conditions, such as cancer. These antibodies, accordingly, can be used to treat these and other disorders and conditions. Antibodies of the disclosure also can be used in the diagnostics field, as well as for further investigating the role of MST1R in the progression of disorders associated with tumors. The disclosure also provides nucleic acid sequences encoding the foregoing antibodies, vectors containing the same, pharmaceutical compositions and kits with instructions for use.

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DESCRIPTION

Title of Invention

ANTI-MST1R ANTIBODIES AND USES THEREOF

Background

[0001]

MST1R (macrophage stimulating 1 receptor; human MST1R is shown in GenBank accession No; NM_002447.2), also described as RON or CDw136, is a c-Met related tyrosine kinase found on cells of epithelial origin. The 1400 amino acid single chain precursor is cleaved into a disulfide-linked heterodimer consisting of an extracellular 40kDa α-chain and a 150kDa β-chain, which includes the intracellular tyrosine kinase domain. Similar to c-Met, MST1R induces invasive cell growth, migration, cell dissociation and matrix invasion. [Wang, et al., *Carcinogenesis* 24, 1291-1300, 2003; Lee, et al., *Clin Cancer Res* 11, 2222-2228, 2005]. Both tyrosine kinases are overexpressed on a variety of malignant tumors, such as breast, lung or prostate cancer [O'Toole, et al., *Cancer Res* 66, 9162-9170, 2006]. MSP, macrophage stimulatory protein, is the only ligand to MST1R known so far. MSP binding triggers autophosphorylation of the MST1R tyrosine kinase domain. Thereby activated MST1R transduces a variety of different pathway cascades. [Wang, et al., *Carcinogenesis* 24, 1291-1300, 2003; O'Toole, et al., *Cancer Res* 66, 9162-9170, 2006]. Generation of biologically active, truncated MST1R variants through mRNA splicing has also been reported [Wang, et al., *Carcinogenesis* 24, 1291-1300, 2003]. For example, MST1R⊿160 variant was found in some of colorectal carcinoma samples, and its overexpression without ligand mediated tumor formation in nude mice [Zhou et al., *Oncogene* 22, 186-197, 2003]. Anti-MST1R antibodies like IMC-41A10 block the ligand-receptor interaction and are potent inhibitors of receptor and downstream signaling, cell migration and tumorigenesis [O'Toole, et al., *Cancer Res* 66, 9162-9170, 2006].

[0002]

In conclusion, antibodies blocking MST1R activity are of potential therapeutic relevance in human cancer.

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Summary

[0003]

In an aspect the present invention provides human and humanized antibodies against MST1R.

[0004]

In another aspect the present invention provides antibodies that are safe for human administration.

[0005]

In an aspect the present invention provides methods for treating disease or and/or conditions associated with MST1R up-regulation by using one or more antibodies of the invention. These and other aspects are more fully described herein.

[0006]

In one embodiment, an isolated antibody or functional fragment that contains an antigen-binding region is specific for MST1R.

[0007]

Such an antibody or functional fragment thereof may contain an antigen-binding region that contains an H-CDR3 (heavy chain CDR3) region having the amino acid sequence of SEQ ID NO: 1 or 4; the antigen-binding region may further include an H-CDR2 (heavy chain CDR2) region having the amino acid sequence of SEQ ID NO: 2 or 5; and the antigen-binding region also may contain an H-CDR1 (heavy chain CDR1) region having the amino acid sequence of SEQ ID NO: 3 or 6. Such an antibody or functional fragment thereof may contain an antigen-binding region that contains an L-CDR3 (light chain CDR3) region having the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12; the antigen-binding region may further include an L-CDR1 (light chain CDR1) region having the amino acid sequence of SEQ ID NO: 13 or 15; and the antigen-binding region also may contain an L-CDR2 (light chain CDR2) region having the amino acid sequence of SEQ ID NO: 14 or 16.

[0008]

Antibodies (and functional fragments thereof) described herein may contain an antigen-binding region that is specific for an epitope of MST1R, which epitope contains one or more amino acid

residues of amino acid having the amino acid sequence of SEQ ID NO: 17. For certain antibodies, the epitope may be linear, whereas for others, it may be conformational (*i.e.*, discontinuous). An antibody or functional fragment thereof having one or more of these properties may contain an antigen-binding region that contains an H-CDR3 region having the amino acid sequence of SEQ ID NO: 1 or 4; the antigen-binding region may further include an H-CDR2 region having the amino acid sequence of SEQ ID NO: 2 or 5; and the antigen-binding region also may contain an H-CDR1 region having the amino acid sequence of SEQ ID NO: 3 or 6. Such a MST1R -specific antibody of the invention may contain an antigen-binding region that contains an L-CDR3 region having the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12; the antigen-binding region may further include an L-CDR1 region shown in SEQ ID NO: 13 or 15; and the antigen-binding region also may contain an L-CDR2 region having the amino acid sequence of SEQ ID NO: 14 or 16.

[0009]

Peptide variants of the sequences disclosed herein are also embraced by various embodiments of the disclosure. Accordingly, the embodiments include anti-MST1R antibodies having a heavy chain amino acid sequence with: at least 60 percent sequence identity in the CDR regions with the CDR regions having the amino acid sequence of SEQ ID NO: 1, 2, 3, 4, 5 or 6; and/or at least 80 percent sequence homology in the CDR regions with the CDR regions having the amino acid sequence of SEQ ID NO: 1, 2, 3, 4, 5 or 6. Further included are anti-MST1R antibodies having a light chain amino acid sequence with: at least 60 percent sequence identity in the CDR regions with the CDR regions having the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11, 12, 13, 14, 15 or 16; and/or at least 80 percent sequence homology in the CDR regions with the CDR regions having the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11, 12, 13, 14, 15 or 16.

[0010]

An antibody disclosed herein may be an IgG (*e.g.*, IgG_1), while an antibody fragment may be a Fab or scFv, for example. An inventive antibody fragment, accordingly, may be, or may contain, an antigen-binding region that behaves in one or more ways as described herein.

[0011]

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Another embodiment also relates to isolated nucleic acid sequences, each of which can encode an antigen-binding region of a human antibody or functional fragment thereof that is specific for an epitope of MST1R. Such a nucleic acid sequence may encode a variable heavy chain of an antibody and include a sequence selected from the group consisting of SEQ ID NOS: 18, 20 or a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of SEQ ID NO: 18 or 20. The nucleic acid might encode a variable light chain of an isolated antibody or functional fragment thereof, and may contain a sequence selected from the group consisting of SEQ ID NOS: 22, 24, 26, 28, 30, 32, or a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of SEQ ID NO: 22, 24, 26, 28, 30 or 32.

[0011a]

Another embodiment of the invention provides an isolated human or humanized antibody or functional fragment thereof comprising an antigen-binding region that is specific for a partial peptide of MST1R, having an amino acid sequence of SEQ ID NO: 17, wherein one of the following requirements applies:

a. said antibody or functional fragment thereof inhibits ligand -dependent and/or -independent phosphorylation of MST1R;
b. said antibody or functional fragment thereof has an affinity against said partial peptide of MST1R as a KD of less than about 10 nM, less than about 5 nM, less than about 1 nM, less than about 0.5 nM or less than about 0.1 nM as determined by surface plasmon resonance, or by Solution Equilibrium Titration.
c. said antibody or functional fragment thereof suppresses MSP-promoted cell proliferation or migration of tumor cells that express MST1R;
d. said antibody or functional fragment thereof internalizes MST1R; or
e. said antibody or functional fragment thereof is cross-reactive with human and at least one other species and

wherein said antigen-binding region comprises

an H-CDR3 region having:

(i) the amino acid sequence of SEQ ID NO: 1 or 4, or
(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 1 or 4;

an H-CDR2 region having:

(i) the amino acid sequence of SEQ ID NO: 2 or 5, or

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(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 2 or 5;

an H-CDR1 region having:

(i) the amino acid sequence of SEQ ID NO: 3 or 6, or

(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 3 or 6;

an L-CDR3 region having:

(i) the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12, or

(ii) an amino acid sequence having at least 80% sequence identity with the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12;

an L-CDR1 region having:

(i) the amino acid sequence of SEQ ID NO: 13 or 15, or

(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 13 or 15; and

an L-CDR2 region having:

(i) the amino acid sequence of SEQ ID NO: 14 or 16, or

(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 14 or 16.

[0011b]

Another embodiment of the invention provides an isolated antibody having

a. a heavy chain having an amino acid sequence of SEQ ID NO: 49 and a light chain having an amino acid sequence of SEQ ID NO: 53; or

b. a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 55, 57, 59, 61 or 63.

[0011c]

A further embodiment of the invention provides a variable heavy chain of an isolated antibody or functional fragment thereof that is encoded by

(i) a nucleic acid sequence comprising SEQ ID NO: 18 or 20, or

(ii) a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of SEQ ID NO: 18 or 20, wherein said antibody or functional fragment thereof is specific for an epitope of MST1R.

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[0011d]

Another embodiment of the invention provides a variable light chain of an isolated antibody or functional fragment thereof that is encoded by

(i) a nucleic acid sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32, or

(ii) a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of a sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32, wherein said antibody or functional fragment thereof is specific for an epitope of MST1R.

[0012]

Nucleic acids described herein are suitable for recombinant production. Thus, vectors and host cells containing a nucleic acid sequence disclosed herein are also further embodiments.

[0013]

Compositions described herein may be used for therapeutic or prophylactic applications. These embodiments, therefore, include a pharmaceutical composition containing an inventive antibody (or functional antibody fragment) and a pharmaceutically acceptable carrier or excipient thereof. In a related aspect, another embodiment includes methods for treating a disorder or condition associated with the undesired presence of MST1R or MST1R expressing cells. Such method contains the steps of administering to a subject in need thereof an effective amount of the pharmaceutical composition that contains an inventive antibody as described or contemplated herein.

[0014]

Yet other embodiments relate to isolated epitopes of MST1R, either in linear or conformational form, and their use for the isolation of an antibody or functional fragment thereof, which antibody of antibody fragment comprises an antigen-binding region that is specific for said epitope. In this regard, a conformational epitope may contain one or more amino acid residues in SEQ ID NO: 17. An epitope of MST1 R can be used, for example, for the isolation of antibodies or functional fragments thereof (each of which antibodies or antibody fragments comprises an

antigen-binding region that is specific for such epitope), comprising the steps of contacting said epitope of MST1R with an antibody library and isolating the antibody(ies) or functional fragment(s) thereof.

[0015]

In another embodiment, the disclosure provides an isolated epitope of MST1R, which consists essentially of an amino acid sequence in SEQ ID NO: 17. As used herein, such an epitope "consists essentially of" one of the immediately preceding amino acid sequences plus additional features, provided that the additional features do not materially affect the basic and novel characteristics of the epitope.

[0016]

The disclosure is also directed to a kit having (i) an isolated epitope of MST1R comprising one or more amino acid residues of the amino acid sequence in SEQ ID NO: 17; (ii) an antibody library; and (iii) instructions for using the antibody library to isolate one or more members of such library that binds specifically to such epitope.

(DETAILED DESCRIPTION)

[0017]

The present disclosure is based on the discovery of novel antibodies that are specific to or have a high affinity for MST1R and can deliver a therapeutic benefit to a subject. The antibodies disclosed herein, which may be human or humanized, can be used in many contexts, which are more fully described herein.

[0018]

A "human" antibody or functional human antibody fragment is hereby defined as one that is not chimeric (*e.g.*, not "humanized") and not from (either in whole or in part) a non-human species. A human antibody or functional antibody fragment can be derived from a human or can be a synthetic human antibody. A "synthetic human antibody" is defined herein as an antibody having a sequence derived, in whole or in part, *in silico* from synthetic sequences that are based on the analysis of known human antibody sequences. *In silico* design of a human antibody sequence or fragment thereof can be achieved, for example, by analyzing a database of human

antibody or antibody fragment sequences and devising a polypeptide sequence utilizing the data obtained therefrom. Another example of a human antibody or functional antibody fragment, is one that is encoded by a nucleic acid isolated from a library of antibody sequences of human origin (*i.e.*, such library being based on antibodies taken from a human natural source).

[0019]

A "humanized antibody" or functional humanized antibody fragment is defined herein as one that is (i) derived from a non-human source (*e.g.*, a transgenic mouse which bears a heterologous immune system), which antibody is based on a human germline sequence; or (ii) chimeric, wherein the variable domain is derived from a non-human origin and the constant domain is derived from a human origin or (iii) complementarity determining regions (CDR)-grafted, wherein the CDRs of the variable domain are from a non-human origin, while one or more frameworks of the variable domain are of human origin and the constant domain (if any) is of human origin.

[0020]

As used herein, an antibody "binds specifically to," is "specific to/for" or "specifically recognizes" an antigen (here, MST1R) if such antibody is able to discriminate between such antigen and one or more reference antigen(s), since binding specificity is not an absolute, but a relative property. In its most general form (and when no defined reference is mentioned), "specific binding" is referring to the ability of the antibody to discriminate between the antigen of interest and an unrelated antigen, as determined, for example, in accordance with one of the following methods. Such methods comprise, but are not limited to Western blots, ELISA-, RIA-, ECL-, IRMA-tests and peptide scans. For example, a standard ELISA assay can be carried out. The scoring may be carried out by standard color development (e.g. secondary antibody with horseradish peroxide and tetramethyl benzidine with hydrogenperoxide). The reaction in certain wells is scored by the optical density, for example, at 450 nm. Typical background (=negative reaction) may be 0.1 OD; typical positive reaction may be 1 OD. This means the difference positive/negative can be more than 10-fold. Typically, determination of binding specificity is performed by using not a single reference antigen, but a set of about three to five unrelated antigens, such as milk powder, BSA, transferrin or the like.

[0021]

However, "specific binding" also may refer to the ability of an antibody to discriminate between the target antigen and one or more closely related antigen(s), which are used as reference points, e.g. between target MST1R and target semaphorin. Additionally, "specific binding" may relate to the ability of an antibody to discriminate between different parts of its target antigen, e.g. different domains or regions of MST1R, such as epitopes in the N-terminal or in the C-terminal region of target MST1R, or between one or more key amino acid residues or stretches of amino acid residues of target MST1R.

[0022]

Also, as used herein, an "immunoglobulin" (Ig) hereby is defined as a protein belonging to the class IgG, IgM, IgE, IgA, or IgD (or any subclass thereof), and includes all conventionally known antibodies and functional fragments thereof. A "functional fragment" of an antibody/immunoglobulin hereby is defined as a fragment of an antibody/immunoglobulin (e.g., a variable region of an IgG) that retains the antigen-binding region. An "antigen-binding region" of an antibody typically is found in one or more hypervariable region(s) of an antibody, i.e., the CDR-1, -2, and/or –3 regions; however, the variable "framework" regions can also play an important role in antigen binding, such as by providing a scaffold for the CDRs. In various embodiments, the "antigen-binding region" comprises at least amino acid residues 4 to 103 of the variable light (VL) chain and 5 to 109 of the variable heavy (VH) chain, amino acid residues 3 to 107 of VL and 4 to 111 of VH, and are the complete VL and VH chains (amino acid positions 1 to 109 of VL and 1 to 113 of VH; numbering according to WO 97/08320). A exemplary class of immunoglobulins for use in the embodiments described herein is IgG. "Functional fragments" of the invention include the domain of a $F(ab')_2$ fragment, a Fab fragment and scFv. The $F(ab')_2$ or Fab may be engineered to minimize or completely remove the intermolecular disulphide interactions that occur between the CH1 and CL domains.

[0023]

An antibody described herein may be derived from a recombinant antibody library that is based on amino acid sequences that have been designed in silico and encoded by nucleic acids that are synthetically created. *In silico* design of an antibody sequence is achieved, for example, by analyzing a database of human sequences and devising a polypeptide sequence utilizing the data obtained therefrom. Methods for designing and obtaining *in silico*-created sequences are

described, for example, in Knappik et al., *J. Mol. Biol.* 296: 57-86, 2000; Krebs et al., *J. Immunol. Methods*. 254:67-84, 2001; and U.S. Patent No. 6,300,064 issued to Knappik et al., which hereby are incorporated by reference in their entirety.

(Antibodies Described Herein)

[0024]

Throughout this disclosure, reference is made to the following representative antibodies: "antibody nos." or "LACS" or "MOR" X. MOR X represents an antibody having a variable heavy region corresponding to SEQ ID NO: 18 or 20 (DNA)/SEQ ID NO: 19 or 21 (protein) and a variable light region selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32 (DNA)/SEQ ID NOs: 23, 25, 27, 29, 31 and 33 (protein).

[0025]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 18 (DNA)/SEQ ID: NO: 19 (protein) and a variable light chain corresponding to SEQ ID NO: 22 (DNA)/SEQ ID NO: 23 (protein).

[0026]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 20 (DNA)/SEQ ID: NO: 21 (protein) and a variable light chain corresponding to SEQ ID NO: 24 (DNA)/SEQ ID NO: 25 (protein).

[0027]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 18 (DNA)/SEQ ID: NO: 19 (protein) and a variable light chain corresponding to SEQ ID NO: 26 (DNA)/SEQ ID NO: 27 (protein).

[0028]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 18 (DNA)/SEQ ID: NO: 19 (protein) and a variable light chain corresponding to SEQ ID NO: 28 (DNA)/SEQ ID NO: 29 (protein).

[0029]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 18 (DNA)/SEQ ID: NO: 19 (protein) and a variable light chain corresponding to SEQ ID NO: 30(DNA)/SEQ ID NO: 31 (protein).

[0030]

In one example, the disclosure provides an antibody having a variable heavy region corresponding to SEQ ID NO: 18 (DNA)/SEQ ID: NO: 19 (protein) and a variable light chain corresponding to SEQ ID NO: 32 (DNA)/SEQ ID NO: 33 (protein).

[0031]

In another aspect, the disclosure provides following antibodies.

[0032]

In one example, the disclosure provides an antibody containing an antigen-binding region that contains an H-CDR3 (heavy chain CDR3) region having the amino acid sequence of SEQ ID NO: 1 or 4; the antigen-binding region may further include an H-CDR2 (heavy chain CDR2) region having the amino acid sequence of SEQ ID NO: 2 or 5; and the antigen-binding region also may contain an H-CDR1 (heavy chain CDR1) region having the amino acid sequence of SEQ ID NO: 3 or 6. Such an antibody thereof may contain an antigen-binding region that contains an L-CDR3 (light chain CDR3) region having the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12; the antigen-binding region may further include an L-CDR1 (light chain CDR1) region having the amino acid sequence of SEQ ID NO: 13 or 15; and the antigen-binding region also may contain an L-CDR2 (light chain CDR2) region having the amino acid sequence of SEQ ID NO: 14 or 16.

[0033]

The disclosure also provides an antibody containing the antigen-binding region (i) H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2 and H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, (ii) H-CDR3 region having the amino acid sequence of SEQ ID NO: 4, H-CDR2 region having the amino acid sequence of SEQ ID NO: 5 and H-CDR1 region having the amino acid sequence of SEQ ID NO: 6.

[0034]

One embodiment also provides an antibody containing an antigen-binding region selected from the group consisting of (i) L-CDR3 region having the amino acid sequence of SEQ ID NO: 7, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (ii) L-CDR3 region having the amino acid sequence of SEQ ID NO: 8, L-CDR1 region having the amino acid sequence of SEQ ID NO: 15 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 16, (iii) L-CDR3 region having the amino acid sequence of SEQ ID NO: 9, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (iv) L-CDR3 region having the amino acid sequence of SEQ ID NO: 10, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (v) L-CDR3 region having the amino acid sequence of SEQ ID NO: 11, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, or (vi) L-CDR3 region having the amino acid sequence of SEQ ID NO: 12, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14.

[0035]

Another embodiment provides an antibody containing an antigen-binding region selected from the group consisting of (i) H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2, H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, L-CDR3 region having the amino acid sequence of SEQ ID NO: 7, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (ii) H-CDR3 region having the amino acid sequence of SEQ ID NO: 4, H-CDR2 region having the amino acid sequence of SEQ ID NO: 5, H-CDR1 region having the amino acid sequence of SEQ ID NO: 6, L-CDR3 region having the amino acid sequence of SEQ ID NO: 8, L-CDR1 region having the amino acid sequence of SEQ ID NO: 15 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 16,, (iii) H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2, H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, L-CDR3 region having the amino acid sequence of SEQ ID NO: 9, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (iv) H-CDR3 region having the amino acid

sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2, H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, L-CDR3 region having the amino acid sequence of SEQ ID NO: 10, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, (v) H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2, H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, L-CDR3 region having the amino acid sequence of SEQ ID NO: 11, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14, and (vi) H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, H-CDR2 region having the amino acid sequence of SEQ ID NO: 2, H-CDR1 region having the amino acid sequence of SEQ ID NO: 3, L-CDR3 region having the amino acid sequence of SEQ ID NO: 12, L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and L-CDR2 region having the amino acid sequence of SEQ ID NO: 14,

[0036]

In another aspect, the disclosure provides the following antibodies.

[0037]

One embodiment also provides an antibody comprising (i) a heavy chain having an amino acid sequence of SEQ ID NO: 49 or 51; and (ii) a light chain having an amino acid sequence selected from the group of SEQ ID NOs: 53, 55, 57, 59, 61 and 63.

[0038]

Yet another embodiment provides an antibody selected from the group of (i) a heavy chain having an amino acid sequence of SEQ ID NO: 49 and a light chain having an amino acid sequence of SEQ ID NO: 53 (named as "MOR07919"), (ii) a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 55 (named as "MOR07692"), (iii) a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 57 (named as "MOR07923"), (iv) a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 59 (named as "MOR07924"), (v) a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ

ID NO: 61 (named as "MOR07925"), (vi) a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 63 named as "MOR07926").

[0039]

In one aspect, the disclosure provides antibodies having an antigen-binding region that can bind specifically to or has a high affinity for one or more regions of target MST1R, having the amino acid sequence of SEQ ID NO: 17. An antibody is said to have a "high affinity" for an antigen if the affinity measurement is at least 100 nM (monovalent affinity of Fab fragment) as a K_D. An antibody or antigen-binding region described herein can, for example, bind to MST1R with an affinity of about less than 100 nM, less than about 60 nM, or less than about 30 nM. Further embodiments include antibodies that bind to MST1R with an affinity of less than about 10 nM or less than about 3 nM. In particular, isolated human or humanized antibodies or functional fragments thereof comprising an antigen-binding region that is specific for a partial peptide of MST1R, having an amino acid sequence of SEQ ID NO: 17, where the antibody or functional fragment thereof has an affinity against the partial peptide of MST1R as a K_D of less than about 10 nM, less than about 5 nM, less than about 1 nM, less than about 0.5 nM or less than about 0.1 nM as determined by surface plasmon resonance. While, the affinity against the partial peptide of MST1R as a K_D less than about 10 nM, less than about 5 nM, less than about 1 nM, less than about 0.5 nM or less than 0.1 nM as determined by Solution Equilibrium Titration. For instance, the affinity of an antibody, described herein, against MST1R may be about 0.98 nM or 0.02 nM (monovalent affinity of Fab fragment).

Table 1 provides a summary of affinities of representative antibodies disclosed herein, as determined by surface plasmon resonance (Biacore) and Solution Equilibrium Titration (SET):

[0040]

[Table 1]

TABLE 1: Antibody Affinities

Antibody (Fab)	BIACORE (Fab) K_D [nM]	SET (Fab) K_D [nM]
MOR07692	0.80	0.25
MOR07919	0.98	0.27
MOR07923	0.07	0.02
MOR07924	0.20	0.03
MOR07925	0.02	0.01
MOR07926	0.13	0.04

[0041]

With reference to Table 1, the affinity of MOR X antibodies was measured by surface plasmon resonance (Biacore) on immobilized recombinant human MST1R. The Biacore studies were performed on directly immobilized antigen. The Fab format of MORs X exhibit a monovalent affinity range between about 0.02 and 0.98 nM on immobilized MST1R protein with Fab MOR07925 showing the highest affinity, followed by Fabs MOR07923 and MOR07926. In addition, the Fab format of MORs X exhibit affinity range between about 0.01 and 0.27 nM with Fab MOR07925 showing the highest affinity, followed by Fabs MOR07923 and MOR07924 in SET studies.

[0042]

Another feature of antibodies described herein is their specificity for an area within the N-terminal region of MST1R. For example, MOR X disclosed herein can bind specifically to the N-terminal region of MST1R.

[0043]

The type of epitope to which an antibody as described herein binds may be linear (i.e. one consecutive stretch of amino acids) or conformational (i.e. multiple stretches of amino acids). In order to determine whether the epitope of a particular antibody is linear or conformational, the skilled practitioner can analyze the binding of antibodies to overlapping peptides (e.g., 13-mer peptides with an overlap of 11 amino acids) covering different domains of MST1R. ELISA analysis was performed using a recombinant MST1R partial peptide, having the amino acid sequence of SEQ ID NO: 17. Since MOR X was not applicable to immunoblot analysis in order to detect denatured form of the same recombinant MST1R protein, then MOR X must have conformational epitopes within amino acids residues of SEQ ID NO: 17.

[0044]

An antibody disclosed herein is species cross-reactive with humans and at least one other species, which may be, for example, a monkey or a mouse. An antibody that is cross reactive with at least cynomolgus monkey, for example, can provide greater flexibility and benefits over known anti-target MST1R antibodies, for purposes of conducting *in vivo* studies in multiple species with the same antibody.

[0045]

In one embodiment, the described antibody not only is able to bind to MST1R, but also is able to inhibit activation of the MST1R. Inhibition of the receptor leads to suppression of intrinsic kinase activity of the receptor and down-regulates signal transduction. Such down regulation can occur for example by limiting ligand binding to MST1R, changing conformation of MST1R, or internalization of MST1R. More specifically, the antibody disclosed herein can mediate its therapeutic effect by MST1R via antibody-effector functions.

[0046]

Yet another embodiment relates to the inhibition of ligand-dependent MST1R phosphorylation activity of MST1R by antibodies described herein. The disclosed antibody IC50 value of at least 100 ng/ml, at least 50 ng/ml, at least 20 ng/ml, at least 10 ng /ml or at least 5 ng/ml in MSP-dependent MST1R signal transduction assay system such as an "Elk1 luciferase assay".

[0047]

Another antibody described herein also inhibits ligand-independent MST1R activation

[0048]

A further antibody disclosed herein also inhibits ERK phosphorylation in response to MST1R ligand MSP.

[0049]

Yet another antibody described herein also suppresses MSP-promoted proliferation of tumor cells that express MST1R.

(Peptide Variants)

[0050]

Antibodies described throughout the disclosure are not limited to the specific peptide sequences provided herein. Rather, variants of these polypeptides are also embodied. With reference to the instant disclosure and conventionally available technologies and references, the skilled practitioner will be able to prepare, test and utilize functional variants of the antibodies disclosed herein, while appreciating these variants having the ability to suppress both/either ligand - dependent and/or -independent activation of MST1R fall within the scope of the present invention.

[0051]

A variant can include, for example, an antibody that has at least one altered complementarity determining region (CDR) (hyper-variable) and/or framework (FR) (variable) domain/position, vis-à-vis a peptide sequence disclosed herein. To better illustrate this concept, a brief description of antibody structure follows.

[0052]

An antibody is composed of two peptide chains, each containing one (light chain) or three (heavy chain) constant domains and a variable region (VL, VH), the latter of which is in each case made up of four FR regions and three interspaced CDRs. The antigen-binding site is formed by one or more CDRs, yet the FR regions provide the structural framework for the CDRs and, hence, play an important role in antigen binding. By altering one or more amino acid residues in a CDR or

FR region, the skilled worker routinely can generate mutated or diversified antibody sequences, which can be screened against the antigen, for new or improved properties, for example.

[0053]

FIG. 1 (VH) and FIG. 2: (VL) delineate the CDR and FR regions (according to Kabat definition) for certain antibodies disclosed herein and compare amino acids at a given position to each other and to corresponding HuCAL "master gene" sequences (as described in U.S. Patent No. 6,300,064):

[0054]

The skilled practitioner can use the data in FIG. 1 and FIG. 2 to design peptide variants that are within the scope of the embodiments disclosed herein. In one embodiment, variants are constructed by changing amino acids within one or more CDR regions; a variant might also have one or more altered framework regions. With reference to a comparison of the novel antibodies to each other, candidate residues that can be changed include residues of the variable light and residues of the variable heavy chains of MORs X. Alterations also may be made in the framework regions. For example, a peptide FR domain might be altered where there is a deviation in a residue compared to a germline sequence.

[0055]

With reference to a comparison of the novel antibodies to the corresponding consensus or "master gene" sequence, candidate residues that can be changed including residues of the variable light chain of MOR X, such as residues of VLλ3 and including residues of the variable heavy chain of MOR X, such as residues of VH3. Alternatively, the skilled worker could make the same analysis by comparing the amino acid sequences disclosed herein to known sequences of the same class of such antibodies, using, for example, the procedure described by Knappik *et al.* (*J. Mol. Biol.* 296, 57-86, 2000) and U.S. Patent No. 6,300,064 issued to Knappik *et al.*

[0056]

Furthermore, variants may be obtained by using one MOR X as a starting point for optimization by diversifying one or more amino acid residues in the MOR X sequence, preferably amino acid residues in one or more CDRs, and by screening the resulting collection of antibody variants for variants with improved properties. Diversification of one or more amino acid residues in the

CDR-3 of VL, the CDR-3 of VH, the CDR-1 of VL and/or the CDR-2 of VH may be accomplished by synthesizing a collection of DNA molecules using trinucleotide mutagenesis (TRIM) technology (Virnekäs, B., Ge, L., Plückthun, A., Schneider, K.C., Wellnhofer, G., and Moroney S.E. (1994) "Trinucleotide phosphoramidites: ideal reagents for the synthesis of mixed oligonucleotides for random mutagenesis." *Nucl. Acids Res.* 22, 5600.).

(Conservative Amino Acid Variants)

[0057]

Polypeptide variants may be made that conserve the overall molecular structure of an antibody peptide sequence described herein. Given the properties of the individual amino acids, some rational substitutions will be recognized by the skilled worker. Amino acid substitutions, *i.e.*, "conservative substitutions," may be made, for instance, on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, and/or the amphipathic nature of the residues involved.

[0058]

For example, (a) nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan, and methionine; (b) polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, asparagine, and glutamine; (c) positively charged (basic) amino acids include arginine, lysine, and histidine; and (d) negatively charged (acidic) amino acids include aspartic acid and glutamic acid. Substitutions typically may be made within groups (a)-(d). In addition, glycine and proline may be substituted for one another based on their ability to disrupt α-helices. Similarly, certain amino acids, such as alanine, cysteine, leucine, methionine, glutamic acid, glutamine, histidine and lysine are more commonly found in α-helices, while valine, isoleucine, phenylalanine, tyrosine, tryptophan and threonine are more commonly found in β-pleated sheets. Glycine, serine, aspartic acid, asparagine, and proline are commonly found in turns. Some preferred substitutions may be made among the following groups: (i) S and T; (ii) P and G; and (iii) A, V, L and I. Given the known genetic code, and recombinant and synthetic DNA techniques, the skilled practitioner readily can construct DNAs encoding the conservative amino acid variants.

[0059]

As used herein, "sequence identity" between two polypeptide sequences, indicates the percentage of amino acids that are identical between the sequences. "Sequence homology" indicates the percentage of amino acids that either is identical or that represents conservative amino acid substitutions. Polypeptide sequences of the invention have a sequence identity in the CDR regions of at least 60%, at least 70% or 80%, at least 90% or at least 95%. Embodied antibodies also have a sequence homology in the CDR regions of at least 80%, at least 90% or at least 95%.

(DNA Molecules)

[0060]

The present disclosure also relates to the DNA molecules that encode an antibody described herein. These sequences include, but are not limited to, those DNA molecules set forth in FIGs. 3A, 3B, and 4A to 4F.

[0061]

DNA molecules of the disclosure are not limited to the sequences disclosed herein, but also include variants thereof. DNA variants within the various embodiments may be described by reference to their physical properties in hybridization. The skilled practitioner will recognize that DNA can be used to identify its complement and, since DNA is double stranded, its equivalent or homolog, using nucleic acid hybridization techniques. It also will be recognized that hybridization can occur with less than 100% complementarity. However, given appropriate choice of conditions, hybridization techniques can be used to differentiate among DNA sequences based on their structural relatedness to a particular probe. For guidance regarding such conditions see, Sambrook et al., 1989 (Sambrook, J., E.F. Fritsch, and T. Maniatis (1989) *Molecular Cloning: A laboratory manual*, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, USA) and Ausubel et al., 1995 (Ausubel, F.M., R. Brent, R.E. Kingston, D.D. Moore, J.G. Sedman, J.A. Smith, & K. Struhl. eds. (1995). *Current Protocols in Molecular Biology*. New York: John Wiley and Sons).

[0062]

Structural similarity between two polynucleotide sequences can be expressed as a function of "stringency" of the conditions under which the two sequences will hybridize with one

another. As used herein, the term "stringency" refers to the extent that the conditions disfavor hybridization. Stringent conditions strongly disfavor hybridization, and only the most structurally related molecules will hybridize to one another under such conditions. Conversely, non-stringent conditions favor hybridization of molecules displaying a lesser degree of structural relatedness. Hybridization stringency, therefore, directly correlates with the structural relationships of two nucleic acid sequences. The following relationships are useful in correlating hybridization and relatedness (where T_m is the melting temperature of a nucleic acid duplex):

[0063]

a. $T_m = 69.3 + 0.41(G+C)\%$

[0064]

b. The T_m of a duplex DNA decreases by 1℃ with every increase of 1% in the number of mismatched base pairs.

[0065]

c. $(T_m)_{\mu 2} - (T_m)_{\mu 1} = 18.5 \log_{10} \mu 2/\mu 1$

where $\mu 1$ and $\mu 2$ are the ionic strengths of two solutions.

[0066]

Hybridization stringency is a function of many factors, including overall DNA concentration, ionic strength, temperature, probe size and the presence of agents which disrupt hydrogen bonding. Factors promoting hybridization include high DNA concentrations, high ionic strengths, low temperatures, longer probe size and the absence of agents that disrupt hydrogen bonding. Hybridization typically is performed in two phases: the "binding" phase and the "washing" phase.

[0067]

First, in the binding phase, the probe is bound to the target under conditions favoring hybridization. Stringency is usually controlled at this stage by altering the temperature. For high stringency, the temperature is usually between 65℃ and 70℃, unless short (< 20 nt) oligonucleotide probes are used. A representative hybridization solution comprises 6 X SSC, 0.5% SDS, 5 X Denhardt's solution and 100 μg of non-specific carrier DNA. See Ausubel *et al.*,

section 2.9, supplement 27 (1994). Of course, many different, yet functionally equivalent, buffer conditions are known. Where the degree of relatedness is lower, a lower temperature may be chosen. Low stringency binding temperatures are between about 25℃ and 40℃. Medium stringency is between at least about 40℃ to less than about 65℃. High stringency is at least about 65℃.

[0068]

Second, the excess probe is removed by washing. It is at this phase that more stringent conditions usually are applied. Hence, it is this "washing" stage that is most important in determining relatedness via hybridization. Washing solutions typically contain lower salt concentrations. One exemplary medium stringency solution contains 2 X SSC and 0.1% SDS. A high stringency wash solution contains the equivalent (in ionic strength) of less than about 0.2 X SSC, with a preferred stringent solution containing about 0.1 X SSC. The temperatures associated with various stringencies are the same as discussed above for "binding." The washing solution also typically is replaced a number of times during washing. For example, typical high stringency washing conditions comprise washing twice for 30 minutes at 55℃and three times for 15 minutes at 60℃.

[0069]

Accordingly, the present disclosure includes nucleic acid molecules that hybridize to the molecules of set forth in FIGs. 3A, 3B, and 4A to 4F under high stringency binding and washing conditions, where such nucleic molecules encode an antibody or functional fragment thereof having properties as described herein. Embodied molecules (from an mRNA perspective) are those that have at least 75% or 80% (preferably at least 85%, more preferably at least 90% and most preferably at least 95%) homology or sequence identity with one of the DNA molecules described herein. In one particular example of a variant of the disclosure, nucleic acid position 7 in SEQ ID NO: 18 or 20 can be substituted from a C to a G, thereby changing the codon from CAA to GAA.

(Functionally Equivalent Variants)

[0070]

Yet another class of DNA variants within the scope of the invention may be described with reference to the product they encode (see the peptides listed in FIGs. 3C, 3D, and 4G to 4L). These functionally equivalent genes are characterized by the fact that they encode the same peptide sequences found in FIGs. 3C, 3D, and 4G to 4L due to the degeneracy of the genetic code. The amino acid sequence in FIG. 3C is also shown as SEQ ID NO: 19. The amino acid sequence in FIG. 3D is also shown as SEQ ID: NO:21. The amino acid sequence in FIG. 4G is also shown as SEQ ID: NO:23. The amino acid sequence in FIG. 4H is also shown as SEQ ID: NO:25. The amino acid sequence in FIG. 4I is also shown as SEQ ID: NO: 27. The amino acid sequence in FIG. 4J is also shown as SEQ ID: NO: 29. The amino acid sequence in FIG. 4K is also shown as SEQ ID: NO: 31. The amino acid sequence in FIG. 4L is also shown as SEQ ID: NO: 33.

[0071]

It is recognized that variants of DNA molecules provided herein can be constructed in several different ways. For example, they may be constructed as completely synthetic DNAs. Methods of efficiently synthesizing oligonucleotides in the range of 20 to about 150 nucleotides are widely available. See Ausubel et al., section 2.11, Supplement 21 (1993). Overlapping oligonucleotides may be synthesized and assembled in a fashion first reported by Khorana et al., *J. Mol. Biol.* 72:209-217 (1971); see also Ausubel et al., supra, Section 8.2. Synthetic DNAs preferably are designed with convenient restriction sites engineered at the 5' and 3' ends of the gene to facilitate cloning into an appropriate vector.

[0072]

As indicated, a method of generating variants is to start with one of the DNAs disclosed herein and then to conduct site-directed mutagenesis. *See* Ausubel et al., *supra,* chapter 8, Supplement 37 (1997). In a typical method, a target DNA is cloned into a single-stranded DNA bacteriophage vehicle. Single-stranded DNA is isolated and hybridized with an oligonucleotide containing the desired nucleotide alteration(s). The complementary strand is synthesized and the double stranded phage is introduced into a host. Some of the resulting progeny will contain the desired mutant, which can be confirmed using DNA sequencing. In addition, various methods are available that increase the probability that the progeny phage will be the desired

mutant. These methods are well known to those in the field and kits are commercially available for generating such mutants.

(Recombinant DNA constructs and expression)

[0073]

The present disclosure further provides recombinant DNA constructs comprising one or more of the nucleotide sequences described herein. These recombinant constructs are used in connection with a vector, such as a plasmid, phagemid, phage or viral vector, into which a DNA molecule encoding any disclosed antibody is inserted.

[0074]

The encoded gene may be produced by techniques described in Sambrook et al., 1989, and Ausubel et al., 1989. Alternatively, the DNA sequences may be chemically synthesized using, for example, synthesizers. See, for example, the techniques described in *Oligonucleotide Synthesis* (1984, Gait, ed., IRL Press, Oxford), which is incorporated by reference herein in its entirety. Recombinant constructs of the disclosure are comprised with expression vectors that are capable of expressing the RNA and/or protein products of the encoded DNA(s). The vector may further comprise regulatory sequences, including a promoter operably linked to the open reading frame (ORF). The vector may further comprise a selectable marker sequence. Specific initiation and bacterial secretory signals also may be required for efficient translation of inserted target gene coding sequences.

[0075]

The present disclosure further provides host cells containing at least one of the DNAs described herein. The host cell can be virtually any cell for which expression vectors are available. It may be, for example, a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, and may be a prokaryotic cell, such as a bacterial cell. Introduction of the recombinant construct into the host cell can be effected by calcium phosphate transfection, lipofection, DEAE, dextran mediated transfection, electroporation or phage infection.

(Bacterial Expression)

[0076]

Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and, if desirable, to provide amplification within the host. Suitable prokaryotic hosts for transformation include *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*.

[0077]

Bacterial vectors may be, for example, bacteriophage-, plasmid- or phagemid-based. These vectors can contain a selectable marker and bacterial origin of replication derived from commercially available plasmids typically containing elements of the well known cloning vector pBR322 (ATCC Accession No. 37017). Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter is de-repressed/induced by appropriate means (*e.g.*, temperature shift or chemical induction) and cells are cultured for an additional period. Cells are typically harvested by centrifugation, disrupted by physical or chemical means, and the resulting crude extract retained for further purification.

[0078]

In bacterial systems, a number of expression vectors may be advantageously selected depending upon the use intended for the protein being expressed. For example, when a large quantity of such a protein is to be produced, for the generation of antibodies or to screen peptide libraries, for example, vectors which direct the expression of high levels of fusion protein products that are readily purified may be desirable.

(Therapeutic Methods)

[0079]

Therapeutic methods involve administering to a subject in need of treatment a therapeutically effective amount of an antibody contemplated by the disclosure. A "therapeutically effective"

amount hereby is defined as the amount of an antibody that is of sufficient quantity to deplete MST1R -positive cells in a treated area of a subject—either as a single dose or according to a multiple dose regimen, alone or in combination with other agents, which leads to the alleviation of an adverse condition, yet which amount is toxicologically tolerable. The subject may be a human or non-human animal (*e.g.*, rabbit, rat, mouse, monkey or other lower-order primate).

[0080]

An antibody of the disclosure might be co-administered with known medicaments, and in some instances the antibody might itself be modified. For example, an antibody could be conjugated to an immunotoxin or a radio labeled antibody to potentially further increase efficacy.

[0081]

The antibodies described herein can be used as a therapeutic or a diagnostic tool in a variety of situations where MST1R is undesirably expressed or found. Disorders and conditions particularly suitable for treatment with an antibody of the disclosure are MST1R- expressing malignant tumors and neoplasma, for example, breast, lung, colon, bladder, skin, pancreatic, glioma, lymphoma, prostate, thyroid, ovary, gastric, liver, stomach and on the like.

[0082]

To treat any of the foregoing disorders, pharmaceutical compositions for use in accordance with the present disclosure may be formulated in a conventional manner using one or more physiologically acceptable carriers or excipients. Any antibody described herein can be administered by any suitable means, which can vary, depending on the type of disorder being treated. Possible administration routes include parenteral (e.g., intramuscular, intravenous, intraarterial, intraperitoneal, or subcutaneous), intrapulmonary and intranasal, and, if desired for local immunosuppressive treatment, intralesional administration. In addition, any disclosed antibody may be administered by pulse infusion, with, e.g., declining doses of the antibody. The dosing can be administered by injections, such as for example, intravenous or subcutaneous injections, depending in part on whether the administration is brief or chronic. The amount to be administered will depend on a variety of factors such as the clinical symptoms, weight of the individual, whether other drugs are administered. The skilled practitioner will recognize that the route of administration will vary depending on the disorder or condition to be treated and will

understand which route would be most appropriate for the individual based on the specific factors for each individual.

[0083]

Determining a therapeutically effective amount of the novel polypeptide, according to this invention, largely will depend on particular patient characteristics, route of administration, and the nature of the disorder being treated. General guidance can be found, for example, in the publications of the *International Conference on Harmonisation* and in *Remington's Pharmaceutical Sciences*, chapters 27 and 28, pp. 484-528 (18th ed., Alfonso R. Gennaro, ED., Easton, PA.: Mack Pub. Co., 1990). More specifically, determining a therapeutically effective amount will depend on such factors as toxicity and efficacy of the medicament. Toxicity may be determined using methods well known in the art and found in the foregoing references. Efficacy may be determined utilizing the same guidance in conjunction with the methods described below in the Examples.

(Diagnostic Methods)

[0084]

MST1R is highly expressed on cancer cells in certain malignancies; thus, an anti-MST1R antibody of the disclosure may be employed in order to image or visualize a site or location of possible MST1R in a patient. In this regard, an antibody can be detectably labeled, through the use of radioisotopes, affinity labels (such as biotin, avidin, etc.), fluorescent labels, paramagnetic atoms, etc. Procedures for accomplishing such labeling are well known to the art. Clinical applications of antibodies in diagnostic imaging are reviewed by Grossman, H.B., *Urol. Clin. North Amer.* 13:465-474 (1986)), Unger, E.C. et al., *Invest. Radiol.* 20:693-700 (1985)), and Khaw, B. A. et al., *Science* 209:295-297 (1980)).

[0085]

The detection of foci of such detectably labeled antibodies might be indicative of MST1R, for example. In one embodiment, this examination is done by removing samples of tissue or blood and incubating such samples in the presence of the detectably labeled antibodies. In a one embodiment, this technique is done in a non-invasive manner through the use of magnetic

imaging, fluorography, etc. Such a diagnostic test may be employed in monitoring the success of treatment of diseases, where presence or absence of a MST1R-positive cell is a relevant indicator.

(Therapeutic And Diagnostic Compositions)

[0086]

The antibodies of the present disclosure can be formulated according to known methods to prepare pharmaceutically useful compositions, where an antibody described herein (including any functional fragment thereof) is combined in a mixture with a pharmaceutically acceptable carrier vehicle. Suitable vehicles and their formulation are described, for example, in *REMINGTON'S PHARMACEUTICAL SCIENCES* (18th ed., Alfonso R. Gennaro, ED., Easton, PA.: Mack Pub. Co., 1990). In order to form a pharmaceutically acceptable composition suitable for effective administration, such compositions will contain an effective amount of one or more of the antibodies of the present disclosure, together with a suitable amount of carrier vehicle.

[0087]

Preparations may be suitably formulated to give controlled-release of the active compound. Controlled-release preparations may be achieved through the use of polymers to complex or absorb anti-MST1R antibody. The controlled delivery may be exercised by selecting appropriate macromolecules (for example polyesters, polyamino acids, polyvinyl, pyrrolidone, ethylenevinyl-acetate, methylcellulose, carboxymethylcellulose, or protamine, sulfate) and the concentration of macromolecules as well as the methods of incorporation in order to control release. Another possible method to control the duration of action by controlled release preparations is to incorporate anti-MST1R antibody into particles of a polymeric material such as polyesters, polyamino acids, hydrogels, poly(lactic acid) or ethylene vinylacetate copolymers. Alternatively, instead of incorporating these agents into polymeric particles, it is possible to entrap these materials in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxymethylcellulose or gelatine-microcapsules and poly(methylmethacylate) microcapsules, respectively, or in colloidal drug delivery systems, for example, liposomes, albumin microspheres, microemulsions, nanoparticles,

and nanocapsules or in macroemulsions. Such techniques are disclosed in *Remington's Pharmaceutical Sciences* (1980).

[0088]

The compounds may be formulated for parenteral administration by injection, *e.g.*, by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, *e.g.*, in ampules, or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilizing and/or dispersing agents. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, *e.g.*, sterile pyrogen-free water, before use.

[0089]

The compositions may, if desired, be presented in a pack or dispenser device, which may contain one or more unit dosage forms containing the active ingredient. The pack may for example comprise metal or plastic foil, such as a blister pack. The pack or dispenser device may be accompanied by instructions for administration. Moreover, the pack or dispenser device and compositions may be presented in a kit for commercial distribution.

[0090]

The various embodiments of the invention may further be understood by reference to the following working examples, which are intended to illustrate and, hence, not limit the scope of the inventive disclosure.

Brief Description of Drawings

[0091]

[Fig. 1]

FIG. 1 provides amino acid sequences of various novel antibody variable heavy regions, and which delineates the CDR and framework (FR) regions. The VH3 sequence (SEQ ID NO: 80) is aligned with the MOR07692, MOR07923, MOR07924, MOR07925, MOR07926 variable heavy region sequence (SEQ ID NO: 19), and the VH5 sequence (SEQ ID NO: 81) is aligned with MOR07919 variable heavy region sequence (SEQ ID NO: 21).

[0092]

[Fig. 2]

FIG. 2: FIG. 2A and FIG. 2B provide amino acid sequences of various novel antibody variable light regions, and which delineates the CDR and framework (FR) regions. The VLκ3 sequence (VLk3; SEQ ID NO: 82) is aligned with the MOR07692 (SEQ ID NO: 23), MOR07923 (SEQ ID NO: 27), MOR07924 (SEQ ID NO: 29), MOR07925 (SEQ ID NO: 31), MOR07926 (SEQ ID NO: 33) variable light region sequences, and the VLλ3 sequence (VLl3; SEQ ID NO: 83) is aligned with MOR07919 variable light region sequence (SEQ ID NO: 25).

[0093]

[Fig. 3]

FIG. 3: FIG. 3A (MOR07692, MOR07923, MOR07924, MOR07925, MOR07926; SEQ ID NO: 18) and FIG. 3B (MOR07919; SEQ ID NO: 20) provide nucleic acid sequences of various novel antibody variable heavy regions. FIG. 3C (MOR07692, MOR07923, MOR07924, MOR07925, MOR07926; SEQ ID NO: 19) and FIG. 3D (MOR07919; SEQ ID NO: 21) provide amino acid sequences of various novel antibody variable heavy regions. CDR regions H-CDR1, H-CDR2 and H-CDR3 are designated from N- to C-terminus in boldface and underlined.

[0094]

[Fig. 4]

FIG. 4: FIG. 4A (MOR07692; SEQ ID NO: 22), FIG. 4B (MOR07919; SEQ ID NO: 24), FIG. 4C (MOR07923; SEQ ID NO: 26), FIG. 4D (MOR07924; SEQ ID NO: 28), FIG. 4E (MOR07925; SEQ ID NO: 30) and FIG. 4F (MOR07926; SEQ ID NO: 32) provide nucleic acid sequences of various novel antibody variable light regions. FIG. 4G (MOR07692; SEQ ID NO: 23), FIG. 4H (MOR07919; SEQ ID NO: 25), FIG. 4I (MOR07923; SEQ ID NO: 27), FIG. 4J (MOR07924; SEQ ID NO: 29), FIG. 4K (MOR07925; SEQ ID NO: 31), and FIG. 4L (MOR07926; SEQ ID NO: 33) provide amino acid sequences of various novel antibody variable light regions. CDR regions L-CDR1, L-CDR2 and L-CDR3 are designated from N- to C-terminus in boldface and underlined.

[0095]

[Fig. 5]

FIG. 5 provides amino acid sequences of variable heavy regions of various consensus-based Human Combinatorial Antibody Library (HuCAL®) antibody master gene sequences. CDR regions H-CDR1, H-CDR2 and H-CDR3 are underlined as designated from N- to C-terminus. The upper line is MOR07919 (SEQ ID NO: 21) while the lower line is MOR07692/7923/7924/7925/7926 (SEQ ID NO: 19).

[0096]

[Fig. 6]

FIG. 6 provides amino acid sequences of variable light regions of various consensus-based HuCAL antibody master gene sequences. CDR regions L-CDR1, L-CDR2 and L-CDR3 are underlined as designated from N- to C-terminus. The two sets of lines from top to bottom are as follows: MOR07919 (SEQ ID NO: 25); MOR07692 (SEQ ID NO: 23); MOR07923 (SEQ ID NO: 27); MOR07924 (SEQ ID NO: 29); MOR07925 (SEQ ID NO: 31); MOR07926 (SEQ ID NO: 33), respectively.

[0097]

[Fig. 7]

FIG. 7: FIG. 7A and FIG. 7B provide the nucleic acid and the amino acid sequence of various novel antibody heavy chains (FIG. 7A: MOR07919; FIG. 7B: MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926; respectively) as expressed from pMORPH®2_h_IgG1f. CDR regions are in boldface and underlined. Amino acid sequence of VH leader and heavy chain constant region is indicated in italics or italics and boldface, respectively. Restriction sites and priming sites of sequencing primers are designated above or below the sequence. The nucleic acid sequence of FIG. 7A is represented by SEQ ID NO: 64, while the amino acid sequence is SEQ ID NO: 65. The nucleic acid sequence of FIG. 7B is represented by SEQ ID NO: 66, while the amino acid sequence is SEQ ID NO: 67.

[0098]

[Fig. 8]

FIG. 8 provides the nucleic acid sequence (SEQ ID NO: 68) and the amino acid sequence (SEQ ID NO: 69) of a novel antibody lambda light chain (MOR07919) as expressed from pMORPH®2_h_Ig_lambda2. CDR regions are in boldface and underlined. Amino acid sequence of VL leader and lambda light chain constant region is indicated in italics or italics and boldface, respectively. Restriction sites and priming sites of sequencing primers are designated above or below the sequence.

[0099]

[Fig. 9]

FIG. 9: FIG. 9A to FIG. 9E provide the nucleic acid and the amino acid sequence of various novel antibody kappa light chains as expressed from pMORPH®2_h_Ig_kappa. CDR regions are in boldface and underlined. Amino acid sequence of VL leader and kappa light chain constant region is indicated in italics or italics and boldface, respectively. Restriction sites and priming sites of sequencing primers are designated above or below the sequence. The nucleic acid sequence of FIG. 9A is represented by SEQ ID NO: 70, while the amino acid sequence is SEQ ID NO: 71 (MOR07692). The nucleic acid sequence of FIG. 9B is represented by SEQ ID NO: 72, while the amino acid sequence is SEQ ID NO: 73 (MOR07923). The nucleic acid sequence of FIG. 9C is represented by SEQ ID NO: 74, while the amino acid sequence is SEQ ID NO: 75 (MOR07924). The nucleic acid sequence of FIG. 9D is represented by SEQ ID NO: 76, while the amino acid sequence is SEQ ID NO: 77 (MOR07925). The nucleic acid sequence of FIG. 9E is represented by SEQ ID NO: 78, while the amino acid sequence is SEQ ID NO: 79 (MOR07926).

[0100]

[Fig. 10]

FIG. 10 provides a FACS analysis demonstrating crossreactivity of isolated antibodies (MorphoSys IgG1- 2 μg/ml) to MST1R orthologs.

[0101]

[Fig. 11]

FIG. 11 shows binding activity of MOR07692, MOR07919, MOR07923, MOR07924, MOR07925 and MOR07926 to the 25-571 portion of human MST1R compared to PBS

control. Using the t-test analysis for n=3, the p values are as follows: MOR07692: 4.56E-07; MOR07919: 1.43E-05; MOR07923: 2.10E-05; MOR07924: 1.42E-06; MOR07925: 9.74E-07; and MOR07926: 1.53E-06.

[0102]

[Fig. 12]

FIG. 12 shows inhibitory Elk1 trans reporter activity in the absence of ligand. Using the t-test analysis and an antibody concentration of 5 μg/ml, the p values are as follows: MOR07692: 5.39E-06; MOR07919: 3.19E-04; and MOR07925: 3.78E-05.

[0103]

[Fig. 13]

FIG. 13 shows inhibition of 200 ng/ml MSP-induced phosphorylation by MOR07692 compared to hIgG control at various time points (min) at an absorbance of 450 nm with a 570 nm reference) At the 5 min time point, the p value is 5.43E-05, while at the 15 min time point, the p value is 4.76E-06.

[0104]

[Fig. 14]

FIG. 14 is a western blot illustrating inhibition of 100 ng/ml MSP-induced phosphorylation of ERK by 1 μg/ml MOR07692 compared to no antibody and hIgG controls in the presence or absence of 1 μg/ml cross-linked antibody.

[0105]

[Fig. 15]

FIG. 15 shows inhibitory activity of the specified antibodies or without antibody control on MSP-induced cell proliferation (%) in the presence or absence of 100 ng/ml MSP. For the various antibodies, the p values are as follows: MOR07692: 0.0001; MOR07919: 0.2037; MOR07923: 0.0106; MOR07924: 0.0203; MOR07925: 0.0042; and MOR07926: 0.0044.

[0106]

[Fig. 16]

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FIG. 16 shows inhibition of MSP-induced migration by indicated anti-MST1R antibodies.

[0107]

[Fig. 17]

FIG. 17 shows potential of indicted anti-MST1R antibodies to induce internalization.

[0107a]

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.

[0107b]

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.

Examples

(Cell culture and transient transfection)

[0108]

Human embryonic kidney (HEK) 293FreeStyle™ cells were grown in Freestyle 293 Medium (Invitrogen). 293α was a stable transfectant obtained by transfection with integrinαv and integrinβ3 expression vectors into HEK293 cells. HEK293 and 293α cells were propagated in DMEM containing 10% FCS. PC3 and T47D were cultured in RPMI containing 10% FCS. For pannings, screenings and functional assays, HEK 293FreeStyle™ cells were transfected with plasmid DNAs using 293fectin (Invitrogen). 293T and 293α cells were transfected with plasmid DNAs using Lipofectamine 2000 (Invitrogen) according to the supplier's instructions.

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(Flow cytometry ("FACS"))

[0109]

Cells (5 x 10^5 cells/well) were incubated with Fab or IgG antibodies at the indicated concentrations in 50 µl FACS buffer (PBS, 5% FCS) for 60 min at 4°C in round bottom 96-well culture plates (Corning). Cells were washed twice and then incubated with Fluorescein Isothiocyanate (FITC) conjugated detection antibody for 30 min at 4°C. Cells were washed again, resuspended in 0.3 ml FACS buffer and then analyzed by flow cytometry in a Cytomics FC500 (Beckman Coulter, Inc.). Data were analysed via FlowJo software (Tomy digital biology Co., Ltd.). Polyclonal goat anti-hMSP R IgG (R&D systems) or anti-FLAG M2 antibody

(Sigma) was used as a positive control and MOR03207 (anti-lysozyme) antibody was used as a negative control.

(Surface plasmon resonance)

[0110]

The kinetic constants k_{on} and k_{off} were determined with serial dilutions of the respective Fab binding to covalently immobilized MST1R-Fc fusion protein (R&D systems) using the BIAcore 3000 instrument (Biacore). For covalent antigen immobilization standard EDC-NHS amine coupling chemistry was used. For direct coupling of MST1R-Fc fusion protein CM5 senor chips (Biacore) were coated with ~600-700 RU in 10 mM acetate buffer, pH 4.5. For the reference flow cell a respective amount of HSA (human serum albumin) was used. Kinetic measurements were done in PBS (136 mM NaCl, 2.7 mM KCl, 10 mM Na_2HPO_4, 1.76 mM KH_2PO_4 pH 7.4) at a flow rate of 20 µl/min using Fab concentration range from 15.6-500 nM. Injection time for each concentration was 1 min, followed by 3 min dissociation phase. For regeneration 5 µl 10 mM HCl was used. All sensograms were globally fitted using BIA evaluation software 3.2 (Biacore).

(Solution Equilibrium Titration (SET))

[0111]

Affinity determination in solution was basically performed as described in the literature (Friguet, B., Chaffotte, A. F., Djavadi-Ohaniance, L., and Goldberg, M. E. (1985) *J Immunol Methods* 77, 305-319.). In order to improve the sensitivity and accuracy of the SET method, the method was modified from classical ELISA to ECL based technology (Haenel, C., Satzger, M., Ducata, D. D., Ostendorp, R., and Brocks, B. (2005) *Anal Biochem* 339, 182-184).

Example 1

(ANTIBODY GENERATION FROM HuCAL LIBRARIES)

[0112]

For the generation of therapeutic antibodies against MST1R, selections with the MorphoSys HuCAL GOLD phage display library were carried out. HuCAL GOLD® is a Fab library based on the HuCAL® concept (Knappik et al. (*J. Mol. Biol.*, 296, 57-86,. 2000); Krebs et al., *J. Immunol. Methods*, 254, 67-84, 2001; Rothe et al., *J. Mol. Biol.*, 376(4):1182-200, 2008), in which all six CDRs are diversified, and which employs the CysDisplay™ technology for linking Fab fragments to the phage surface (WO 01/05950).

(A. Phagemid Rescue, Phage Amplification And Purification)

[0113]

HuCAL GOLD® phagemid library was amplified in 2x YT medium containing 34 µg/ml chloramphenicol and 1% glucose (2x YT-CG). After helper phage infection (VCSM13) at an OD_{600nm} of 0.5 (30 min at 37℃without shaking; 30 min at 37℃ shaking at 250 rpm), cells were spun down (4120 g; 5 min; 4℃), resuspended in 2x YT/ 34 µg/ml chloramphenicol / 50 µg/ml kanamycin / 0.25 mM IPTG and grown overnight at 22℃. Phages were PEG-precipitated from the supernatant, resuspended in PBS/ 20% glycerol and stored at -80℃. Phage amplification between two panning rounds was conducted as follows: mid-log phase TG1 cells were infected with eluted phages and plated onto LB-agar supplemented with 1% of glucose and 34 µg/ml of chloramphenicol (LB-CG). After overnight incubation at 30℃, colonies were scraped off, and used to inoculate 2xYT-CG until an OD_{600nm} of 0.5 was reached and VCSM13 helper phages added for infection as described above.

(B. Pannings with HuCAL GOLD®)

[0114]

For the selections HuCAL GOLD® antibody-phages were divided into six pools comprising different combinations of VH master genes (pool 1: VH1/3/5 κ, pool 2: VH1/3/5 λ, pool 3: VH2/4/6 κ, pool 4: VH2/4/6 λ, pool 5: VH1-6 κ, pool 6: VH1-6 λ). These pools were individually subjected to 3 rounds of whole cell panning on MST1R expression vector-transfected HEK 293FreeStyle™ cells followed by pH-elution and a post-adsorption step on MST1R-negative HEK 293FreeStyle™ cells for depletion of irrelevant antibody-phages. Finally, the remaining antibody phages were used to infect *E. coli* TG1 cells which were then plated on

agar plates and incubated overnight at 30°C. The next day, the bacterial colonies were scraped off the plates, phages were rescued and amplified as described above. The second and the third round of selections were performed as the initial one. In addition to standard pannings, the LCDR3- RapMAT®technology was applied to potentially identify clones with higher affinities. RapMAT® represents a built-in affinity maturation process for the rapid selection of high affinity antibodies. This technology is based on the modular design of the HuCAL GOLD® Fab library. For the RapMAT® method two rounds of standard panning were performed with separate pools of lambda and kappa libraries. The selected 2nd round Fab pools were diversified via exchange of the LCDR3 with LCDR3 library cassettes. The resulting Fab libraries were subjected to two further rounds of pannings under stringent conditions.

(C. Subcloning And Expression Of Soluble Fab Fragments)

[0115]

The Fab encoding inserts of the selected HuCAL GOLD® phagemids were subcloned into the expression vector pMORPH®x9_Fab_FS (Rauchenberger et al., *J. Biol. Chem.* 278(40):38194-205, 2003) to facilitate rapid expression of soluble Fab. For this purpose, the Fab encoding insert (ompA-VLCL and phoA-Fd) of the selected clones was cut out of the plasmid DNA with *Xba*I and *Eco*RI, and cloned into the *Xba*I / *Eco*RI cut vector pMORPH®x9_ FS. Fabs expressed in this vector carry two C-terminal tags (FLAG™ and Strep-tag® II) for detection and purification.

(D. Expression of HuCAL GOLD® Fab Antibodies in *E. coli* and Purification)

[0116]

Expression of Fab fragments encoded by pMORPH®x9_Fab_FS in *E. coli* TG-1 cells was carried out in shaker flask cultures using 750 ml of 2x YT medium supplemented with 34 µg/ml chloramphenicol. Cultures were shaken at 30°C until the OD600nm reached 0.5. Expression was induced by addition of 0.75 mM IPTG for 20 hr at 30°C. Bacteria were harvested by centrifugation and the periplasmic fraction prepared using 30-35ml BBS. Fabs were purified via Strep-tag® II using Step-Tactin sepharose columns. Purity of the samples was analyzed together with calibration standards by SDS-PAGE in denatured, reduced state and by size exclusion

chromatography (SEC) in native state. Protein concentrations were determined by UV-spectrophotometry (Krebs et al., *J. Immunol. Methods* 254, 67-84, 2001).

Example 2

(CLONING, EXPRESSION AND PURIFICATION OF HuCAL® IgG1)

[0117]

In order to express full length IgG1, variable domain fragments of heavy (VH) and light chains (VL) were subcloned from Fab expression vector into pMORPH®2_hIg vectors. Restriction enzymes *Mfe*I and *Blp*I were used for subcloning of VH fragments. Restriction enzymes *Eco*RV and *Bsi*WI or *Hpa*I were used for subcloning of VL kappa or VL lambda fragments, respectively. After digestion, VH and VL fragments were isolated from preparative agarose gel and ligated into the respective IgG expression vectors (VH fragment into pMORPH®2_h_IgG1f; Vkappa fragment into pMORPH®2_h_Igκ; Vlambda fragment into pMORPH®2_h_Igλ2). The resulting IgG expression plasmids were characterized by restriction analysis and sequencing. Transient expression of full length human IgG was performed in HKB11 cells, which were transfected with IgG heavy and light chain expression vectors. IgGs were purified from cell culture supernatants by affinity chromatography *via* Protein A Sepharose column. Further down stream processing included a buffer exchange by gel filtration and sterile filtration of purified IgG. Quality control revealed a purity of >90 % by reducing SDS-PAGE and >90 % monomeric IgG as determined by analytical size exclusion chromatography.

Example 3

(ELISA SCREENING OF HuCAL® Fab CLONES AND HuCAL® IgG1)

[0118]

Wells of a 384-well MaxiSorp™ microtiter plate were coated with 0.5 µg/ml recombinant MST1R-Fc fusion protein diluted in PBS. The plate was incubated overnight at 4℃. Next day, the wells were washed 3 times with PBST (0.05% Tween20 in PBS) and then blocked with MPBST (5% milk powder in PBST) for 30 min at room temperature on a microtiter plate shaker. The wells were washed 3 times with PBST before adding the primary antibody,

i.e. preblocked BEL extracts of HuCAL® Fab clones or purified HuCAL® antibodies and control antibodies. The plate was incubated for 2 hr at room temperature on a microtiter plate shaker and then washed 3 times with PBST. For detection of HuCAL® antibodies, goat anti-human IgG alkaline phosphatase (Dianova, diluted 1:5,000 in 0.5% milkpowder in PBST) was added and the plate incubated for 1h at room temperature on a microtiter plate shaker. Subsequently, the plate was washed 5 times with TBST (0.05% Tween20 in TBS). Attophos (AttoPhos Substrate Set, Roche) was added (diluted 1:10 in TBS) and fluorescence was measured in a TECAN microtiter plate reader (emission: 535nm, excitation: 430nm).

Example 4

(CROSS-REACTIVITY ANALYSIS BY FACS)

[0119]

FACS-analysis of MST1R ortholog-expressing cells: Human MST1R (cDNA nucleotide sequence is shown as GenBank Accession No: NM_002447.2), cynomolgus monkey MST1R and mouse MST1R (cDNA nucleotide sequence is shown as GeneBank Accession No: NM_009074.1) expression vector containing N-terminal Flag tag (pFLAG-myc-CMV-19, Sigma) were constructed. cDNA encoding cynomolgus monkey MST1R was amplified by PCR using cynomolgus monkey stomach cDNA as a template with forward and reverse primers having nucleotide sequences of SEQ ID NO: 34 and 35, respectively. By sequencing analysis of PCR product, the cynomolgus MST1R ORF nucleotide sequence was identified as shown in SEQ ID NO: 36. The corresponding amino acid sequence was shown in SEQ ID NO: 37. Then human, cynomolgus monkey and mouse MST1R ORF cDNA, excluding signal peptide regions, were amplified using respective forward and reverse primers having nucleotide sequences of SEQ ID NO: 38 and 39 (human), 40 and 41 (cynomolgus monkey), and 42 and 43 (mouse) with appropriate cloning sites and then cloned into pFLAG-myc-CMV-19. An amplified human MST1R fragment encodes amino acids corresponding to GenBank Accession No: NP_002438.2 (SEQ ID NO: 45). A Mouse MST1R fragment encodes amino acids corresponding to GenBank Accession No: NP_033100.1 (SEQ ID NO: 47) except for those amino acid differences at positions: 688 (Leu to Pro), 713 (Ile to Val), 714 (Ala to Gly) and 719 (Ala to Val). These expression vectors were transfected into HEK293T cells. For FACS-analysis, cells were

incubated with 2 μg/ml primary antibodies followed by incubation with FITC-labeled secondary antibody as described above. In FIG. 10, anti-Flag antibody confirmed expression of each (human, cynomolgus monkey and mouse MST1R) protein. MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 showed binding to both human and monkey MST1R. On the other hand, MOR07919 also exhibited binding to mouse MST1R besides human and monkey MST1R.

[0120]

The nucleotide sequence of these antibodies was decided by DNA sequencer. The nucleotide sequence of variable heavy chain of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 was decided as shown in FIG. 3A and SEQ ID NO: 18. The nucleotide sequence of variable heavy chain of MOR07919 is shown in FIG. 3B and SEQ ID NO: 20. The amino acid sequence of variable heavy chain of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 was decided as shown in FIG. 3C and SEQ ID NO: 19. The amino acid sequence of variable heavy chain of MOR07919 is shown in FIG. 3D and SEQ ID NO: 21.

[0121]

The nucleotide sequence of variable light chain of MOR07692 is shown in FIG. 4A and SEQ ID NO: 22. The amino acid sequence of variable light chain of MOR07692 is shown in FIG 4G and SEQ ID NO: 23. The nucleotide sequence of variable light chain of MOR07919 is shown in FIG. 4B and SEQ ID NO: 24. The amino acid sequence of variable light chain of MOR07919 is shown in FIG. 4H and SEQ ID NO: 25. The nucleotide sequence of variable light chain of MOR07923 is shown in FIG. 4C and SEQ ID NO: 26. The amino acid sequence of variable light chain of MOR07923 is shown in FIG. 4I and SEQ ID NO: 27. The nucleotide sequence of variable light chain of MOR07924 is shown in FIG. 4D and SEQ ID NO: 28. The amino acid sequence of variable light chain of MOR07924 is shown in FIG. 4J and SEQ ID NO: 29. The nucleotide sequence of variable light chain of MOR07925 is shown in FIG. 4E and SEQ ID NO: 30. The amino acid sequence of variable light chain of MOR07925 is shown in FIG. 4K and SEQ ID NO: 31. The nucleotide sequence of variable light chain of MOR07926 is shown in FIG. 4F and SEQ ID NO: 32. The amino acid sequence of variable light chain of MOR07926 is shown in FIG. 4L and SEQ ID NO: 33.

[0122]

The amino acid sequence of variable heavy chain CDR3 (H-CDR3) of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 is shown in SEQ ID NO: 1. The amino acid sequence of variable heavy chain CDR3 (H-CDR3) of MOR07919 is shown in SEQ ID NO: 4.

[0123]

The amino acid sequence of variable heavy chain CDR2 (H-CDR2) of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 is shown in SEQ ID NO: 2. The amino acid sequence of variable heavy chain CDR2 (H-CDR2) of MOR07919 is shown in SEQ ID NO: 5.

[0124]

The amino acid sequence of variable heavy chain CDR1 (H-CDR1) of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 is shown in SEQ ID NO: 3. The amino acid sequence of variable heavy chain CDR1 (H-CDR1) of MOR07919 is shown in SEQ ID NO: 6.

[0125]

The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07692, is shown in SEQ ID NO: 7. The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07919 is shown in SEQ ID NO: 8. The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07923 is shown in SEQ ID NO: 9. The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07924 is shown in SEQ ID NO: 10. The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07925 is shown in SEQ ID NO: 11 The amino acid sequence of variable light chain CDR3 (L-CDR3) of MOR07926 is shown in SEQ ID NO: 12.

[0126]

The amino acid sequence of variable light chain CDR2 (L-CDR2) of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 is shown in SEQ ID NO: 14. The amino acid sequence of variable light chain CDR2 (L-CDR2) of MOR07919 is shown in SEQ ID NO: 16.

[0127]

The amino acid sequence of variable light chain CDR1 (L-CDR1) of MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 is shown in SEQ ID NO: 13. The amino acid sequence of variable light chain CDR1 (L-CDR1) of MOR07919 is shown in SEQ ID NO: 15.

[0128]

The nucleotide sequence of heavy chain of MOR07692 is shown in SEQ ID: NO: 50. The amino acid sequence of heavy chain of MOR07692 is shown in SEQ ID NO: 51. The nucleotide sequence of light chain of MOR07692 is shown in SEQ ID: NO: 54. The amino acid sequence of light chain of MOR07692 is shown in SEQ ID NO: 55.

[0129]

The nucleotide sequence of heavy chain of MOR07923 is shown in SEQ ID: NO: 50. The amino acid sequence of heavy chain of MOR07923 is shown in SEQ ID NO: 51. The nucleotide sequence of light chain of MOR07923 is shown in SEQ ID: NO: 56. The amino acid sequence of light chain of MOR07923 is shown in SEQ ID NO: 57.

[0130]

The nucleotide sequence of heavy chain of MOR07924 is shown in SEQ ID: NO: 50. The amino acid sequence of heavy chain of MOR07924 is shown in SEQ ID NO: 51. The nucleotide sequence of light chain of MOR07924 is shown in SEQ ID: NO: 58. The amino acid sequence of light chain of MOR07924 is shown in SEQ ID NO: 59.

[0131]

The nucleotide sequence of heavy chain of MOR07925 is shown in SEQ ID: NO: 50. The amino acid sequence of heavy chain of MOR07925 is shown in SEQ ID NO: 51. The nucleotide sequence of light chain of MOR07925 is shown in SEQ ID: NO: 60. The amino acid sequence of light chain of MOR07925 is shown in SEQ ID NO: 61.

[0132]

The nucleotide sequence of heavy chain of MOR07926 is shown in SEQ ID: NO: 50. The amino acid sequence of heavy chain of MOR07926 is shown in SEQ ID NO: 51. The nucleotide sequence of light chain of MOR07926 is shown in SEQ ID: NO: 62. The amino acid sequence of light chain of MOR07926 is shown in SEQ ID NO: 63

[0133]

The nucleotide sequence of heavy chain of MOR07919 is shown in SEQ ID: NO: 48. The amino acid sequence of heavy chain of MOR07919 is shown in SEQ ID NO: 49. The nucleotide

sequence of light chain of MOR07919 is shown in SEQ ID: NO: 52. The amino acid sequence of light chain of MOR07919 is shown in SEQ ID NO: 53.

Example 5

(BINDING ACTIVITY ANALYSIS BY ELISA)

[0134]

Wells of a 96-well MaxiSorp™ microtiter plate were coated with 1 μg/ml recombinant MST1R-Fc fusion protein (containing 25-571 amino acid sequence of human MST1R, R&D) diluted in PBS. The plate was incubated overnight at 4℃. Next day, the wells were washed once with PBS-FCS buffer (5% FCS in PBS) and then blocked with PBS-FCS buffer for 1 hr at room temperature. After removal of the PBS-FCS buffer 4 μg/ml primary antibody was added to the MST1R-Fc coated wells and incubated for 1 hr at room temperature. After washing once with PBS-FCS buffer, the secondary antibody was added and allowed to incubate for 1hr at room temperature. After washing 3 times with PBS-FCS buffer, substrate of HRP (0.4 mg/ml *o*-Phenylenediamine Dihydrochloride and 0.006% Hydrogen peroxide in substrate buffer (50 mM tri-sodium citrate dehydrate, 100mM di- sodium Hydorogen Phosphate, pH4.5)) was added. After yellow color developed, 1 M HCl was further added to stop reaction. Absorbance at 490 nm was measured in EnVision microtiter plate reader. In FIG. 11, all of the obtained antibodies (MOR07692, MOR07919, MOR07923, MOR07924, MOR07925 and MOR07926) showed binding to 25-571 portion of human MST1R. Each antibody was applicable for immunoprecipitation of non-reduced and non-denatured MST1R, but not for Western blotting to detect reduced and denatured MST1R (data not shown). It indicates that these antibodies recognize native conformation within amino acids residues in SEQ ID NO: 17.

Example 6

(BIOLOGICAL ASSAYS)

(A. Elk1 Luciferase Reporter Gene Assay)

[0135]

Functionality of antibodies was tested via Elk1 luciferase reporter gene assay. The principle of the assay is based on the co-transfection of 293α cells with several vectors. MST1R is integrated into the cell membrane and becomes activated (phosphorylated) to transduce signal to ERK (extracellullar signal-regulated kinase) when it is overexpressed or stimulated with MSP. To test functionality of antibodies, Elk1 luciferase reporter gene assay was established as follows: First we constructed pFR-Luc2CP vector. To construct pFR-Luc2CP, pFR-Luc vector (Stratagene) was digested with *Hind*III, treated with T4 DNA polymerase for blunting, and digested with *Bam*HI to obtain about 140 bp fragment containing the 5 x GAL4 binding element and TATA box. pGL4.12[*luc2CP*] (Promega) was digested with *Eco*ICRI/*Bgl*II, dephosphorylated, and ligated with the above fragment to generate pFR-Luc2CP. Then, 293α cells were transiently co-transfectd with pcDNA-DEST40 MST1R, pcDNA-DEST40, pFA2-Elk1(Stratagene), pFR-Luc2CP and pGL4.74 [hRluc/TK] (Promega) using a Lipofectamine 2000 (Invitrogen) transfection procedure and seeded onto white 96-well cell culture plates. The next day after transfection, the cells were preincubated with the antibodies for 1 hr and then the ligand (human MSP) was added to the wells. After 6 hr incubation, cell lysates were prepared and the firefly luciferase activity (specific signal) and the Renilla luciferase activity (signal for normalization) were measured using the Dual-luciferase reporter assay system (Promega). The firefly/Renilla ratio was calculated to normalize the data of each well. Table 2 shows IC_{50} values in the presence of 100 ng/ml MSP ligand. MOR07692, MOR07919, MOR07923, MOR07924, MOR07925 and MOR07926 showed low IC_{50} value ranging between 4 and 100 ng/ml. As shown in FIG. 12, overexpression of MST1R by itself induced ligand-independent activation of MST1R. MOR07925, MOR07919 and MOR07692 also suppressed this type of activation of MST1R.

[0136]

[Table 2]

Table 2: IC_{50} value of Elk1 luciferase reporter gene assay

Clone ID	Reporter assay IC50 (ng/ml)
MOR07692	4.4
MOR07919	87.6
MOR07923	9
MOR07924	15.7
MOR07925	5.9
MOR07926	11.4

(B. ELISA for Detection of Phosphorylation of MST1R)

[0137]

The change in phosphorylation status of MST1R after treatment with ligand and/or antibody was determined by ELISA system. After overnight incubation of PC3 cells (1×10^6) on 6 cm-diameter dishes, cells were washed with PBS, and incubated with 0.1% BSA-RPMI medium. After overnight incubation, cells were treated with 1 μg/ml MOR07692 antibody for 1hr at 37℃, and then stimulated with 200 ng/ml of recombinant MSP (R&D systems) for 0 min to 15 min. Then cell lysates were prepared and phosphorylated forms of MST1R were measured by Human Phospho-MSP R/Ron ELISA system (R&D systems) according to the supplier's instruction. MOR07692 showed complete inhibition of MST1R phosphorylation promoted by addition of MSP ligand as shown in FIG. 13.

(C. Western blotting for activated ERK)

[0138]

The change in phosphorylation status of ERK after treatment with ligand and/or antibody was determined by Western blotting. After overnight cultivation of PC3 cells (2×10^5) on 12 wells plate, cells were washed with PBS, and incubated with 0.1% BSA-RPMI medium. After overnight incubation, cells were treated with 1 μg/ml MOR07692 antibody with or without 1 μg/ml goat affinity purified antibody to human IgG-Fc (Cappel) for 1hr at 37℃. After the incubation, 100 ng/ml of recombinant MSP (R&D systems) was added, and further incubated for 30 min. Then cells were lysed with RIPA buffer containing complete mini (Roche) and phosphatase inhibitor (Nakarai tesque). Lysates were cleared from cellular debris by centrifugation, and protein concentrations were determined using BCA protein assay (PIERCE). Lysates were resuspended in buffer containing β-mercaproethanol and denatured at 99℃ for 5 minutes. Protein (10 μg/lane) was resolved by SDS-PAGE on 5-20% gels. Proteins were blotted onto PVDF membrane (BioRad). Membranes were blocked with Blockace (Yukijirushi), for 1 hr at room temperature and incubated overnight at 4℃ with polyclonal antibodies against ERK or phospho-ERK antibody. After washing, membranes were incubated with secondary anti-rabbit horseradish peroxidase-conjugated antibody (Amersham). Immunoreactive bands were visualized on X-ray films using ECL plus substrate (GE Healthcare). FIG. 14 represented ERK phosphorylation in response to ligand MSP. The increase was almost completely inhibited by the addition of MOR07692 in the presence and absence of cross link antibody to human IgG-Fc.

(D. Cell Proliferation assay)

[0139]

T-47D cells (5000 cells/well) suspended in RPMI medium containing 2% charcoal/dextran-treated FCS (Hyclone) were seeded onto 96-well plates. Cells were incubated with 1 μg/ml antibodies for 1 hr at 37℃, and then stimulated with 100 ng/ml recombinant MSP. After 5 days incubation, cellular ATP was measured by CellTiter-Glo luminescent cell viability assay kit (Promega), according to the supplier's instruction. As shown in FIG. 15, MOR07692,

MOR07923, MOR07924, MOR07925 and MOR07926 clearly suppressed MSP-promoted proliferation of T-47D cells. MOR07919 had a weaker inhibitory activity than other antibodies.

(E. Migration Assay)

[0140]

BxPC-3 cells (5 x 10^4 cells/well) suspended in RPMI medium containing 10% FCS were seeded onto 96-well Oris™ Cell Migration Assay plates (Platypus Technologies, LLC.). After overnight cultivation, the stoppers were removed from test wells and medium was replaced with 2% charcoal/dextran-treated FCS (Hyclone). Cells were incubated with 10 μg/ml antibodies for 1 hr at 37℃, and then stimulated with 300 ng/ml recombinant MSP. After 24 hr incubation, migrated cells were observed using bright field microscopy (Nikon) and then their images were analyzed by Image J software to calculate cell-free area. As shown in FIG 16, MOR07919, MOR07692 and MOR07925 clearly suppressed MSP-promoted cell migration of BxPC-3 cells. MOR07692 and MOR07925 had stronger inhibitory activity compared to MOR07919.

(F. Internalization assay)

[0141]

In order to evaluate the ability of antibodies to internalize, Hum-ZAP secondary conjugate (affinity-purified goat anti-human IgG-saporin provided by ADVANCED TARGETING SYSTEMS) was used as the secondary antibody to cause protein synthesis inhibition and, ultimately, cell death after internalization into cells. PC3 cells (2000 cells/well) suspended in RPMI medium containing 10% FCS were seeded onto 96-well flat clear bottom white culture plates. The next day, the cells were preincubated with antibodies for 1 hr at 4℃. After removal of the medium containing the antibodies, 0.5 μg/ml Hum-ZAP secondary conjugate was added to the wells. The plates were incubated for 1 hr at 4℃ and then for 3 days at 37℃. The cellular ATP was measured as readout for cell viability by CellTiter-Glo luminescent cell viability assay kit (Promega), according to the supplier's instruction. As shown in FIG. 17, viability of PC3 cells was greatly reduced by the treatment with MOR07692, MOR07919, MOR07923, MOR07924, MOR07925 and MOR07926, suggesting the potential of these antibodies to internalize.

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[0142]

The contents of all patents, patent applications, published PCT applications and articles, books, references, reference manuals and abstracts cited herein are hereby incorporated by reference in their entirety to more fully describe the state of the art to which the invention pertains.

[0143]

As various changes can be made in the above-described subject matter without departing from the scope and spirit of the present invention, it is intended that all subject matter contained in the above description, or defined in the appended claims, be interpreted as descriptive and illustrative of the present invention. Many modifications and variations of the present invention are possible in light of the above teachings.

[0144]

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.

[0145]

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.

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THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. An isolated human or humanized antibody or functional fragment thereof comprising an antigen-binding region that is specific for a partial peptide of MST1R, having an amino acid sequence of SEQ ID NO: 17, wherein one of the following requirements applies:
   a. said antibody or functional fragment thereof inhibits ligand -dependent and/or independent phosphorylation of MST1R.;
   b. said antibody or functional fragment thereof has an affinity against said partial peptide of MST1R as a KD of less than about 10 nM, less than about 5 nM, less than about 1 nM, less than about 0.5 nM or less than about 0.1 nM as determined by surface plasmon resonance, or by Solution Equilibrium Titration.
   c. said antibody or functional fragment thereof suppresses MSP-promoted cell proliferation or migration of tumor cells that express MST1R.;
   d. said antibody or functional fragment thereof internalizes MST1R.; or
   e. said antibody or functional fragment thereof is cross-reactive with human and at least one other species. and

   wherein said antigen-binding region comprises

   an H-CDR3 region having:
      (i) the amino acid sequence of SEQ ID NO: 1 or 4, or
      (ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 1 or 4;

   an H-CDR2 region having:
      (i) the amino acid sequence of SEQ ID NO: 2 or 5, or
      (ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 2 or 5;

   an H-CDR1 region having:
      (i) the amino acid sequence of SEQ ID NO: 3 or 6, or
      (ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 3 or 6;

   an L-CDR3 region having:
      (i) the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12, or

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(ii) an amino acid sequence having at least 80% sequence identity with the amino acid sequence of SEQ ID NO: 7, 8, 9, 10, 11 or 12;

an L-CDR1 region having:

(i) the amino acid sequence of SEQ ID NO: 13 or 15, or

(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 13 or 15; and

an L-CDR2 region having:

(i) the amino acid sequence of SEQ ID NO: 14 or 16, or

(ii) an amino acid sequence having at least 80% sequence identity with SEQ ID NO: 14 or 16.

2. The isolated human or humanized antibody or functional fragment thereof according to claim 1, comprising:

(i) the H-CDR3 region having the amino acid sequence of SEQ ID NO: 1, the H-CDR2 region having the amino acid sequence of SEQ ID NO: 2 and the H-CDR1 region having the amino acid sequence of SEQ ID NO: 3; or

(ii) the H-CDR3 region having the amino acid sequence of SEQ ID NO: 4, the H-CDR2 region having the amino acid sequence of SEQ ID NO: 5 and the H-CDR1 region having the amino acid sequence of SEQ ID NO: 6.

3. The isolated human or humanized antibody or functional fragment thereof according to claim 1, comprising:

(i) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 7, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 14;

(ii) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 8, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 15 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 16;

(iii) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 9, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 14;

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(iv) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 10, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 14;

(v) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 11, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 14; or

(vi) the L-CDR3 region having the amino acid sequence of SEQ ID NO: 12, the L-CDR1 region having the amino acid sequence of SEQ ID NO: 13 and the L-CDR2 region having the amino acid sequence of SEQ ID NO: 14.

4. The isolated human or humanized antibody or functional fragment thereof according to claim 1, wherein said antibody inhibits phosphorylation of ERK and/or Akt.

5. The isolated human or humanized antibody or functional fragment thereof according to claim 1 part (e), wherein said other species is a mouse or a monkey.

6. An isolated antigen binding region of an antibody or functional fragment thereof according to any one of claims 1 to 5.

7. The isolated antigen binding region according to claim 6, comprising:

a. a variable heavy chain having the amino acid sequence of SEQ ID NO: 19 or 21; or

b. a variable light chain having the amino acid sequence selected from the group consisting of SEQ ID NOs: 23, 25, 27, 29, 31 and 33; or

c. a heavy chain amino acid sequence of (a) SEQ ID NO: 49 or 51; or (b) a sequence having at least 80 percent sequence identity in the CDR regions with the CDR regions of SEQ ID NO: 49 or 51; or

d. a light chain amino acid sequence of (a) SEQ ID NO: 53, 55, 57, 59, 61 or 63; or (b) a sequence having at least 80 percent sequence identity in the CDR regions with the CDR regions of SEQ ID NO: 53, 55, 57, 59, 61 or 63.

8. The isolated human or humanized antibody according to any one of claims 1 to 5,

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wherein the isolated human or humanized antibody is an IgG.

9. The isolated human or humanized antibody according to claim8, wherein the isolated human or humanized antibody is an IgG1.

10. The isolated antibody or functional fragment thereof according to any one of claims 1 to 5, wherein the isolated antibody or functional fragment thereof comprises:
    a. a variable heavy chain of SEQ ID NO: 19 or 21;
    b. a variable light chain having the amino acid sequence of SEQ ID NO: 23, 25, 27, 29, 31 or 33;
    c. a heavy chain amino acid sequence selected from the group consisting of (a) SEQ ID NO: 49 and 51; and (b) a sequence having at least 80 percent sequence identity in the CDR regions with the CDR regions of SEQ ID NOs: 49 and 51; or
    d. a light chain amino acid sequence selected from the group consisting of (a) SEQ ID NOs: 53, 55, 57, 59, 61 and 63; and (b) a sequence having at least 80 percent sequence identity in the CDR regions with the CDR regions of SEQ ID NOs: 53, 55, 57, 59, 61 and 63.

11. An isolated antibody having
    a. a heavy chain having an amino acid sequence of SEQ ID NO: 49 and a light chain having an amino acid sequence of SEQ ID NO: 53; or
    b. a heavy chain having an amino acid sequence of SEQ ID NO: 51 and a light chain having an amino acid sequence of SEQ ID NO: 55, 57, 59, 61 or 63.

12. The isolated functional fragment according to any one of claims 1 to 5 or 10, which is a Fab or scFv antibody fragment.

13. A variable heavy chain of an isolated antibody or functional fragment thereof that is encoded by
    (i) a nucleic acid sequence comprising SEQ ID NO: 18 or 20, or
    (ii) a nucleic acid sequence that hybridizes under high stringency conditions to the

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complementary strand of SEQ ID NO: 18 or 20, wherein said antibody or functional fragment thereof is specific for an epitope of MST1R.

14. A variable light chain of an isolated antibody or functional fragment thereof that is encoded by
    (i) a nucleic acid sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32, or
    (ii) a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of a sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32, wherein said antibody or functional fragment thereof is specific for an epitope of MST1R.

15. An isolated nucleic acid sequence that encodes the isolated human or humanized antibody, functional fragment thereof, or isolated antigen-binding region thereof according to any one of claims 1 to 10.

16. An isolated nucleic acid sequence encoding a variable heavy chain of an isolated antibody or functional fragment thereof, comprising:
    (i) a sequence of SEQ ID NO: 18 or 20 or
    (ii) a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of SEQ ID NO: 18 or 20, wherein said antibody or functional fragment thereof is specific for a partial peptide of MST1R, having an amino acid sequence of SEQ ID NO: 17.

17. An isolated nucleic acid sequence encoding a variable light chain of an isolated antibody or functional fragment thereof, comprising
    (i) a sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32 or
    (ii) a nucleic acid sequence that hybridizes under high stringency conditions to the complementary strand of a sequence selected from the group consisting of SEQ ID NOs: 22, 24, 26, 28, 30 and 32, wherein said antibody or functional fragment thereof is specific for a partial peptide of MST1R, having an amino

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acid sequence of SEQ ID NO: 17.

18. A vector comprising the nucleic acid sequence according to claims 15 to 17.

19. An isolated cell comprising the vector according to claim 18.

20. The isolated cell according to claim 19, wherein said cell is bacterial or mammalian.

21. A method for producing an antibody or functional fragment thereof by using the vector of claim 20.

22. A method for producing an antibody or functional fragment thereof by culturing the cell of claim 19 or claim 20.

23. An antibody or functional fragment thereof which is obtainable by a method according to claims 21 or 22.

24. The human antibody according to any one of claims 1 to 11 or 23, wherein the human antibody is a synthetic human antibody.

25. A pharmaceutical composition comprising an antibody or functional fragment thereof according to any one of claims 1 to 12 or 23 to 24 and a pharmaceutically acceptable carrier or excipient thereof.

26. A method for treating a disorder or condition associated with the undesired presence of MST1R, comprising administering to a subject in need thereof an effective amount of the pharmaceutical composition according to claim 25.

27. Use of a pharmaceutical composition according to claim 25 for the manufacture of a medicament for treating a disorder or condition associated with the undesired presence of MST1R.

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28. The method according to claim 26 or the use according to claim 27, wherein said disorder or condition is caused by MST1R phosphorylation.

29. The method or use according to claim 28, wherein said disorder or condition is a malignant tumor and/or neoplasm.

30. The method or use according to claim 29, wherein said disorder or condition is: breast cancer, lung cancer, colon cancer, bladder cancer, skin cancer, pancreatic cancer, glioma, lymphoma, prostate cancer, thyroid cancer, ovary cancer, gastric cancer, liver cancer or stomach cancer.

31. A method for targeting MST1R+ cells in a subject or a cell sample, comprising the step of contacting said MST1R+ cells and the antibody or functional fragment thereof according to any one of claims 1 to 12 or 23 to 24.

32. Use of the antibody or functional fragment thereof according to any one of claims 1 to 12 or 23 to 24 for the manufacture of a medicament for targeting MST1R+ cells in a subject, comprising the step of contacting said MST1R+ cells and said medicament.

33. An isolated antibody or functional fragment thereof according to any one of claims 1 to 5, or 10, or an isolated antigen binding region according to claim 6 or 7, or an isolated antibody according to any one of claims 8, 9 or 11, or an isolated functional fragment according to claim 12, or a variable chain according to claim 13 or claim 14, or a nucleic acid sequence according to claim 15 to 17, or a vector according to claim 18, or a cell according to claim 19 or claim 20, or a composition according to claim 25, or a method according to any one of claims 21 to 22, 26, 28 to 31, or use according to any one of claims 27 to 30 or 32, or an antibody according to claim 23 or 24 substantially as hereinbefore defined.

FIG. 1

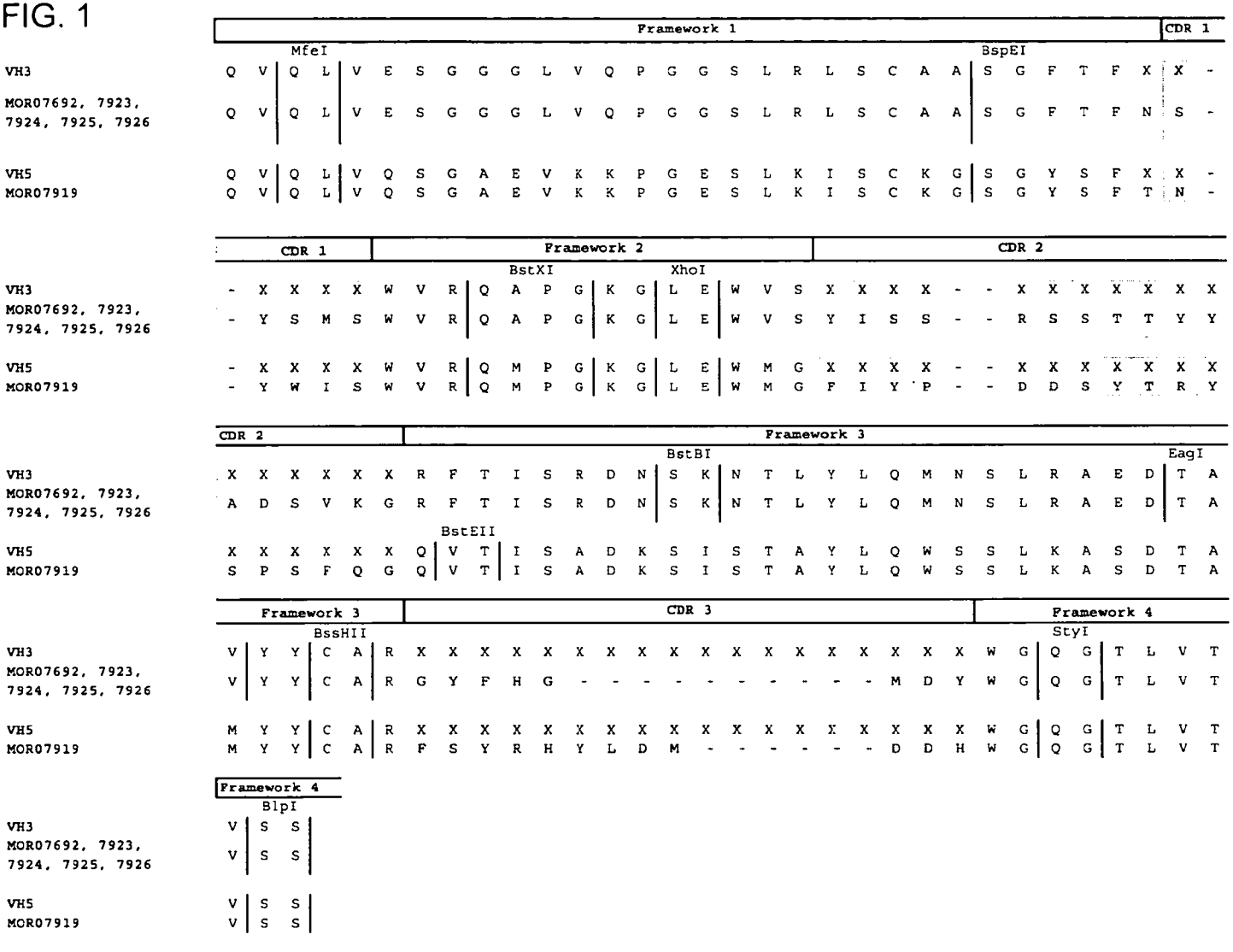
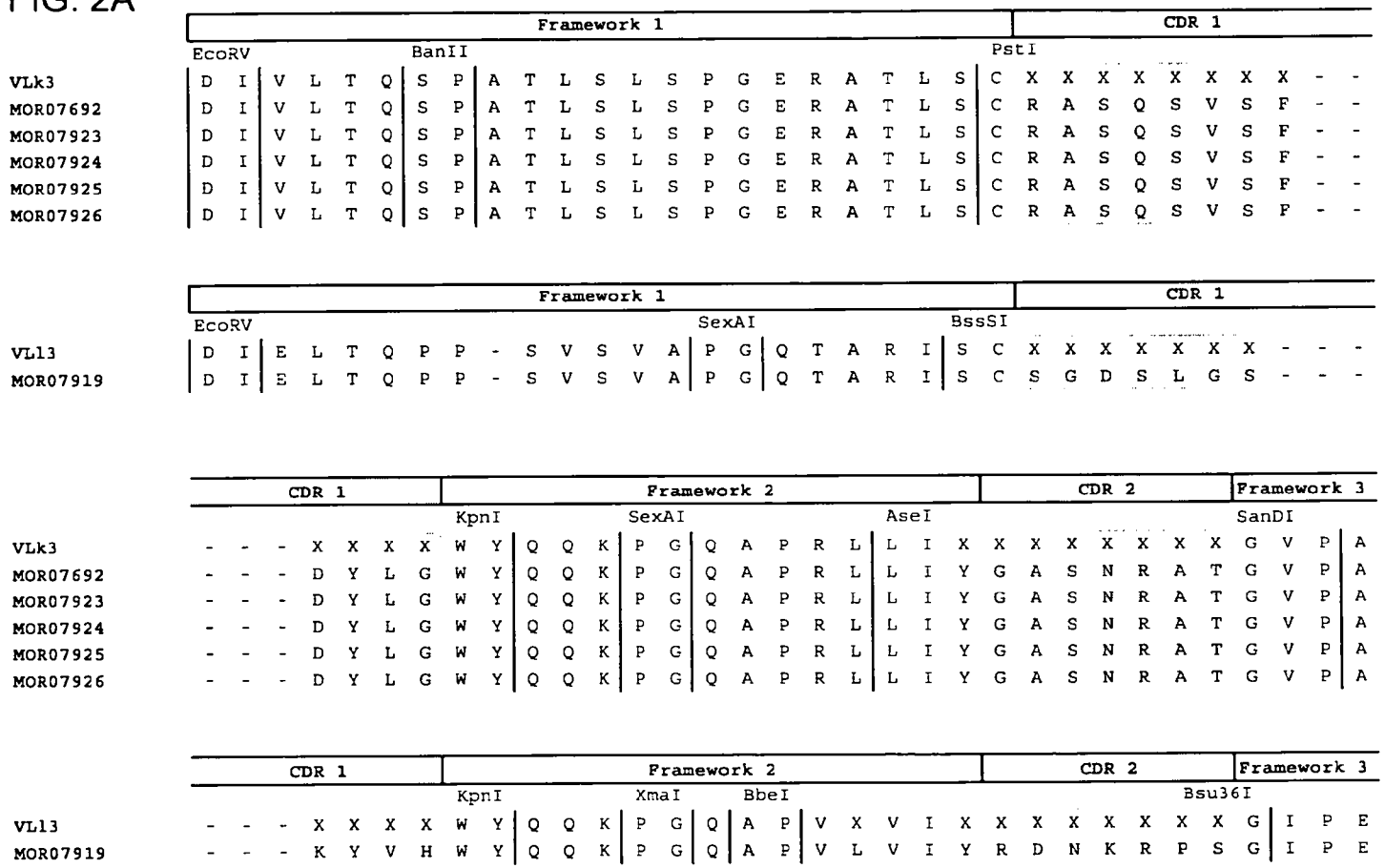

Framework 1
CDR 1
MfeI
BspEI
VH3
MOR07692, 7923, 7924, 7925, 7926
VH5
MOR07919
CDR 1
Framework 2
CDR 2
BstXI
XhoI
CDR 2
Framework 3
BstBI
EagI
BstEII
Framework 3
CDR 3
Framework 4
BssHII
StyI
Framework 4
BlpI

FIG. 2A


Framework 1
CDR 1
EcoRV
BanII
PstI
VLk3 D I V L T Q S P A T L S L S P G E R A T L S C X X X X X X X X - -
MOR07692 D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F - -
MOR07923 D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F - -
MOR07924 D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F - -
MOR07925 D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F - -
MOR07926 D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F - -
Framework 1
CDR 1
EcoRV
SexAI
BsssSI
VL13 D I E L T Q P P - S V S V A P G Q T A R I S C X X X X X X X - - -
MOR07919 D I E L T Q P P - S V S V A P G Q T A R I S C S G D S L G S - - -
CDR 1
Framework 2
CDR 2
Framework 3
KpnI
SexAI
AseI
SanDI
VLk3 - - - X X X X W Y Q Q K P G Q A P R L L I X X X X X X X G V P A
MOR07692 - - - D Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A
MOR07923 - - - D Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A
MOR07924 - - - D Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A
MOR07925 - - - D Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A
MOR07926 - - - D Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A
CDR 1
Framework 2
CDR 2
Framework 3
KpnI
XmaI
BbeI
Bsu36I
VL13 - - - X X X X W Y Q Q K P G Q A P V X V I X X X X X X X G I P E
MOR07919 - - - K Y V H W Y Q Q K P G Q A P V L V I Y R D N K R P S G I P E

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FIG. 2B

	Framework 3																						CDR 3				
					BamHI														BbsI								
VLk3	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T/V	Y	Y	C	X	X	X	X	X
MOR07692	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T	Y	Y	C	Q	Q	Y	Y	N
MOR07923	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T	Y	Y	C	F	Q	Y	L	I
MOR07924	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T	Y	Y	C	Q	Q	Y	N	I
MOR07925	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T	Y	Y	C	L	Q	Y	F	N
MOR07926	R	F	S	G	S	G	S	G	T	D	F	T	L	T	I	S	S	L	E	P	E	D	F	A	T	Y	Y	C	F	Q	A	L	I

	Framework 3																							CDR 3				
				BamHI															BbsI									
VL13	R	F	S	G	S	N	S	G	N	T	A	T	L	T	I	S	G	T	Q	A	E	D	E	A	D	Y	Y	C	X	X	X	X	X
MOR07919	R	F	S	G	S	N	S	G	N	T	A	T	L	T	I	S	G	T	Q	A	E	D	E	A	D	Y	Y	C	Q	S	Y	D	A

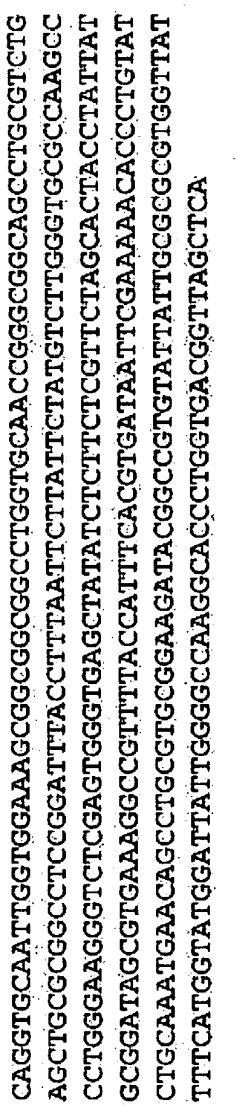
	CDR 3						Framework 4									
							MscI								BsiWI	
VLk3	X	X	-	-	X	X	F	G	Q	G	T	K	V	E	I	K	R	T
MOR07692	M	P	-	-	Y	T	F	G	Q	G	T	K	V	E	I	K	R	T
MOR07923	V	P	-	-	F	T	F	G	Q	G	T	K	V	E	I	K	R	T
MOR07924	N	P	-	-	F	T	F	G	Q	G	T	K	V	E	I	K	R	T
MOR07925	P	P	-	-	H	T	F	G	Q	G	T	K	V	E	I	K	R	T
MOR07926	M	P	-	-	F	T	F	G	Q	G	T	K	V	E	I	K	R	T

	CDR 3						Framework 4									
											HpaI				MscI	
VL13	X	X	X	X	X	X	F	G	G	G	T	K	L	T	V	L	G	Q
MOR07919	T	E	F	T	Y	V	F	G	G	G	T	K	L	T	V	L	G	Q

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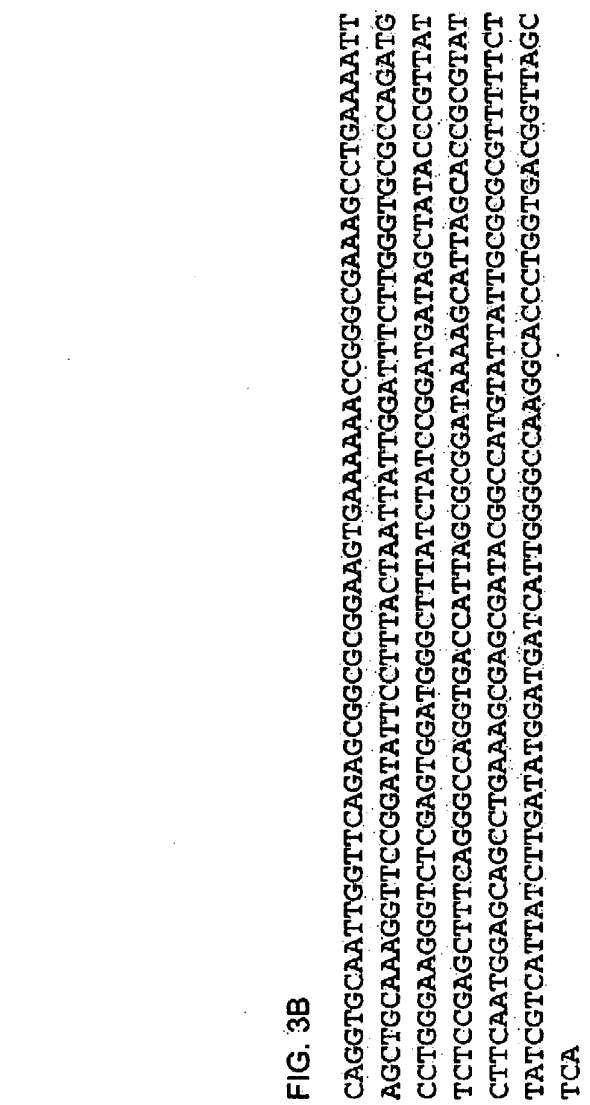
[Fig. 3A]

FIG. 3A

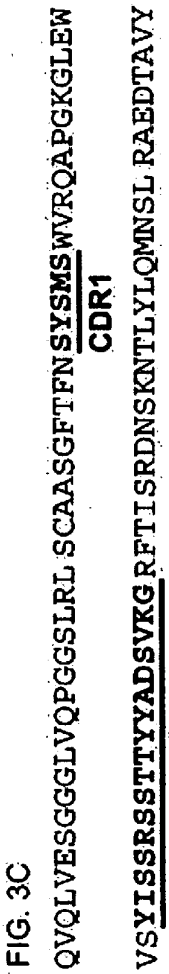

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CCTGGGAAGGGTCTCGAGTGGGTGAGCTATATCTCTTCTCGTTCTAGCACTACCTATTAT
GCGGATAGCGTGAAAGGCCGTTTTACCATTTCACGTGATAATTCGAAAAACACCCTGTAT
CTGCAAATGAACAGCCTGCGTGCGGAAGATACGGCCGTGTATTATTGCGCGCGTGGTTAT
TTTCATGGTATGGATTATTGGGGCCAAGGCACCCTGGTGACGGTTAGCTCA

[Fig. 3B]

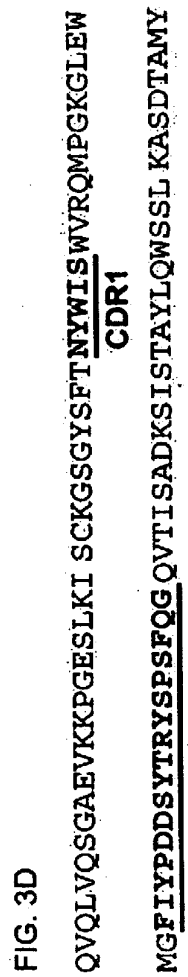
FIG. 3B


CAGGTGCAATTGGTTCAGAGCGGCGCGGAAGTGAAAAAACCGGGCGAAAGCCTGAAAATT
AGCTGCAAAGGTTCCGGATATTCCTTTACTAATTATTGGATTTCTTGGGTGCGCCAGATG
CCTGGGAAGGGTCTCGAGTGGATGGGCTTTATCTATCCGGATGATAGCTATACCCGTTAT
TCTCCGAGCTTTCAGGGCCAGGTGACCATTAGCGCGGATAAAAGCATTAGCACCGCGTAT
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TCA

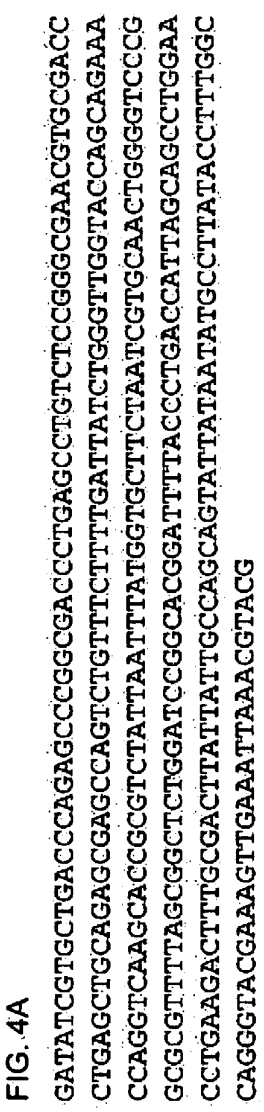
[Fig. 3C]


FIG. 3C
QVQLVESGGGLVQPGGSLRLSCAASGFTFNSYSMSWVRQAPGKGLEW
CDR1
VSYISSRSSTTYYADSVKGRFTISRDNSKNTLYLQMNSLRAEDTAVY
CDR2
YCARGYFHGMDYWGQGTLVTVSS
CDR3

[Fig. 3D]


FIG. 3D
QVQLVQSGAEVKKPGESLKI SCKGSGYSFTNYWIS WVRQMPGKGLEW
CDR1
MGFIYPDDSYTRYSPSFQG QVTISADKSISTAYLQWSSL KASDTAMY
CDR2
YCARFSYRHYLDMDDH WGQGTLVTVSS
CDR3

[Fig. 4A]


FIG. 4A
GATATCGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGTCTCCGGGCGAACGTGCGACC
CTGAGCTGCAGAGCGAGCCAGTCTGTTTCTTTTGATTATCTGGGTTGGTACCAGCAGAAA
CCAGGTCAAGCACCGCGTCTATTTAATTTATGGTGCTTCTAATCGTGCAACTGGGGTCCCG
GCGCGTTTTAGCGGCTCTGGATCCGGCACGGATTTTACCCTGACCATTAGCAGCCTGGAA
CCTGAAGACTTTGCGACTTATTATTGCCAGCAGTATTATAATATGCCTTATACCTTTGGC
CAGGGTACCGAAAGTTGAAATTAAACGTACG

[Fig. 4B]

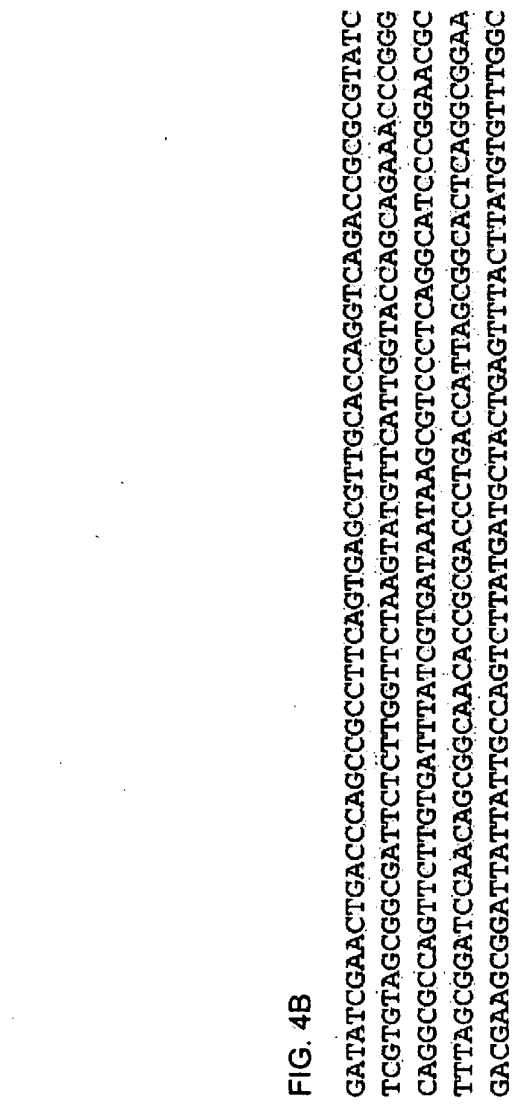

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CAGGCGCCAGTTCTTGTGATTTATCGTGATAATAAGCGTCCCTCAGGCATCCCGGAACGC
TTTAGCGGATCCAACAGCGGCAACACCGCGACCCTGACCATTAGCGGCACTCAGGCGGAA
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GGCGGCACGAAGTTAACCGTTCTTGGCCAG

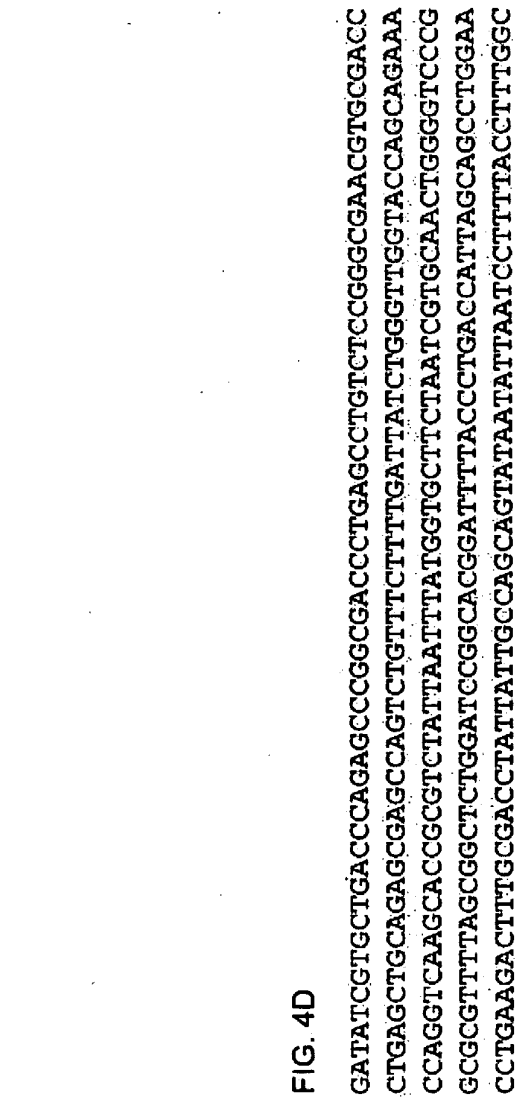
FIG. 4B

[Fig. 4C]

FIG. 4C

GATATCGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGTCTCCGGGCGAACGTGCGACC
CTGAGCTGCAGAGCGAGCCAGTCTGTTTCTTTTGATTATCTGGGTTGGTACCAGCAGAAA
CCAGGTCAAGCACCGCGTCTATTAATTTATGGTGCTTCTAATCGTGCAACTGGGGTCCCG
GCGCGTTTTAGCGGCTCTGGATCCGGCACGGATTTTACCCTGACCATTAGCAGCCTGGAA
CCTGAAGACTTTGCGACCTATTATTGCTTTCAGTATCTTATTGTTCCTTTTACCTTTGGC
CAGGGTACGAAAGTTGAAATTAAACGTACG

[Fig. 4D]


FIG. 4D
GATATCGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGTCTCCGGGCGAACGTGCGACC
CTGAGCTGCAGAGCGAGCCAGTCTGTTTCTTTTGATTATCTGGGTTGGTACCAGCAGAAA
CCAGGTCAAGCACCGCGTCTATTAATTTATGGTGCTTCTAATCGTGCAACTGGGGTCCCG
GCGCGTTTTAGCGGCTCTGGATCCGGCACGGATTTTACCCTGACCATTAGCAGCCTGGAA
CCTGAAGACTTTGCGACCTATTATTGCCAGCAGTATAATATTAATCCTTTTACCTTTGGC
CAGGGTACGAAAGTTGAAATTAAACGTACG

[Fig. 4E]

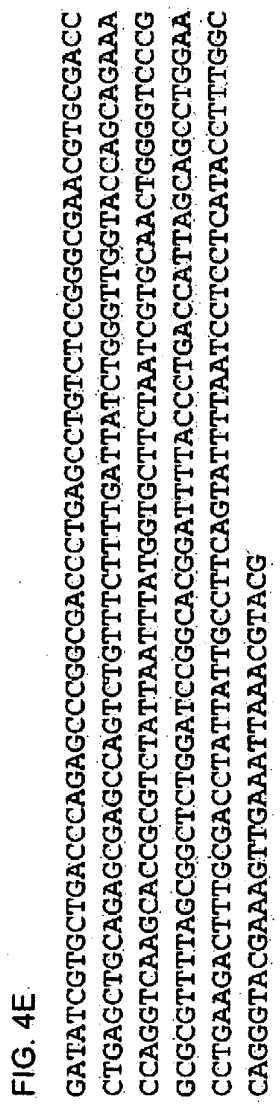
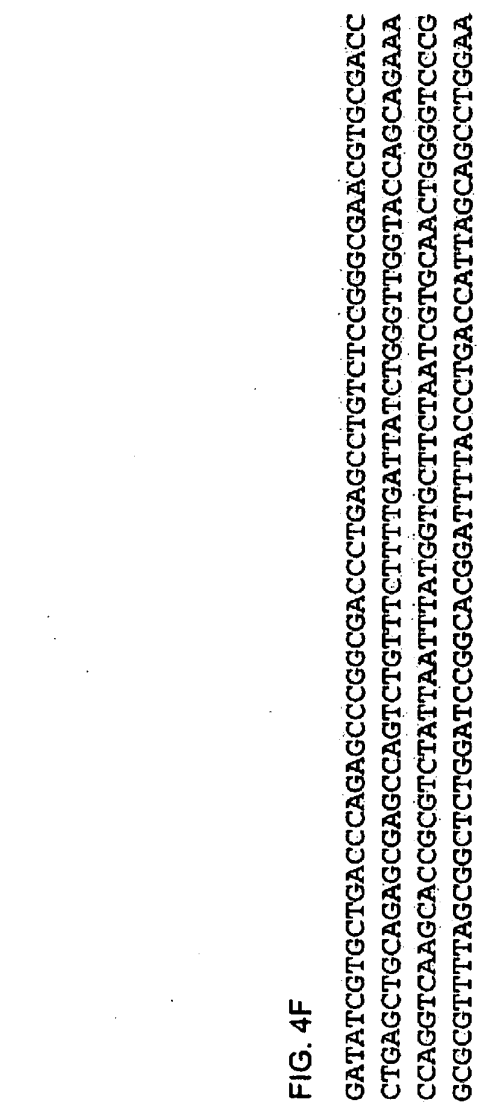


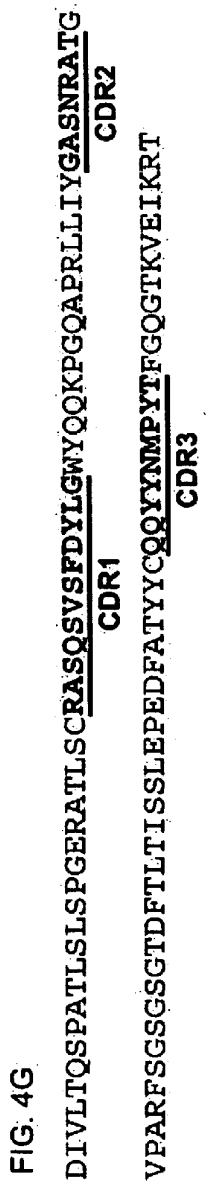
FIG. 4E

GATATCGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGTCTCCGGGCGAACGTGCGACC
CTGAGCTGCAGAGCGAGCCAGTCTGTTTCTTTGATTATCTGGGTTGGTACCAGCAGAAA
CCAGGTCAAGCACCGCGTCTATTAATTTATGGTGCTTCTAATCGTGCAACTGGGGTCCCG
GCGCGTTTTAGCGGCTCTGGATCCGGCACGGATTTTACCCTGACCATTAGCAGCCTGGAA
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CAGGGTACGAAAGTTGAAATTAAACGTACG

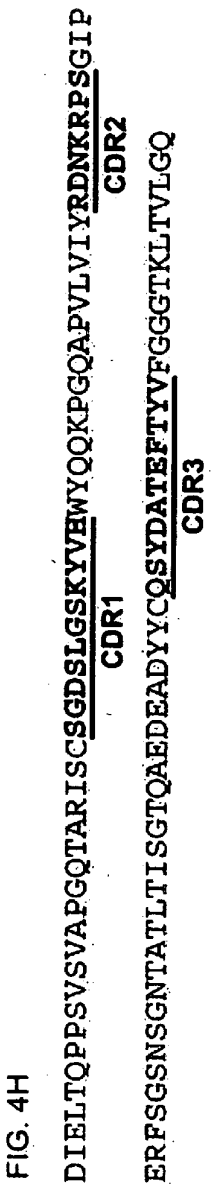
[Fig. 4F]


FIG. 4F
GATATCGTGCTGACCCAGAGCCCGGCGACCCTGAGCCTGTCTCCGGGCGAACGTGCGACC
CTGAGCTGCAGAGCGAGCCAGTCTGTTTCTTTTGATTATCTGGGTTGGTACCAGCAGAAA
CCAGGTCAAGCACCGCGTCTATTAATTTATGGTGCTTCTAATCGTGCAACTGGGGTCCCG
GCGCGTTTTAGCGGCTCTGGATCCGGCACGGATTTTACCCTGACCATTAGCAGCCTGGAA
CCTGAAGACTTTGCGACCTATTATTGCTTTCAGGCTCTTATTATGCCTTTTACCTTTGGC
CAGGGTACGAAAGTTGAAATTAAACGTACG

[Fig. 4G]

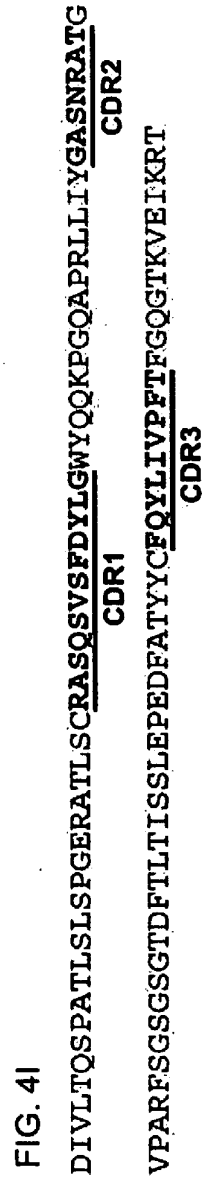

FIG. 4G
DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATG
CDR1
CDR2
VPARFSGSGSGTDFTLTISSLEPEDFATYYCQQYYNMPYTFGQGTKVEIKRT
CDR3

[Fig. 4H]

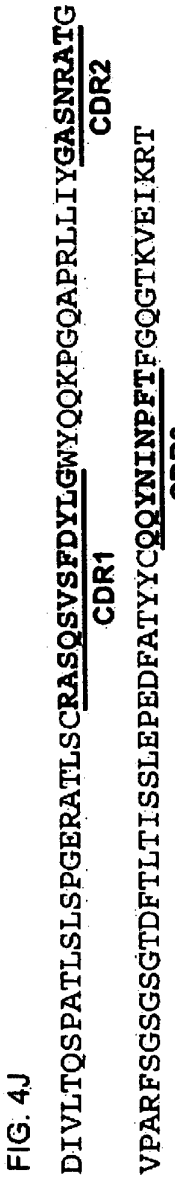

FIG. 4H
DIELTQPPSVSVAPGQTARISCSGDSLGSKYVEWYQQKPGQAPVLVIYRDNKRPSGIP
CDR1
CDR2
ERFSGSNSGNTATLTISGTQAEDEADYYCQSYDATEFTYVFGGGTKLTVLGQ
CDR3

[Fig. 4I]

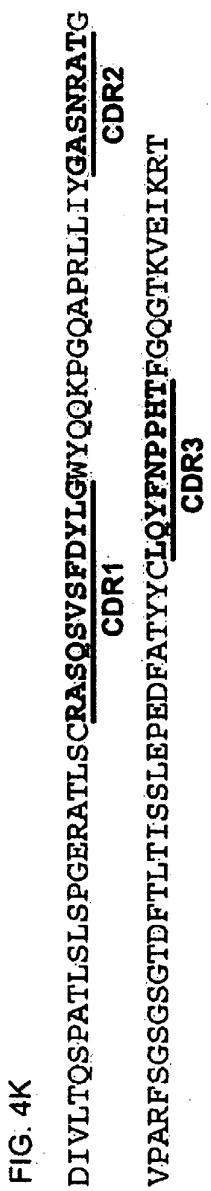
FIG. 4I


DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATG
CDR1
CDR2
VPARFSGSGSGTDFTLTISSLEPEDFATYYCFQYLIVPFTFGQGTKVEIKRT
CDR3

[Fig. 4J]


FIG. 4J
DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATG
CDR1
CDR2
VPARFSGSGSGTDFTLTISSLEPEDFATYYCQQYINNPFTFGQGTKVEIKRT
CDR3

[Fig. 4K]


FIG. 4K
DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATG
CDR1
CDR2
VPARFSGSGSGTDFTLTISSLEPEDFATYYCLQYFNFPHTFGQGTKVEIKRT
CDR3

[Fig. 4L]

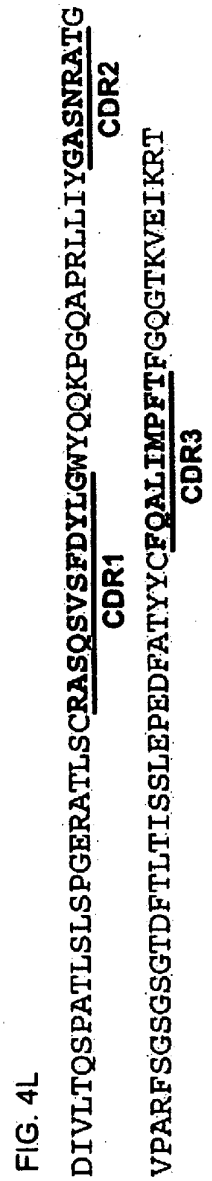

FIG. 4L
DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATG
CDR1
CDR2
VPARFSGSGSGTDFTLTISSLEPEDFATYYCFQALIMPFTFGQGTKVEIKRT
CDR3

FIG. 5

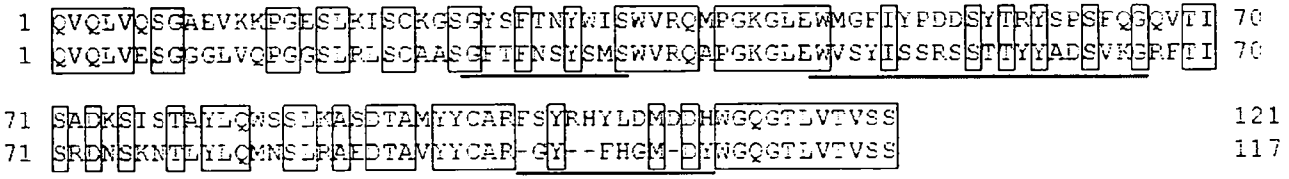

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1 QVQLVESGGGLVQPGGSLRLSCAASGFTFNSYSMSWVRQAPGKGLEWVSYISSRSSTIYYADSVKGRFTI 70
71 SADKSISTAYLQWSSLKASDTAMYYCARFSYRHYLDMDDHWGQGTLVTVSS 121
71 SRDNSKNTLYLQMNSLRAEDTAVYYCAR-GY--FHGM-DYWGQGTLVTVSS 117

FIG. 6

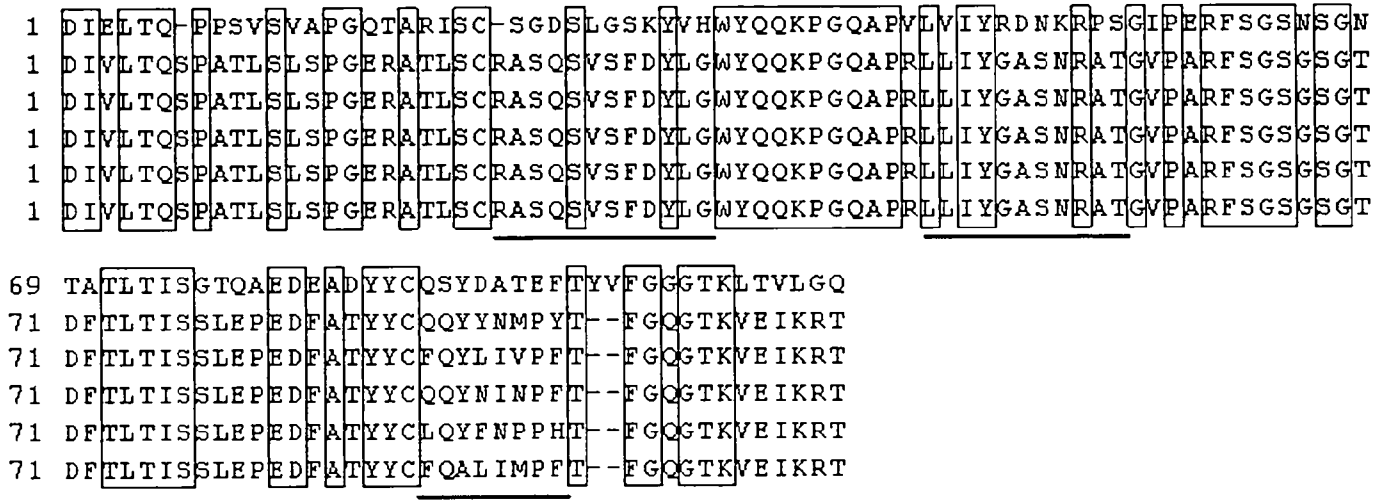

1 DIELTQ-PPSVSVAPGQTARISC-SGDSLGSKYVHWYQQKPGQAPVLVIYRDNKRPSGIPERFSGSNSGN
1 DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATGVPARFSGSGSGT
1 DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATGVPARFSGSGSGT
1 DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATGVPARFSGSGSGT
1 DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATGVPARFSGSGSGT
1 DIVLTQSPATLSLSPGERATLSCRASQSVSFDYLGWYQQKPGQAPRLLIYGASNRATGVPARFSGSGSGT
69 TATLTISGTQAEDEADYYCQSYDATEFTYVFGGGTKLTVLGQ
71 DFTLTISSLEPEDFATYYCQQYYNMPYT--FGQGTKVEIKRT
71 DFTLTISSLEPEDFATYYCFQYLIVPFT--FGQGTKVEIKRT
71 DFTLTISSLEPEDFATYYCQQYNINPFT--FGQGTKVEIKRT
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FIG 7A

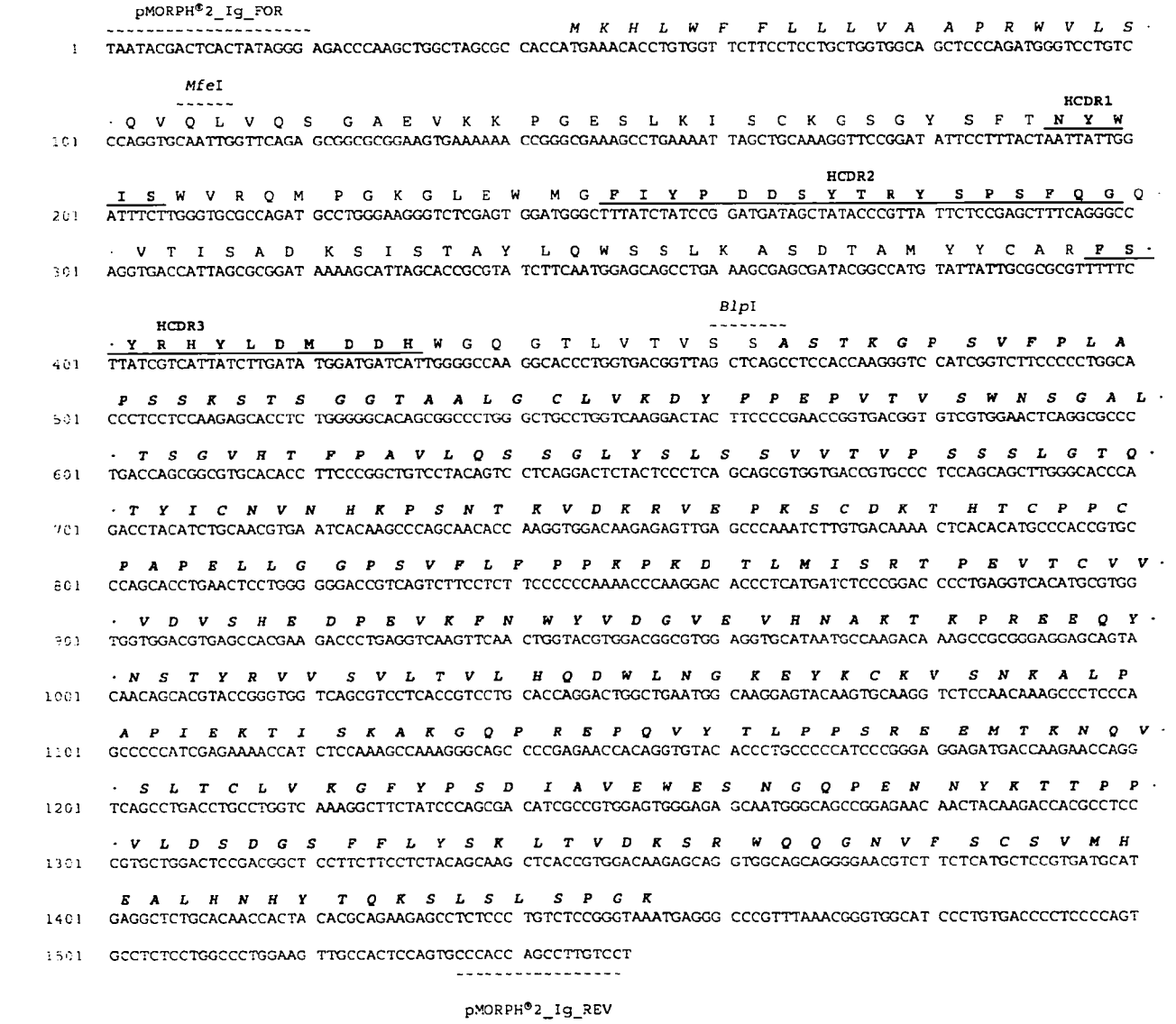


FIG. 7B

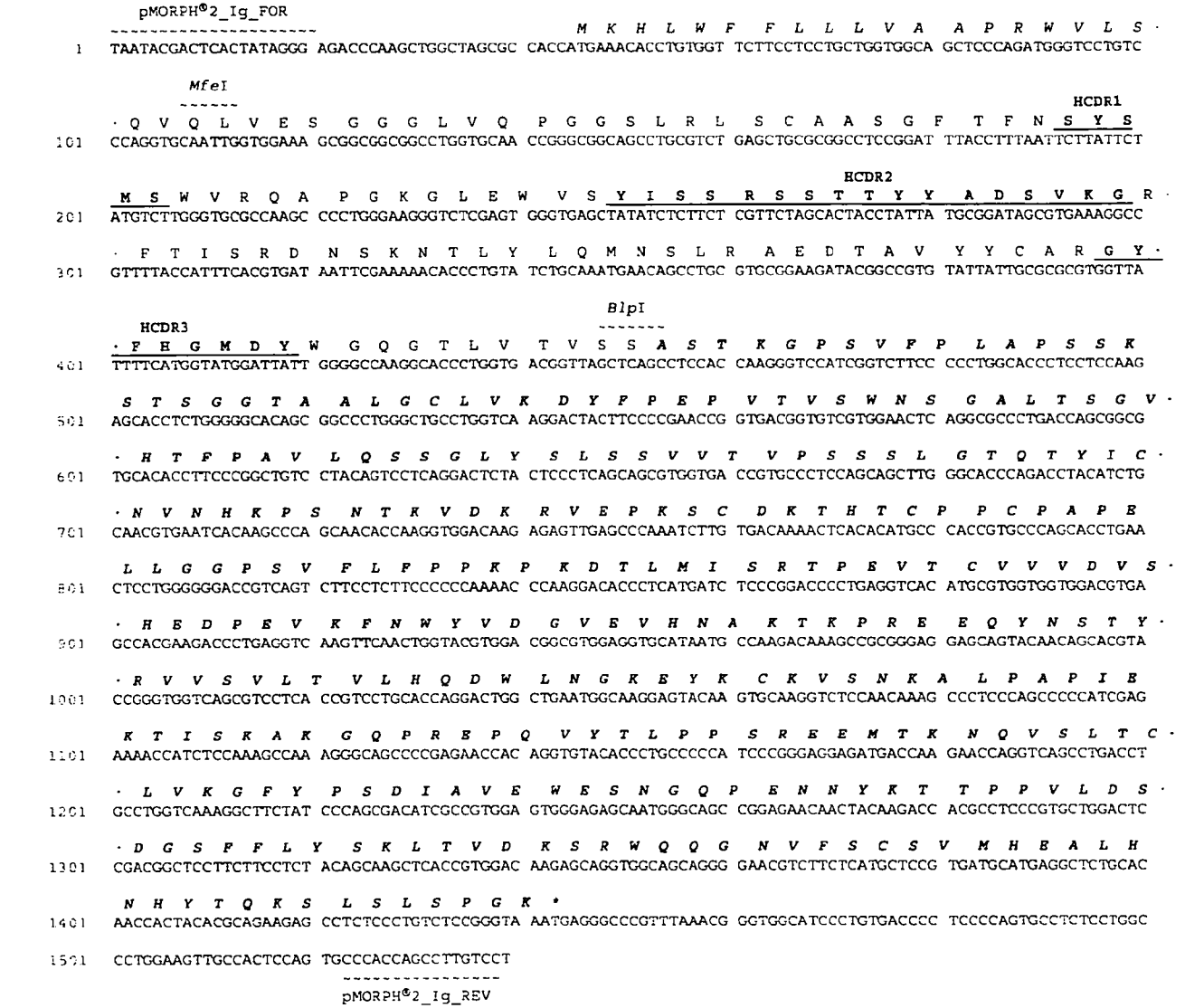


FIG. 8

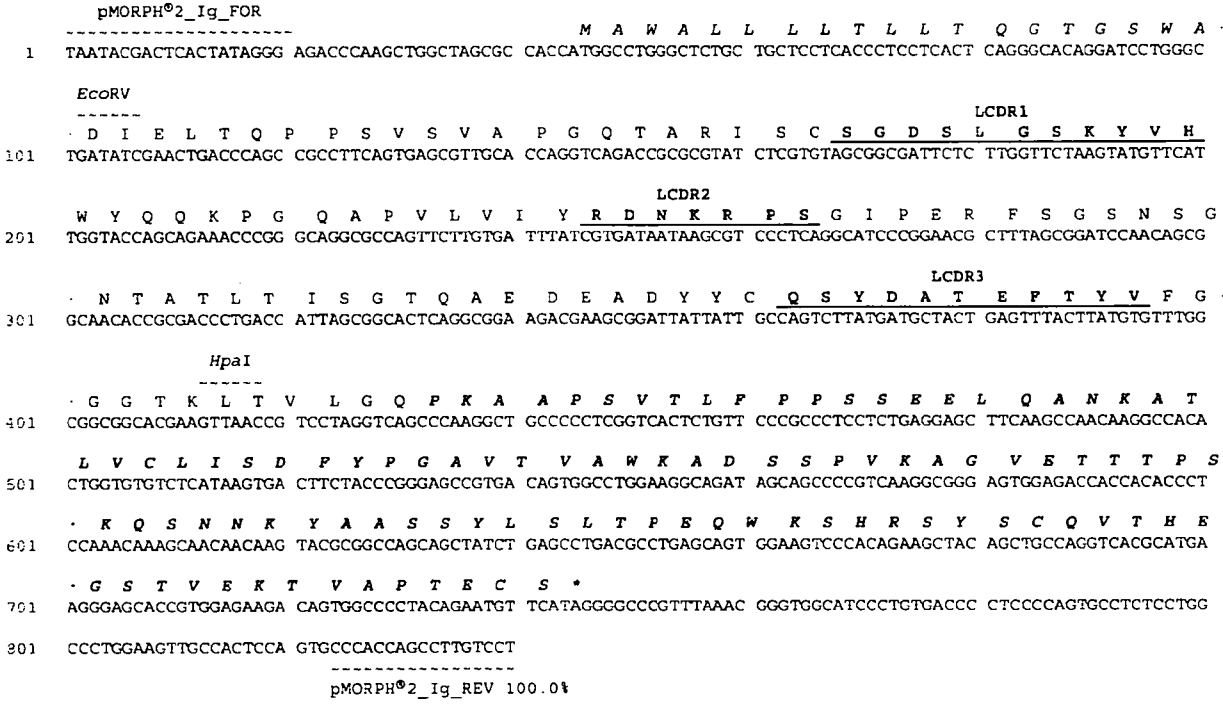
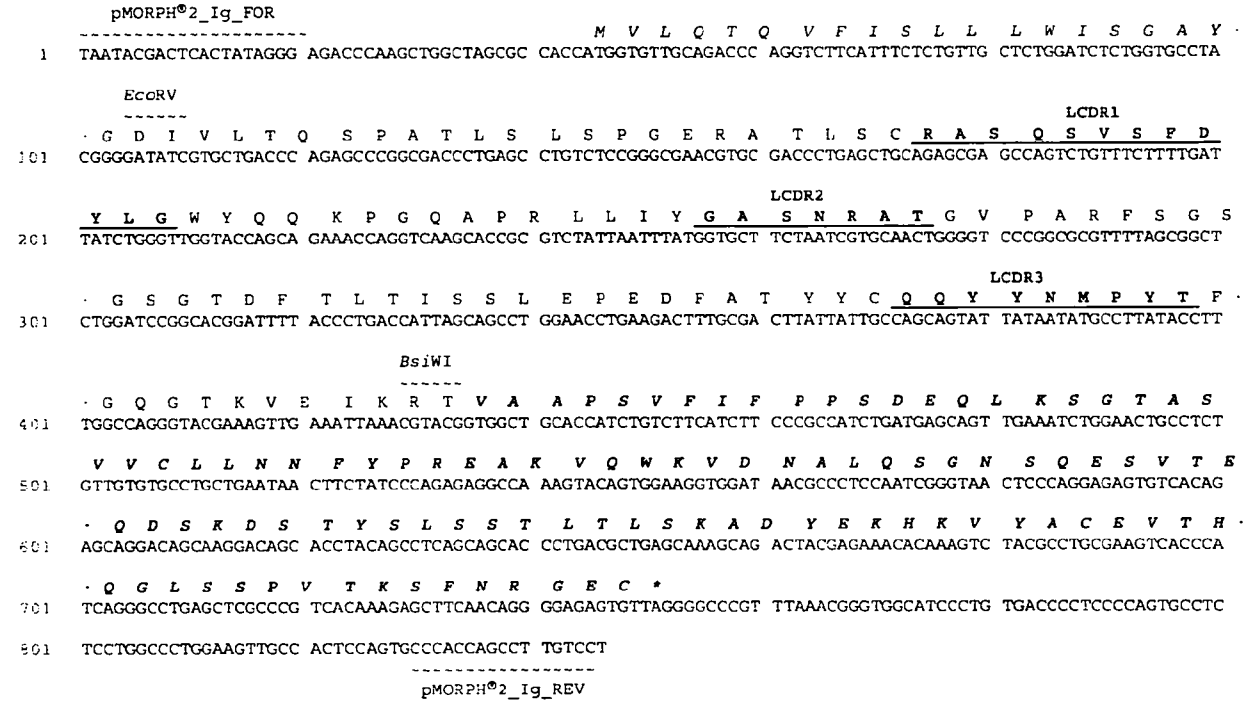

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LCDR1
D I E L T Q P P S V S V A P G Q T A R I S C S G D S L G S K Y V H
101 TGATATCGAACTGACCCAGC CGCCTTCAGTGAGCGTTGCA CCAGGTCAGACCGCGCGTAT CTCGTGTAGCGGCGATTCTC TTGGTTCTAAGTATGTTCAT
LCDR2
W Y Q Q K P G Q A P V L V I Y R D N K R P S G I P E R F S G S N S G
201 TGGTACCAGCAGAAACCCGG GCAGGCGCCAGTTCTTGTGA TTTATCGTGATAATAAGCGT CCCTCAGGCATCCCGGAACG CTTTAGCGGATCCAACAGCG
LCDR3
N T A T L T I S G T Q A E D E A D Y Y C Q S Y D A T E F T Y V F G
301 GCAACACCGCGACCCTGACC ATTAGCGGCACTCAGGCGGA AGACGAAGCGGATTATTATT GCCAGTCTTATGATGCTACT GAGTTTACTTATGTGTTTGG
HpaI
G G T K L T V L G Q P K A A P S V T L F P P S S E E L Q A N K A T
401 CGGCGGCACGAAGTTAACCG TCCTAGGTCAGCCCAAGGCT GCCCCCTCGGTCACTCTGTT CCCGCCCTCCTCTGAGGAGC TTCAAGCCAACAAGGCCACA
L V C L I S D F Y P G A V T V A W K A D S S P V K A G V E T T T P S
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K Q S N N K Y A A S S Y L S L T P E Q W K S H R S Y S C Q V T H E
601 CCAAACAAAGCAACAACAAG TACGCGGCCAGCAGCTATCT GAGCCTGACGCCTGAGCAGT GGAAGTCCCACAGAAGCTAC AGCTGCCAGGTCACGCATGA
G S T V E K T V A P T E C S *
701 AGGGAGCACCGTGGAGAAGA CAGTGGCCCCTACAGAATGT TCATAGGGGCCCGTTTAAAC GGGTGGCATCCCTGTGACCC CTCCCCAGTGCCTCTCCTGG
801 CCCTGGAAGTTGCCACTCCA GTGCCCACCAGCCTTGTCCT
pMORPH®2_Ig_REV 100.0%

FIG. 9A


pMORPH®2_Ig_FOR
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LCDR1
· G D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F D
101 CGGGGATATCGTGCTGACCC AGAGCCCGGCGACCCTGAGC CTGTCTCCGGGCGAACGTGC GACCCTGAGCTGCAGAGCGA GCCAGTCTGTTTCTTTTGAT
LCDR2
Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A R F S G S ·
201 TATCTGGGTTGGTACCAGCA GAAACCAGGTCAAGCACCGC GTCTATTAATTTATGGTGCT TCTAATCGTGCAACTGGGGT CCCGGCGCGTTTTAGCGGCT
LCDR3
· G S G T D F T L T I S S L E P E D F A T Y Y C Q Q Y Y N M P Y T F ·
301 CTGGATCCGGCACGGATTTT ACCCTGACCATTAGCAGCCT GGAACCTGAAGACTTTGCGA CTTATTATTGCCAGCAGTAT TATAATATGCCTTATACCTT
BsiWI
· G Q G T K V E I K R T V A A P S V F I F P P S D E Q L K S G T A S
401 TGGCCAGGGTACGAAAGTTG AAATTAAACGTACGGTGGCT GCACCATCTGTCTTCATCTT CCCGCCATCTGATGAGCAGT TGAAATCTGGAACTGCCTCT
V V C L L N N F Y P R E A K V Q W K V D N A L Q S G N S Q E S V T E ·
501 GTTGTGTGCCTGCTGAATAA CTTCTATCCCAGAGAGGCCA AAGTACAGTGGAAGGTGGAT AACGCCCTCCAATCGGGTAA CTCCCAGGAGAGTGTCACAG
· Q D S K D S T Y S L S S T L T L S K A D Y E K H K V Y A C E V T H ·
601 AGCAGGACAGCAAGGACAGC ACCTACAGCCTCAGCAGCAC CCTGACGCTGAGCAAAGCAG ACTACGAGAAACACAAAGTC TACGCCTGCGAAGTCACCCA
· Q G L S S P V T K S F N R G E C *
701 TCAGGGCCTGAGCTCGCCCG TCACAAAGAGCTTCAACAGG GGAGAGTGTTAGGGGCCCGT TTAAACGGGTGGCATCCCTG TGACCCCTCCCCAGTGCCTC
801 TCCTGGCCCTGGAAGTTGCC ACTCCAGTGCCCACCAGCCT TGTCCT
pMORPH®2_Ig_REV

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FIG. 9B

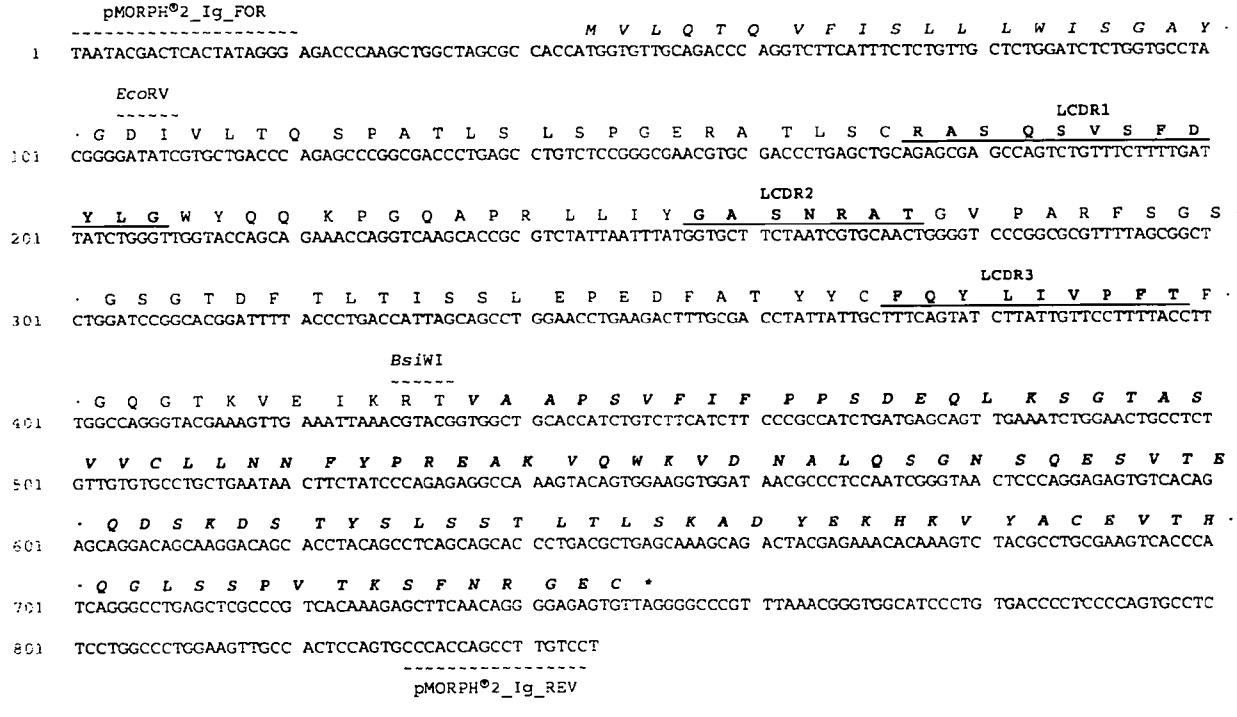

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EcoRV
LCDR1
· G D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F D
101 CGGGGATATCGTGCTGACCC AGAGCCCGGCGACCCTGAGC CTGTCTCCGGGCGAACGTGC GACCCTGAGCTGCAGAGCGA GCCAGTCTGTTTCTTTTGAT
LCDR2
Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A R F S G S ·
201 TATCTGGGTTGGTACCAGCA GAAACCAGGTCAAGCACCGC GTCTATTAATTTATGGTGCT TCTAATCGTGCAACTGGGGT CCCGGCGCGTTTTAGCGGCT
LCDR3
· G S G T D F T L T I S S L E P E D F A T Y Y C F Q Y L I V P F T F ·
301 CTGGATCCGGCACGGATTTT ACCCTGACCATTAGCAGCCT GGAACCTGAAGACTTTGCGA CCTATTATTGCTTTCAGTAT CTTATTGTTCCTTTTACCTT
BsiWI
· G Q G T K V E I K R T V A A P S V F I F P P S D E Q L K S G T A S
401 TGGCCAGGGTACGAAAGTTG AAATTAAACGTACGGTGGCT GCACCATCTGTCTTCATCTT CCCGCCATCTGATGAGCAGT TGAAATCTGGAACTGCCTCT
V V C L L N N F Y P R E A K V Q W K V D N A L Q S G N S Q E S V T E ·
501 GTTGTGTGCCTGCTGAATAA CTTCTATCCCAGAGAGGCCA AAGTACAGTGGAAGGTGGAT AACGCCCTCCAATCGGGTAA CTCCCAGGAGAGTGTCACAG
· Q D S K D S T Y S L S S T L T L S K A D Y E K H K V Y A C E V T H ·
601 AGCAGGACAGCAAGGACAGC ACCTACAGCCTCAGCAGCAC CCTGACGCTGAGCAAAGCAG ACTACGAGAAACACAAAGTC TACGCCTGCGAAGTCACCCA
· Q G L S S P V T K S F N R G E C *
701 TCAGGGCCTGAGCTCGCCCG TCACAAAGAGCTTCAACAGG GGAGAGTGTTAGGGGCCCGT TTAAACGGGTGGCATCCCTG TGACCCCTCCCCAGTGCCTC
801 TCCTGGCCCTGGAAGTTGCC ACTCCAGTGCCCACCAGCCT TGTCCT
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FIG. 9C

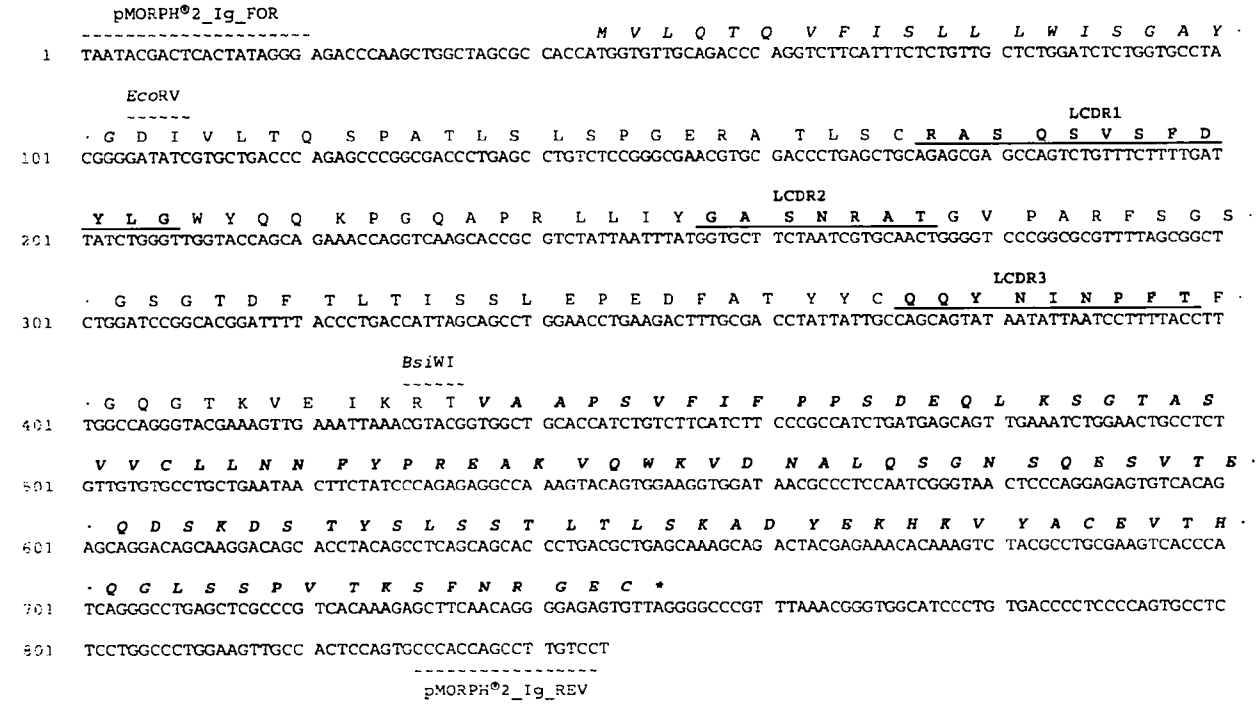

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EcoRV
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G D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F D
101 CGGGGATATCGTGCTGACCC AGAGCCCGGCGACCCTGAGC CTGTCTCCGGGCGAACGTGC GACCCTGAGCTGCAGAGCGA GCCAGTCTGTTTCTTTTGAT
LCDR2
Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A R F S G S
201 TATCTGGGTTGGTACCAGCA GAAACCAGGTCAAGCACCGC GTCTATTAATTTATGGTGCT TCTAATCGTGCAACTGGGGT CCCGGCGCGTTTTAGCGGCT
LCDR3
G S G T D F T L T I S S L E P E D F A T Y Y C Q Q Y N I N P F T F
301 CTGGATCCGGCACGGATTTT ACCCTGACCATTAGCAGCCT GGAACCTGAAGACTTTGCGA CCTATTATTGCCAGCAGTAT AATATTAATCCTTTTACCTT
BsiWI
G Q G T K V E I K R T V A A P S V F I F P P S D E Q L K S G T A S
401 TGGCCAGGGTACGAAAGTTG AAATTAAACGTACGGTGGCT GCACCATCTGTCTTCATCTT CCCGCCATCTGATGAGCAGT TGAAATCTGGAACTGCCTCT
V V C L L N N F Y P R E A K V Q W K V D N A L Q S G N S Q E S V T E
501 GTTGTGTGCCTGCTGAATAA CTTCTATCCCAGAGAGGCCA AAGTACAGTGGAAGGTGGAT AACGCCCTCCAATCGGGTAA CTCCCAGGAGAGTGTCACAG
Q D S K D S T Y S L S S T L T L S K A D Y E K H K V Y A C E V T H
601 AGCAGGACAGCAAGGACAGC ACCTACAGCCTCAGCAGCAC CCTGACGCTGAGCAAAGCAG ACTACGAGAAACACAAAGTC TACGCCTGCGAAGTCACCCA
Q G L S S P V T K S F N R G E C *
701 TCAGGGCCTGAGCTCGCCCG TCACAAAGAGCTTCAACAGG GGAGAGTGTTAGGGGCCCGT TTAAACGGGTGGCATCCCTG TGACCCCTCCCCAGTGCCTC
801 TCCTGGCCCTGGAAGTTGCC ACTCCAGTGCCCACCAGCCT TGTCCT
pMORPH®2_Ig_REV

FIG. 9D

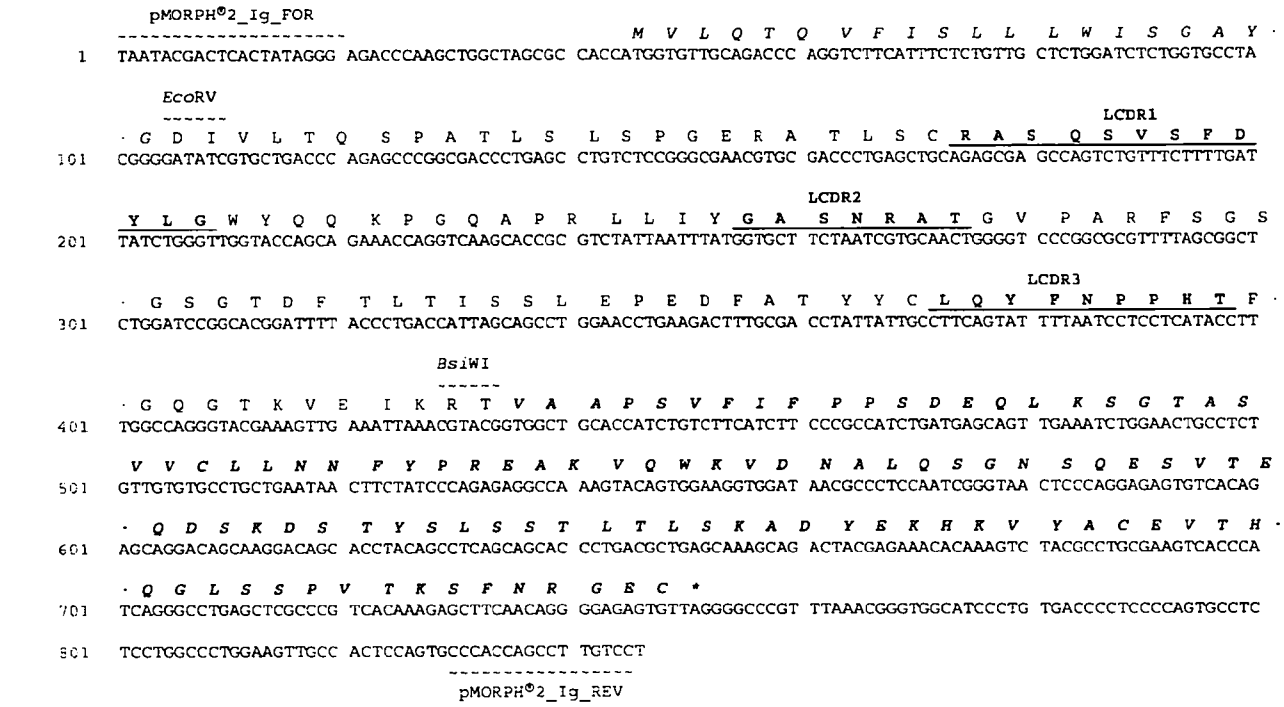
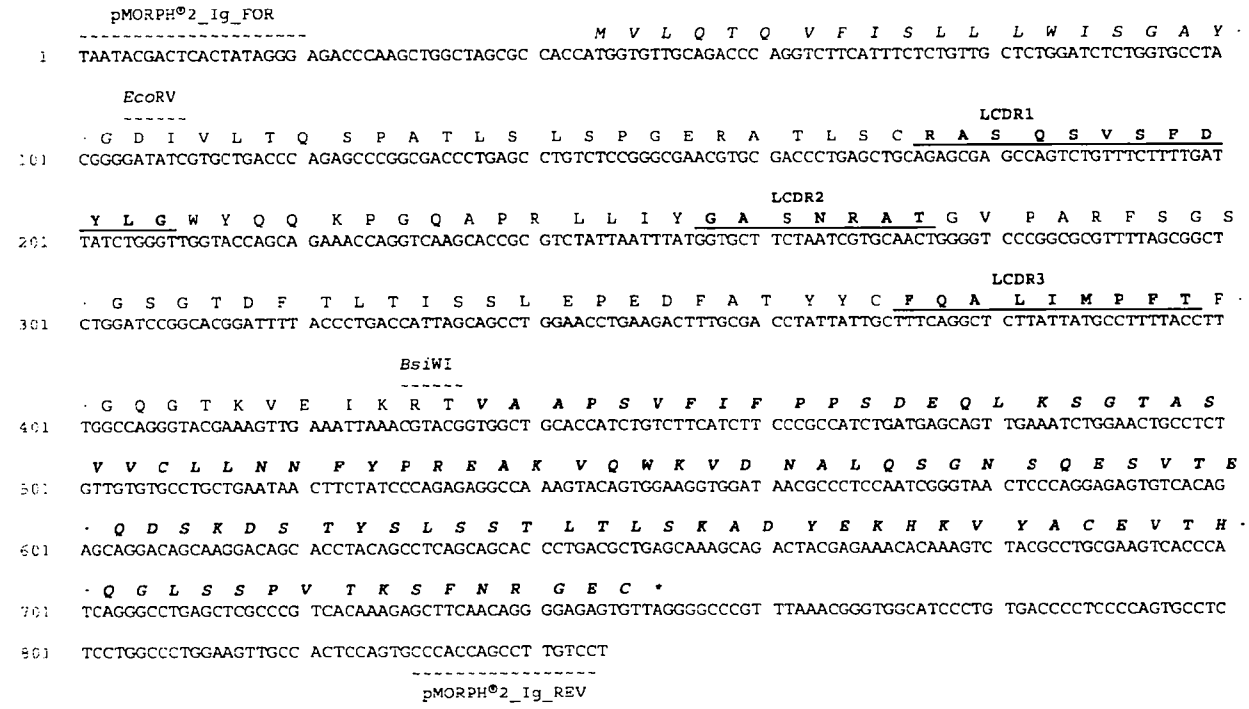

pMORPH®2_Ig_FOR
M V L Q T Q V F I S L L L W I S G A Y
1 TAATACGACTCACTATAGGG AGACCCAAGCTGGCTAGCGC CACCATGGTGTTGCAGACCC AGGTCTTCATTTCTCTGTTG CTCTGGATCTCTGGTGCCTA
EcoRV
LCDR1
G D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F D
101 CGGGGATATCGTGCTGACCC AGAGCCCGGCGACCCTGAGC CTGTCTCCGGGCGAACGTGC GACCCTGAGCTGCAGAGCGA GCCAGTCTGTTTCTTTTGAT
LCDR2
Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A R F S G S
201 TATCTGGGTTGGTACCAGCA GAAACCAGGTCAAGCACCGC GTCTATTAATTTATGGTGCT TCTAATCGTGCAACTGGGGT CCCGGCGCGTTTTAGCGGCT
LCDR3
G S G T D F T L T I S S L E P E D F A T Y Y C L Q Y F N P P H T F
301 CTGGATCCGGCACGGATTTT ACCCTGACCATTAGCAGCCT GGAACCTGAAGACTTTGCGA CCTATTATTGCCTTCAGTAT TTTAATCCTCCTCATACCTT
BsiWI
G Q G T K V E I K R T V A A P S V F I F P P S D E Q L K S G T A S
401 TGGCCAGGGTACGAAAGTTG AAATTAAACGTACGGTGGCT GCACCATCTGTCTTCATCTT CCCGCCATCTGATGAGCAGT TGAAATCTGGAACTGCCTCT
V V C L L N N F Y P R E A K V Q W K V D N A L Q S G N S Q E S V T E
501 GTTGTGTGCCTGCTGAATAA CTTCTATCCCAGAGAGGCCA AAGTACAGTGGAAGGTGGAT AACGCCCTCCAATCGGGTAA CTCCCAGGAGAGTGTCACAG
Q D S K D S T Y S L S S T L T L S K A D Y E K H K V Y A C E V T H
601 AGCAGGACAGCAAGGACAGC ACCTACAGCCTCAGCAGCAC CCTGACGCTGAGCAAAGCAG ACTACGAGAAACACAAAGTC TACGCCTGCGAAGTCACCCA
Q G L S S P V T K S F N R G E C *
701 TCAGGGCCTGAGCTCGCCCG TCACAAAGAGCTTCAACAGG GGAGAGTGTTAGGGGCCCGT TTAAACGGGTGGCATCCCTG TGACCCCTCCCCAGTGCCTC
801 TCCTGGCCCTGGAAGTTGCC ACTCCAGTGCCCACCAGCCT TGTCCT
pMORPH®2_Ig_REV

FIG. 9E


pMORPH®2_Ig_FOR
M V L Q T Q V F I S L L L W I S G A Y ·
1 TAATACGACTCACTATAGGG AGACCCAAGCTGGCTAGCGC CACCATGGTGTTGCAGACCC AGGTCTTCATTTCTCTGTTG CTCTGGATCTCTGGTGCCTA
EcoRV
LCDR1
· G D I V L T Q S P A T L S L S P G E R A T L S C R A S Q S V S F D
101 CGGGGATATCGTGCTGACCC AGAGCCCGGCGACCCTGAGC CTGTCTCCGGGCGAACGTGC GACCCTGAGCTGCAGAGCGA GCCAGTCTGTTTCTTTTGAT
LCDR2
Y L G W Y Q Q K P G Q A P R L L I Y G A S N R A T G V P A R F S G S ·
201 TATCTGGGTTGGTACCAGCA GAAACCAGGTCAAGCACCGC GTCTATTAATTTATGGTGCT TCTAATCGTGCAACTGGGGT CCCGGCGCGTTTTAGCGGCT
LCDR3
· G S G T D F T L T I S S L E P E D F A T Y Y C F Q A L I M P F T F ·
301 CTGGATCCGGCACGGATTTT ACCCTGACCATTAGCAGCCT GGAACCTGAAGACTTTGCGA CCTATTATTGCTTTCAGGCT CTTATTATGCCTTTTACCTT
BsiWI
· G Q G T K V E I K R T V A A P S V F I F P P S D E Q L K S G T A S
401 TGGCCAGGGTACGAAAGTTG AAATTAAACGTACGGTGGCT GCACCATCTGTCTTCATCTT CCCGCCATCTGATGAGCAGT TGAAATCTGGAACTGCCTCT
V V C L L N N F Y P R E A K V Q W K V D N A L Q S G N S Q E S V T E ·
501 GTTGTGTGCCTGCTGAATAA CTTCTATCCCAGAGAGGCCA AAGTACAGTGGAAGGTGGAT AACGCCCTCCAATCGGGTAA CTCCCAGGAGAGTGTCACAG
· Q D S K D S T Y S L S S T L T L S K A D Y E K H K V Y A C E V T H ·
601 AGCAGGACAGCAAGGACAGC ACCTACAGCCTCAGCAGCAC CCTGACGCTGAGCAAAGCAG ACTACGAGAAACACAAAGTC TACGCCTGCGAAGTCACCCA
· Q G L S S P V T K S F N R G E C *
701 TCAGGGCCTGAGCTCGCCCG TCACAAAGAGCTTCAACAGG GGAGAGTGTTAGGGGCCCGT TTAAACGGGTGGCATCCCTG TGACCCCTCCCCAGTGCCTC
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pMORPH®2_Ig_REV

[Fig. 10]

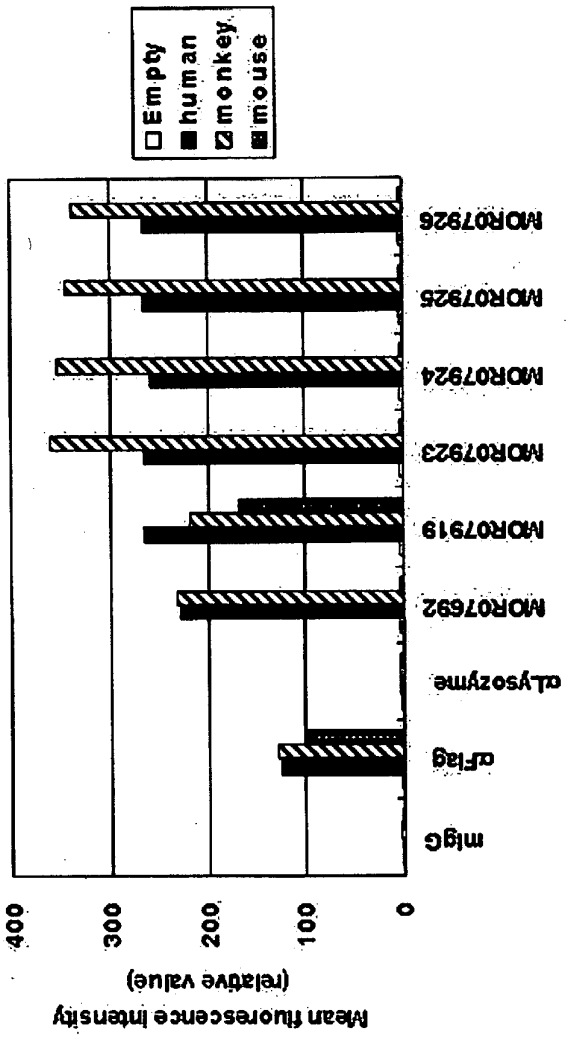
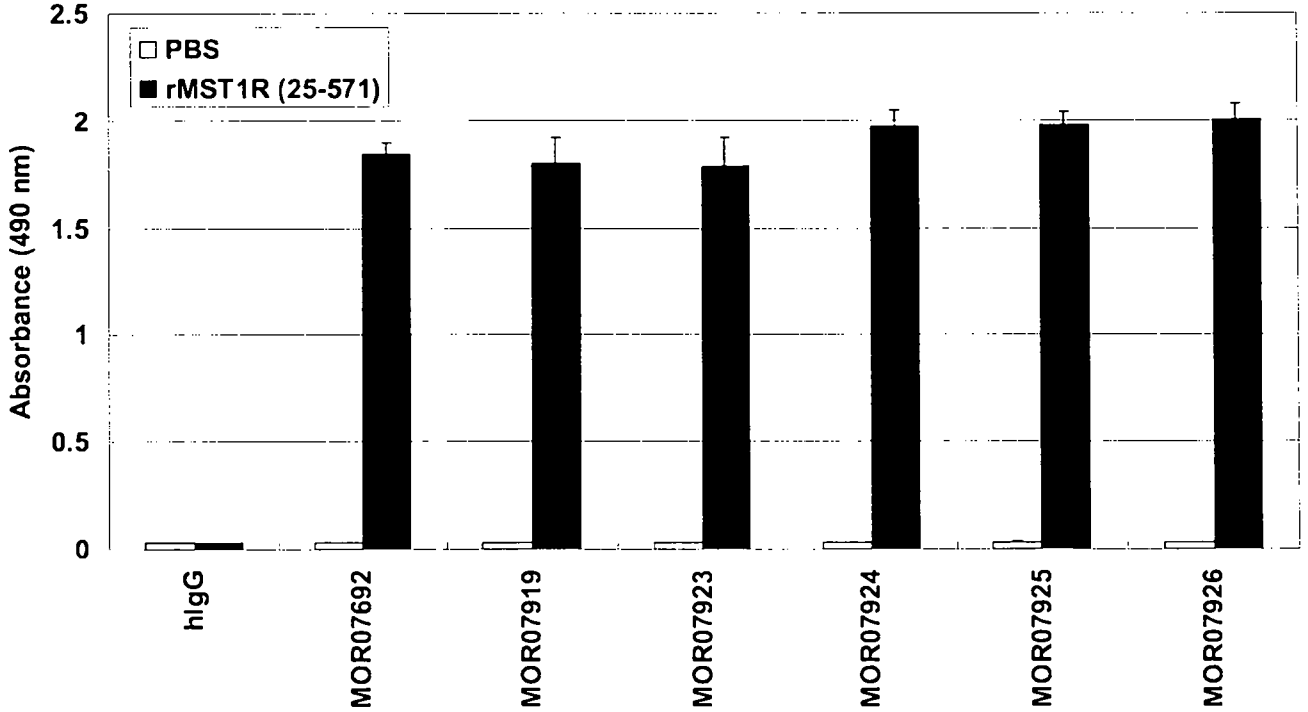

Empty
human
monkey
mouse
hOR7926
hOR7925
hOR7924
hOR7923
hOR7919
hOR7692
αLysozyme
αFlag
mIgG
0
100
200
300
400
Mean fluorescence intensity
(relative value)

FIG. 10

FIG. 11


2.5
2
1.5
1
0.5
0
Absorbance (490 nm)
PBS
rMST1R (25-571)
hIgG
MOR07692
MOR07919
MOR07923
MOR07924
MOR07925
MOR07926

[Fig. 12]

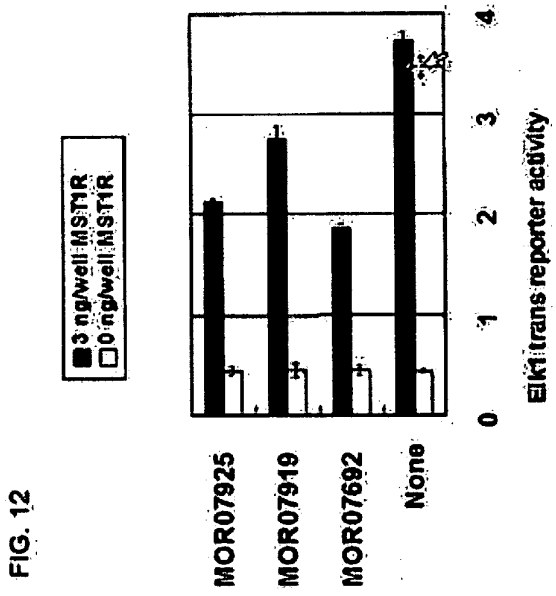

3 ng/well MST1R
0 ng/well MST1R
Elk1 trans-reporter activity
0
1
2
3
4
MOR07925
MOR07919
MOR07692
None

FIG. 12

[Fig. 13]

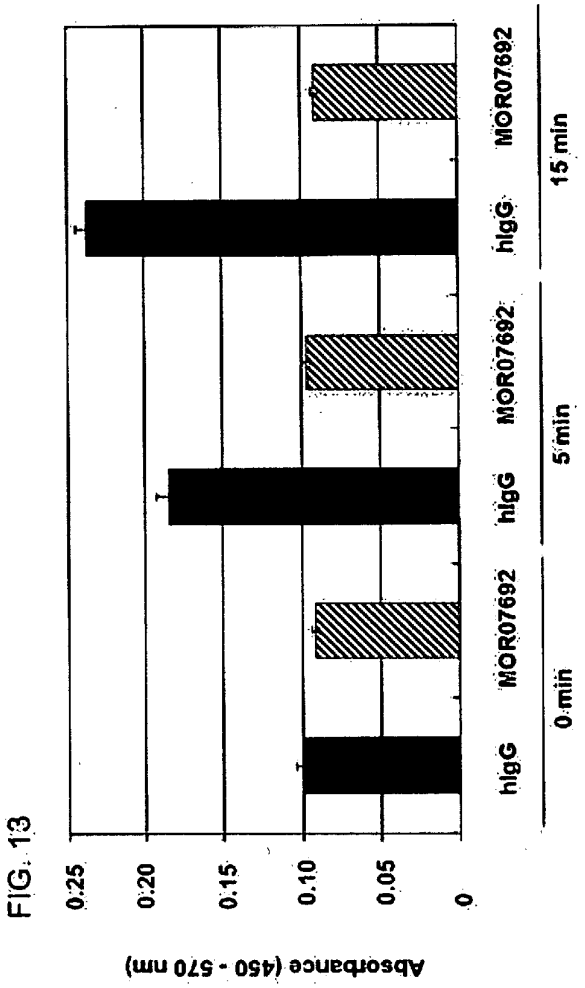
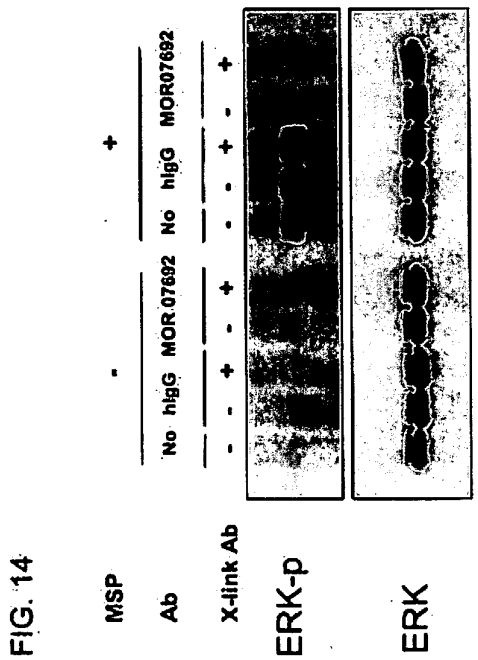

Absorbance (450 - 570 nm)
0.25
0.20
0.15
0.10
0.05
0
hIgG
MOR07692
hIgG
MOR07692
hIgG
MOR07692
0 min
5 min
15 min

FIG. 13

[Fig. 14]



[Fig. 15]

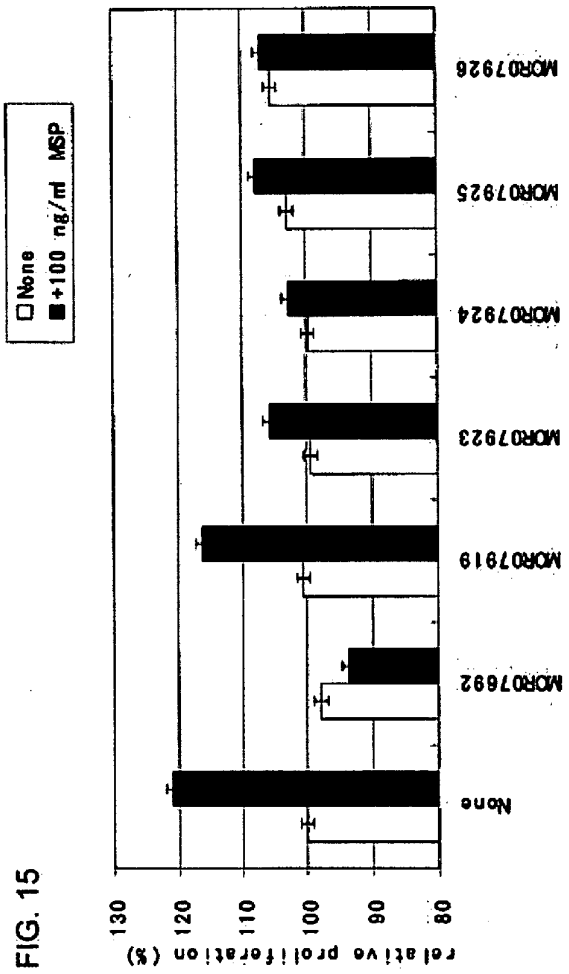

None
+100 ng/ml MSP
relative proliferation (%)
130
120
110
100
90
80
None
MOR07692
MOR07919
MOR07923
MOR07924
MOR07925
MOR07926

FIG. 15

[Fig. 16]

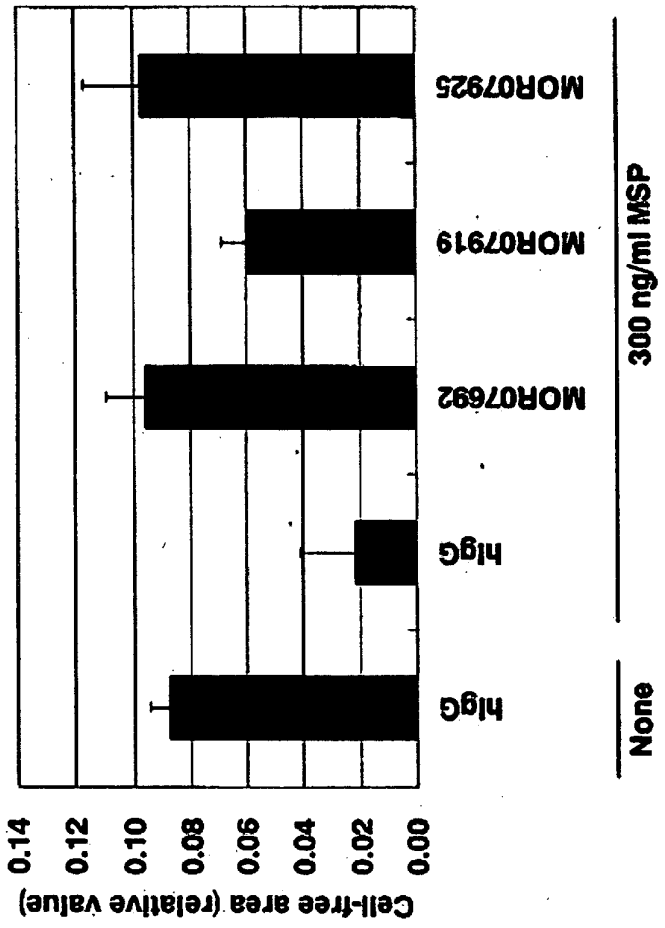

Cell-free area (relative value)
0.14
0.12
0.10
0.08
0.06
0.04
0.02
0.00
hIgG
hIgG
MOR07692
MOR07919
MOR07925
None
300 ng/ml MSP

FIG. 16

[Fig. 17]

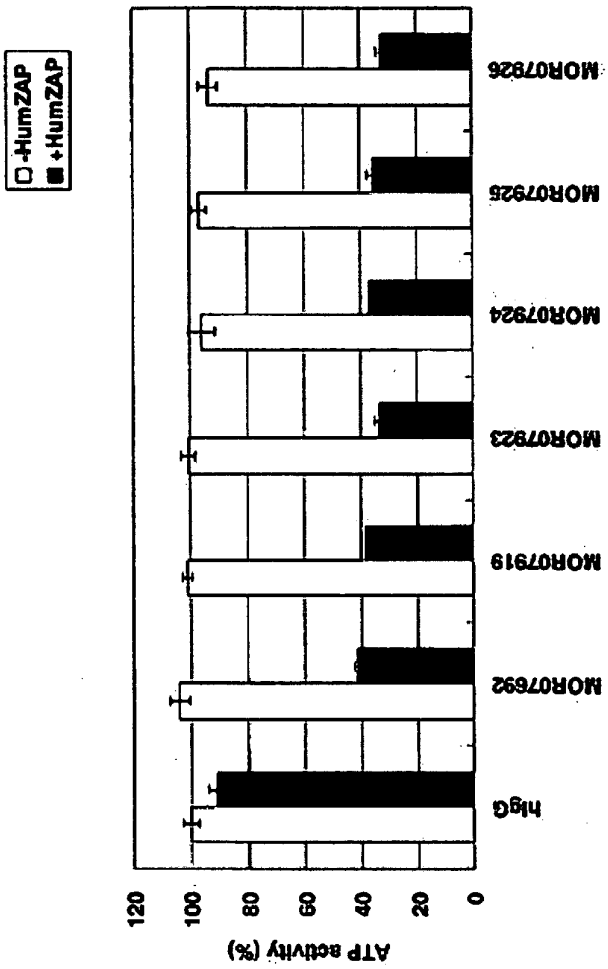

-HumZAP
+HumZAP
MOR07926
MOR07925
MOR07924
MOR07923
MOR07919
MOR07692
hIgG
120
100
80
60
40
20
0
ATP activity (%)

FIG. 17

SEQUENCE LISTING

20102143301 26 Sep 2011

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<120> ANTI-MST1R ANTIBODIES AND USES THEREOF

<130> DSPCT-FP1004

<150> US 61/151,411
<151> 2009-02-10

<160> 83

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Gly

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Asp Val Lys Tyr Val Val Pro Ser Phe Ser Ala Gly Gly Leu Val Gln
            20                  25                  30

Ala Met Val Thr Tyr Glu Gly Asp Arg Asn Glu Ser Ala Val Phe Val
        35                  40                  45

Ala Ile Arg Asn Arg Leu His Val Leu Gly Pro Asp Leu Lys Ser Val
    50                  55                  60

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

Ala Ala Cys Gly Pro Gly Pro His Gly Pro Pro Gly Asp Thr Asp Thr
                85                  90                  95

Lys Val Leu Val Leu Asp Pro Ala Leu Pro Ala Leu Val Ser Cys Gly
            100                 105                 110

Ser Ser Leu Gln Gly Arg Cys Phe Leu His Asp Leu Glu Pro Gln Gly
        115                 120                 125

Thr Ala Val His Leu Ala Ala Pro Ala Cys Leu Phe Ser Ala His His
    130                 135                 140

Asn Arg Pro Asp Asp Cys Pro Asp Cys Val Ala Ser Pro Leu Gly Thr
145                 150                 155                 160

Arg Val Thr Val Val Glu Gln Gly Gln Ala Ser Tyr Phe Tyr Val Ala
                165                 170                 175

Ser Ser Leu Asp Ala Ala Val Ala Gly Ser Phe Ser Pro Arg Ser Val
            180                 185                 190

Ser Ile Arg Arg Leu Lys Ala Asp Ala Ser Gly Phe Ala Pro Gly Phe
        195                 200                 205

Val Ala Leu Ser Val Leu Pro Lys His Leu Val Ser Tyr Ser Ile Glu
    210                 215                 220

Tyr Val His Ser Phe His Thr Gly Ala Phe Val Tyr Phe Leu Thr Val
225                 230                 235                 240

Gln Pro Ala Ser Val Thr Asp Asp Pro Ser Ala Leu His Thr Arg Leu
                245                 250                 255

Ala Arg Leu Ser Ala Thr Glu Pro Glu Leu Gly Asp Tyr Arg Glu Leu
            260                 265                 270
```

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Val Leu Asp Cys Arg Phe Ala Pro Lys Arg Arg Arg Gly Ala Pro
275 280 285

Glu Gly Gly Gln Pro Tyr Pro Val Leu Gln Val Ala His Ser Ala Pro
290 295 300

Val Gly Ala Gln Leu Ala Thr Glu Leu Ser Ile Ala Glu Gly Gln Glu
305 310 315 320

Val Leu Phe Gly Val Phe Val Thr Gly Lys Asp Gly Gly Pro Gly Val
325 330 335

Gly Pro Asn Ser Val Val Cys Ala Phe Pro Ile Asp Leu Leu Asp Thr
340 345 350

Leu Ile Asp Glu Gly Val Glu Arg Cys Cys Glu Ser Pro Val His Pro
355 360 365

Gly Leu Arg Arg Gly Leu Asp Phe Phe Gln Ser Pro Ser Phe Cys Pro
370 375 380

Asn Pro Pro Gly Leu Glu Ala Leu Ser Pro Asn Thr Ser Cys Arg His
385 390 395 400

Phe Pro Leu Leu Val Ser Ser Ser Phe Ser Arg Val Asp Leu Phe Asn
405 410 415

Gly Leu Leu Gly Pro Val Gln Val Thr Ala Leu Tyr Val Thr Arg Leu
420 425 430

Asp Asn Val Thr Val Ala His Met Gly Thr Met Asp Gly Arg Ile Leu
435 440 445

Gln Val Glu Leu Val Arg Ser Leu Asn Tyr Leu Leu Tyr Val Ser Asn
450 455 460

Phe Ser Leu Gly Asp Ser Gly Gln Pro Val Gln Arg Asp Val Ser Arg
465 470 475 480

Leu Gly Asp His Leu Leu Phe Ala Ser Gly Asp Gln Val Phe Gln Val
485 490 495

Pro Ile Arg Gly Pro Gly Cys Arg His Phe Leu Thr Cys Gly Arg Cys
500 505 510

Leu Arg Ala Trp His Phe Met Gly Cys Gly Trp Cys Gly Asn Met Cys
515 520 525

Gly Gln Gln Lys Glu Cys Pro Gly Ser Trp Gln Gln Asp His Cys Pro
530 535 540

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2010214301

```
Pro Lys Leu
545


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

agc ctg cgt ctg agc tgc gcg gcc tcc gga ttt acc ttt aat tct tat          96
Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Ser Tyr
            20                  25                  30

tct atg tct tgg gtg cgc caa gcc cct ggg aag ggt ctc gag tgg gtg         144
Ser Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val
        35                  40                  45

agc tat atc tct tct cgt tct agc act acc tat tat gcg gat agc gtg         192
Ser Tyr Ile Ser Ser Arg Ser Ser Thr Thr Tyr Tyr Ala Asp Ser Val
    50                  55                  60

aaa ggc cgt ttt acc att tca cgt gat aat tcg aaa aac acc ctg tat         240
Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr
65                  70                  75                  80

ctg caa atg aac agc ctg cgt gcg gaa gat acg gcc gtg tat tat tgc         288
Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys
                85                  90                  95

gcg cgt ggt tat ttt cat ggt atg gat tat tgg ggc caa ggc acc ctg         336
Ala Arg Gly Tyr Phe His Gly Met Asp Tyr Trp Gly Gln Gly Thr Leu
            100                 105                 110

gtg acg gtt agc tca                                                    351
Val Thr Val Ser Ser
        115


<210>  19
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<213>  Homo sapiens

<400>  19

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


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


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

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2010214301

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

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

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

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

Val Thr Val Ser Ser
        115
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```
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Gln Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys Pro Gly Glu
1               5                   10                  15

agc ctg aaa att agc tgc aaa ggt tcc gga tat tcc ttt act aat tat          96
Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe Thr Asn Tyr
            20                  25                  30

tgg att tct tgg gtg cgc cag atg cct ggg aag ggt ctc gag tgg atg         144
Trp Ile Ser Trp Val Arg Gln Met Pro Gly Lys Gly Leu Glu Trp Met
        35                  40                  45

ggc ttt atc tat ccg gat gat agc tat acc cgt tat tct ccg agc ttt         192
Gly Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser Pro Ser Phe
    50                  55                  60

cag ggc cag gtg acc att agc gcg gat aaa agc att agc acc gcg tat         240
Gln Gly Gln Val Thr Ile Ser Ala Asp Lys Ser Ile Ser Thr Ala Tyr
65                  70                  75                  80

ctt caa tgg agc agc ctg aaa gcg agc gat acg gcc atg tat tat tgc         288
Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met Tyr Tyr Cys
                85                  90                  95

gcg cgt ttt tct tat cgt cat tat ctt gat atg gat gat cat tgg ggc         336
Ala Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp His Trp Gly
            100                 105                 110

caa ggc acc ctg gtg acg gtt agc tca                                    363
Gln Gly Thr Leu Val Thr Val Ser Ser
        115                 120
```

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20102143O1

<213> Homo sapiens

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1               5                  10                  15

Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe Thr Asn Tyr
            20                  25                  30

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

Gly Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser Pro Ser Phe
    50                  55                  60

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

Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met Tyr Tyr Cys
                85                  90                  95

Ala Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp His Trp Gly
            100                 105                 110

Gln Gly Thr Leu Val Thr Val Ser Ser
        115                 120
```

<210> 22
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<213> Homo sapiens

<220>
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<222> (1)..(330)

<400> 22

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

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat          96
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta         144
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        35                  40                  45

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc         192
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    50                  55                  60

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa         240
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

cct gaa gac ttt gcg act tat tat tgc cag cag tat tat aat atg cct         288
```

```
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Tyr Asn Met Pro
                85                  90                  95

tat acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg              330
Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110


<210>  23
<211>  110
<212>  PRT
<213>  Homo sapiens

<400>  23

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

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

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Tyr Asn Met Pro
                85                  90                  95

Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110


<210>  24
<211>  330
<212>  DNA
<213>  Homo sapiens


<220>
<221>  CDS
<222>  (1)..(330)

<400>  24
gat atc gaa ctg acc cag ccg cct tca gtg agc gtt gca cca ggt cag       48
Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln
1               5                   10                  15

acc gcg cgt atc tcg tgt agc ggc gat tct ctt ggt tct aag tat gtt       96
Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser Lys Tyr Val
            20                  25                  30

cat tgg tac cag cag aaa ccc ggg cag gcg cca gtt ctt gtg att tat      144
His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr
        35                  40                  45

cgt gat aat aag cgt ccc tca ggc atc ccg gaa cgc ttt agc gga tcc      192
Arg Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
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    50                  55                  60

aac agc ggc aac acc gcg acc ctg acc att agc ggc act cag gcg gaa        240
Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
65                  70                  75                  80

gac gaa gcg gat tat tat tgc cag tct tat gat gct act gag ttt act        288
Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr Glu Phe Thr
                85                  90                  95

tat gtg ttt ggc ggc ggc acg aag tta acc gtt ctt ggc cag                330
Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
            100                 105                 110


<210>  25
<211>  110
<212>  PRT
<213>  Homo sapiens

<400>  25

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


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


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


Arg Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser
    50                  55                  60


Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
65                  70                  75                  80


Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr Glu Phe Thr
                85                  90                  95


Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
            100                 105                 110


<210>  26
<211>  330
<212>  DNA
<213>  Homo sapiens


<220>
<221>  CDS
<222>  (1)..(330)

<400>  26
gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc         48
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
1               5                   10                  15

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat         96
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30
```

```
tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta        144
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        35                  40                  45

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc        192
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    50                  55                  60

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa        240
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

cct gaa gac ttt gcg acc tat tat tgc ttt cag tat ctt att gtt cct        288
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr Leu Ile Val Pro
                85                  90                  95

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg                330
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110


<210>  27
<211>  110
<212>  PRT
<213>  Homo sapiens

<400>  27

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


Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30


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


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


Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80


Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr Leu Ile Val Pro
                85                  90                  95


Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110


<210>  28
<211>  330
<212>  DNA
<213>  Homo sapiens


<220>
<221>  CDS
<222>  (1)..(330)

<400>  28
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gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc          48
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
1               5                  10                  15

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat          96
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta         144
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        35                  40                  45

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc         192
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    50                  55                  60

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa         240
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

cct gaa gac ttt gcg acc tat tat tgc cag cag tat aat att aat cct         288
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ile Asn Pro
                85                  90                  95

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg                 330
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110
```

<210> 29
<211> 110
<212> PRT
<213> Homo sapiens

<400> 29

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

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

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ile Asn Pro
                85                  90                  95

Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110
```

<210> 30
<211> 330
<212> DNA
<213> Homo sapiens

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<220>
<221> CDS
<222> (1)..(330)

<400> 30

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gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc          48
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
1               5                  10                  15

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat          96
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta         144
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        35                  40                  45

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc         192
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    50                  55                  60

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa         240
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

cct gaa gac ttt gcg acc tat tat tgc ctt cag tat ttt aat cct cct         288
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr Phe Asn Pro Pro
                85                  90                  95

cat acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg                 330
His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110
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<210> 31
<211> 110
<212> PRT
<213> Homo sapiens

<400> 31

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

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

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr Phe Asn Pro Pro
                85                  90                  95

His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
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100 105 110

<210> 32
<211> 330
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (1)..(330)

<400> 32

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gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc          48
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
1               5                   10                  15

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat          96
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta         144
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        35                  40                  45

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc         192
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    50                  55                  60

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa         240
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80

cct gaa gac ttt gcg acc tat tat tgc ttt cag gct ctt att atg cct         288
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala Leu Ile Met Pro
                85                  90                  95

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg                 330
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
            100                 105                 110
```

<210> 33
<211> 110
<212> PRT
<213> Homo sapiens

<400> 33

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

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            20                  25                  30

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

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65                  70                  75                  80
```

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Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala Leu Ile Met Pro
85 90 95

Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
100 105 110

<210> 34
<211> 24
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR forward primer

<400> 34
ttggcccagt ccagacctcg agtc 24

<210> 35
<211> 25
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR reverse primer

<400> 35
actctgtgga gtgagggacc taatg 25

<210> 36
<211> 4200
<212> DNA
<213> Macaca fascicularis

<220>
<221> CDS
<222> (1)..(4200)

<400> 36
atg gag ctc ctc cct ccg ctg cct cag tcc ttc tta ctg ctg ctg ctg 48
Met Glu Leu Leu Pro Pro Leu Pro Gln Ser Phe Leu Leu Leu Leu Leu
1 5 10 15

ttg cct gcc aag ccc gcg gcg gcc aag gaa tgg cag tgc ccg cgc acc 96
Leu Pro Ala Lys Pro Ala Ala Ala Lys Glu Trp Gln Cys Pro Arg Thr
20 25 30

ccc tac gcg gcc tct cga gac ttt aac gtg aag tac atg gtg ccc agc 144
Pro Tyr Ala Ala Ser Arg Asp Phe Asn Val Lys Tyr Met Val Pro Ser
35 40 45

ttc tcc gcc gga ggc ctg gtg cag acc atg gtg acc tac cag ggc gac 192
Phe Ser Ala Gly Gly Leu Val Gln Thr Met Val Thr Tyr Gln Gly Asp
50 55 60

aaa aat gag agt gct gtg ttt gta gcc ata cgc aat cgc ctg cac gtg 240
Lys Asn Glu Ser Ala Val Phe Val Ala Ile Arg Asn Arg Leu His Val
65 70 75 80

ctt ggg cct gac ctg aag tct gtc cag agc ctg gcc acg ggc cct gct 288
Leu Gly Pro Asp Leu Lys Ser Val Gln Ser Leu Ala Thr Gly Pro Ala
85 90 95

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```
ggg gac cct ggc tgc cag acg tgt gca gcc tgt ggc cca ggc ccc cac          336
Gly Asp Pro Gly Cys Gln Thr Cys Ala Ala Cys Gly Pro Gly Pro His
            100                 105                 110

ggc cct tcc ggt gac aca gac aca aag gtg ctg gtg ctg gag ccc gcg          384
Gly Pro Ser Gly Asp Thr Asp Thr Lys Val Leu Val Leu Glu Pro Ala
        115                 120                 125

ctg cct gcc ctg gtc agt tgt ggc tcc agc ctg cag ggc cgc tgc ttc          432
Leu Pro Ala Leu Val Ser Cys Gly Ser Ser Leu Gln Gly Arg Cys Phe
    130                 135                 140

ctg cat gac cta gat ccc caa ggg aca gcc gtg cat ctg gca gcg cca          480
Leu His Asp Leu Asp Pro Gln Gly Thr Ala Val His Leu Ala Ala Pro
145                 150                 155                 160

gcc tgc ctc ttc tca gcc cac cat aac cgg ccc gat gac tgc ccc gac          528
Ala Cys Leu Phe Ser Ala His His Asn Arg Pro Asp Asp Cys Pro Asp
                165                 170                 175

tgt gtg gcc agc cca ttg ggc acc cgt gtg act gtg gtt gag caa ggc          576
Cys Val Ala Ser Pro Leu Gly Thr Arg Val Thr Val Val Glu Gln Gly
            180                 185                 190

cag gcc tcc tat ttc tac gtg gca tcc tca ctg gac gca gcc gtg gct          624
Gln Ala Ser Tyr Phe Tyr Val Ala Ser Ser Leu Asp Ala Ala Val Ala
        195                 200                 205

gcc agc ttc agc cca cgc tca gtg tct atc agg cgt ctc aag gcc gac          672
Ala Ser Phe Ser Pro Arg Ser Val Ser Ile Arg Arg Leu Lys Ala Asp
    210                 215                 220

gcc tcg gga ttt gca ccg ggc ttt gtg gca ttg tca gtg ctg ccc aag          720
Ala Ser Gly Phe Ala Pro Gly Phe Val Ala Leu Ser Val Leu Pro Lys
225                 230                 235                 240

cat ctt gtc tcc tac agt att gaa tat gtg cac agc ttc cat acg gga          768
His Leu Val Ser Tyr Ser Ile Glu Tyr Val His Ser Phe His Thr Gly
                245                 250                 255

gcc ttc gtc tac ttc ctg act gta cag ccg gcc agc gtg act gat gct          816
Ala Phe Val Tyr Phe Leu Thr Val Gln Pro Ala Ser Val Thr Asp Ala
            260                 265                 270

cct ggt gcc ctg cac aca cgc ctg gca cga ctt agc gcc act gag cca          864
Pro Gly Ala Leu His Thr Arg Leu Ala Arg Leu Ser Ala Thr Glu Pro
        275                 280                 285

gag ttg ggt gac tat cgg gag ctg gtc ctc gac tgc aga ttt gct cca          912
Glu Leu Gly Asp Tyr Arg Glu Leu Val Leu Asp Cys Arg Phe Ala Pro
    290                 295                 300

aaa cgc agg cgc cgg ggg gcc cca gag ggc gga cag ccc tac cct gtg          960
Lys Arg Arg Arg Arg Gly Ala Pro Glu Gly Gly Gln Pro Tyr Pro Val
305                 310                 315                 320

ctg cgg gtg gcc cac tct gct cca gtg ggt gct caa ctt gcc act gag         1008
Leu Arg Val Ala His Ser Ala Pro Val Gly Ala Gln Leu Ala Thr Glu
                325                 330                 335

ctg agc att gct gag ggc cag gaa gtg cta ttt ggg gtc ttt gtg gct         1056
Leu Ser Ile Ala Glu Gly Gln Glu Val Leu Phe Gly Val Phe Val Ala
            340                 345                 350

ggc aag gat agt ggc cct ggc gtg ggc ccc aac tct gtc gtc tgt gcc         1104
Gly Lys Asp Ser Gly Pro Gly Val Gly Pro Asn Ser Val Val Cys Ala
        355                 360                 365
```

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```
ttc ccc att gac ctg ctg gac aca tta att gat gaa ggt gtg gag cgc      1152
Phe Pro Ile Asp Leu Leu Asp Thr Leu Ile Asp Glu Gly Val Glu Arg
    370                 375                 380

tgt tgt gaa tcc cca gtc cat cca ggc ctc cgg cga ggc ctc gac ttc      1200
Cys Cys Glu Ser Pro Val His Pro Gly Leu Arg Arg Gly Leu Asp Phe
385                 390                 395                 400

ttc cag tca ccc agt ttt tgc ccc aac ccg cct ggc ctg gag gcc ccc      1248
Phe Gln Ser Pro Ser Phe Cys Pro Asn Pro Pro Gly Leu Glu Ala Pro
                405                 410                 415

agc ccc aac acc agc tgc cgc cac ttc cct ttg ctg gtc agt agc agc      1296
Ser Pro Asn Thr Ser Cys Arg His Phe Pro Leu Leu Val Ser Ser Ser
            420                 425                 430

ttc tca cgt gtg gac cta ttc aat ggg ctg ttg gga aca gta gag gtc      1344
Phe Ser Arg Val Asp Leu Phe Asn Gly Leu Leu Gly Thr Val Glu Val
        435                 440                 445

act gca ctg tat gtg aca cgc ctt gac aac gtc aca gtg gca cac atg      1392
Thr Ala Leu Tyr Val Thr Arg Leu Asp Asn Val Thr Val Ala His Met
    450                 455                 460

ggc aca gcg gat ggg cgt atc ctg cag gtg gag ctg gcc agg tca ctc      1440
Gly Thr Ala Asp Gly Arg Ile Leu Gln Val Glu Leu Ala Arg Ser Leu
465                 470                 475                 480

aac tac ttg ctg tat gtg tcc aac ttc tca ctg ggt gac agt ggg cag      1488
Asn Tyr Leu Leu Tyr Val Ser Asn Phe Ser Leu Gly Asp Ser Gly Gln
                485                 490                 495

ccc gtg cag cgg gat gtc agt cgc ctt ggg gac cac cta ttc ttc gcc      1536
Pro Val Gln Arg Asp Val Ser Arg Leu Gly Asp His Leu Phe Phe Ala
            500                 505                 510

tct ggg gac cag gtt ttc cag gta cct atc caa ggc cct ggc tgc cgc      1584
Ser Gly Asp Gln Val Phe Gln Val Pro Ile Gln Gly Pro Gly Cys Arg
        515                 520                 525

cac ttc ctc acc tgt ggg cgt tgc cta agg gca cag cgt ttc atg ggc      1632
His Phe Leu Thr Cys Gly Arg Cys Leu Arg Ala Gln Arg Phe Met Gly
    530                 535                 540

tgt ggc tgg tgt ggg aac atg tgt ggc cgg cag aag gag tgt cct ggc      1680
Cys Gly Trp Cys Gly Asn Met Cys Gly Arg Gln Lys Glu Cys Pro Gly
545                 550                 555                 560

tcc tgg caa cag gac cac tgt ccg cct aag ctt act gag ttc cac ccc      1728
Ser Trp Gln Gln Asp His Cys Pro Pro Lys Leu Thr Glu Phe His Pro
                565                 570                 575

cac agt gga cct tta agg ggc agt aca agg ctg acc ctg tgt ggc tcc      1776
His Ser Gly Pro Leu Arg Gly Ser Thr Arg Leu Thr Leu Cys Gly Ser
            580                 585                 590

aac ttc tac ctg cac cct tct ggt ctg gtg cct gag gga acc cat cag      1824
Asn Phe Tyr Leu His Pro Ser Gly Leu Val Pro Glu Gly Thr His Gln
        595                 600                 605

atc acg gtg ggc caa agt ccc tgc cgg cca ctg ccc aag gac agc tca      1872
Ile Thr Val Gly Gln Ser Pro Cys Arg Pro Leu Pro Lys Asp Ser Ser
    610                 615                 620

aaa ctc aga cca gtg ccc cgg aaa gac ttt gta gag gag ttt gag tgt      1920
Lys Leu Arg Pro Val Pro Arg Lys Asp Phe Val Glu Glu Phe Glu Cys
625                 630                 635                 640
```

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gaa ctg gag ccc ttg ggc acc caa gca gtg ggg cct acc aac gtc agc      1968
Glu Leu Glu Pro Leu Gly Thr Gln Ala Val Gly Pro Thr Asn Val Ser
                645                 650                 655

ctc acc gtg act aac atg cca ccg ggc aag cac ttc cgg gta gac ggc      2016
Leu Thr Val Thr Asn Met Pro Pro Gly Lys His Phe Arg Val Asp Gly
            660                 665                 670

acc tcc atg ctg aga ggc ttc ttt ttc atg gag cca gtg ctg ata gca      2064
Thr Ser Met Leu Arg Gly Phe Phe Phe Met Glu Pro Val Leu Ile Ala
        675                 680                 685

gtg caa ccc ctc ttt ggc cca cgg gca gga ggc acc tgt ctc act ctt      2112
Val Gln Pro Leu Phe Gly Pro Arg Ala Gly Gly Thr Cys Leu Thr Leu
    690                 695                 700

gaa ggc cag agt ctg tct gta ggc acc agc cgg gct gtg ctg gtc aat      2160
Glu Gly Gln Ser Leu Ser Val Gly Thr Ser Arg Ala Val Leu Val Asn
705                 710                 715                 720

ggg act gag tgt ctg cta gca cgg gtc agt gag ggg cag ctt tta tgt      2208
Gly Thr Glu Cys Leu Leu Ala Arg Val Ser Glu Gly Gln Leu Leu Cys
                725                 730                 735

gcc aca ccc cct ggg gcc atg gtg gcc agt gtc ccc ctt agc ctg cag      2256
Ala Thr Pro Pro Gly Ala Met Val Ala Ser Val Pro Leu Ser Leu Gln
            740                 745                 750

gtg ggg ggt gcc cag gta cct ggt tcc tgg acc ttc cac tac aga gaa      2304
Val Gly Gly Ala Gln Val Pro Gly Ser Trp Thr Phe His Tyr Arg Glu
        755                 760                 765

gac cct gtc gtg cta agc atc agc ccc aac tgt ggc tac agc aac tcc      2352
Asp Pro Val Val Leu Ser Ile Ser Pro Asn Cys Gly Tyr Ser Asn Ser
    770                 775                 780

cac atc acc atc tgt ggc cag cat cta act tca gca tgg cac tta gtg      2400
His Ile Thr Ile Cys Gly Gln His Leu Thr Ser Ala Trp His Leu Val
785                 790                 795                 800

ctg tca ttc cat gac ggg ctt agg gca gtg gag agc agg tgt gag agg      2448
Leu Ser Phe His Asp Gly Leu Arg Ala Val Glu Ser Arg Cys Glu Arg
                805                 810                 815

cag ctt cca gag cag cag ttg tgc cgc ctg cct gaa tat gtg gtc caa      2496
Gln Leu Pro Glu Gln Gln Leu Cys Arg Leu Pro Glu Tyr Val Val Gln
            820                 825                 830

gac ccc cag gga tgg gtg gca gga aat ctg agt gcc tgg ggg gat gga      2544
Asp Pro Gln Gly Trp Val Ala Gly Asn Leu Ser Ala Trp Gly Asp Gly
        835                 840                 845

gct gct ggc ttt aca ctg cct ggc ttt cgc ttc cta acc cca ccc cat      2592
Ala Ala Gly Phe Thr Leu Pro Gly Phe Arg Phe Leu Thr Pro Pro His
    850                 855                 860

cca ccc agt gcc aac cta att cca ctg aag cct gag gag cat gcc att      2640
Pro Pro Ser Ala Asn Leu Ile Pro Leu Lys Pro Glu Glu His Ala Ile
865                 870                 875                 880

aag ttt gag tat att ggg ctg ggt gct gtg act gac tgc gtg ggt gtc      2688
Lys Phe Glu Tyr Ile Gly Leu Gly Ala Val Thr Asp Cys Val Gly Val
                885                 890                 895

aac gtg acc gtg ggt ggt gag agc tgc cag cac gag ttc cgg ggg gac      2736
Asn Val Thr Val Gly Gly Glu Ser Cys Gln His Glu Phe Arg Gly Asp
            900                 905                 910
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atg gtt gtc tgc ccc ctg ccc cca tcc ctg cag ctt ggc aag gat ggt       2784
Met Val Val Cys Pro Leu Pro Pro Ser Leu Gln Leu Gly Lys Asp Gly
        915                 920                 925

gcc cca ctg cag gtc tgc gtg gat ggt gaa tgt cac atc ctg ggt aga       2832
Ala Pro Leu Gln Val Cys Val Asp Gly Glu Cys His Ile Leu Gly Arg
    930                 935                 940

gtg gtg tgg cca ggg cca gat ggg gtc cca cag agc acg ctc ctt ggt       2880
Val Val Trp Pro Gly Pro Asp Gly Val Pro Gln Ser Thr Leu Leu Gly
945                 950                 955                 960

atc ctg ctg cct ttg ctg ctg ctt gtg gcc gca ttg gcc act gca ctg       2928
Ile Leu Leu Pro Leu Leu Leu Leu Val Ala Ala Leu Ala Thr Ala Leu
                965                 970                 975

gtc ttc agc tac tgg tgg cag agg aag cag cta gtt ctt cct ccc aac       2976
Val Phe Ser Tyr Trp Trp Gln Arg Lys Gln Leu Val Leu Pro Pro Asn
            980                 985                 990

ctg gat gac ctg gca tcc ctg gac  cag act act gga gcc  aca ccc ctg     3024
Leu Asp Asp Leu Ala Ser Leu Asp  Gln Thr Thr Gly Ala  Thr Pro Leu
        995                 1000                1005

cct att  ctc tac tcg ggc tct  gac tac aga agt ggc  ctt gca cgc        3069
Pro Ile  Leu Tyr Ser Gly Ser  Asp Tyr Arg Ser Gly  Leu Ala Arg
    1010                1015                1020

cct gcc  act gat ggt cta gat  tcc act tgt gtc cat  gga gca tcc        3114
Pro Ala  Thr Asp Gly Leu Asp  Ser Thr Cys Val His  Gly Ala Ser
    1025                1030                1035

ttc tcc  aat agt gaa gat gaa  tcc tgt gtt cca ctg  ctg cgg aaa        3159
Phe Ser  Asn Ser Glu Asp Glu  Ser Cys Val Pro Leu  Leu Arg Lys
    1040                1045                1050

gag tcc  atc cag cta agg gac  ctg gac tct gcg ctg  ttg gct gag        3204
Glu Ser  Ile Gln Leu Arg Asp  Leu Asp Ser Ala Leu  Leu Ala Glu
    1055                1060                1065

gtc aag  gat gtg ctg att ccc  cat gag cgg gtg gtc  gcc cac agt        3249
Val Lys  Asp Val Leu Ile Pro  His Glu Arg Val Val  Ala His Ser
    1070                1075                1080

gac cga  gtc att ggc aaa ggc  cac ttt gga gtt gtc  tac cat gga        3294
Asp Arg  Val Ile Gly Lys Gly  His Phe Gly Val Val  Tyr His Gly
    1085                1090                1095

gaa tac  ata gac cag gcc cag  aat cga atc caa tgt  gcc atc aag        3339
Glu Tyr  Ile Asp Gln Ala Gln  Asn Arg Ile Gln Cys  Ala Ile Lys
    1100                1105                1110

tca cta  agt cgc atc aca gag  atg cag cag gtg gag  gcc ttc ctt        3384
Ser Leu  Ser Arg Ile Thr Glu  Met Gln Gln Val Glu  Ala Phe Leu
    1115                1120                1125

cga gag  ggg ctg ctc atg cgt  ggc ctg aac cac ccg  aat gtg ctg        3429
Arg Glu  Gly Leu Leu Met Arg  Gly Leu Asn His Pro  Asn Val Leu
    1130                1135                1140

gct ctc  att ggt atc atg ttg  cca ccc gag ggc ctg  ccc cat gtg        3474
Ala Leu  Ile Gly Ile Met Leu  Pro Pro Glu Gly Leu  Pro His Val
    1145                1150                1155

ctg ctg  ccc tat atg tgc cac  ggt gac ctg ctc cag  ttc atc cgc        3519
Leu Leu  Pro Tyr Met Cys His  Gly Asp Leu Leu Gln  Phe Ile Arg
    1160                1165                1170
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tca cct  cag cgg aac ccc acc  gtg aag gac ctc atc  agc ttt ggc        3564
Ser Pro  Gln Arg Asn Pro Thr  Val Lys Asp Leu Ile  Ser Phe Gly
    1175                 1180                 1185

ctg cag  gta gcc cat ggc atg  gag tac ctc gca gag  cag aag ttt        3609
Leu Gln  Val Ala His Gly Met  Glu Tyr Leu Ala Glu  Gln Lys Phe
    1190                 1195                 1200

gtg cac  agg gac ctg gct gcg  cga aac tgc atg ctg  gac gag tca        3654
Val His  Arg Asp Leu Ala Ala  Arg Asn Cys Met Leu  Asp Glu Ser
    1205                 1210                 1215

ttc act  gtc aaa gtg gct gac  ttt ggt ttg gcc cgt  gac atc ctg        3699
Phe Thr  Val Lys Val Ala Asp  Phe Gly Leu Ala Arg  Asp Ile Leu
    1220                 1225                 1230

gac aag  gaa tac tat agt gtt  caa cag cat cgc cac  gct cgc cta        3744
Asp Lys  Glu Tyr Tyr Ser Val  Gln Gln His Arg His  Ala Arg Leu
    1235                 1240                 1245

cct gtg  aag tgg atg gcg ctg  gag agc ctg cag acc  tat aga ttt        3789
Pro Val  Lys Trp Met Ala Leu  Glu Ser Leu Gln Thr  Tyr Arg Phe
    1250                 1255                 1260

acc acc  aag tct gat gtg tgg  tca ttt ggt gtg ctg  ctg tgg gaa        3834
Thr Thr  Lys Ser Asp Val Trp  Ser Phe Gly Val Leu  Leu Trp Glu
    1265                 1270                 1275

ctg ttg  aca cgg ggt gcc cca  cca tac ccc cac atc  gac cct ttt        3879
Leu Leu  Thr Arg Gly Ala Pro  Pro Tyr Pro His Ile  Asp Pro Phe
    1280                 1285                 1290

gac ctc  acc cac ttc ctg gcc  cag ggt cgg cgc ctg  ccc cag cct        3924
Asp Leu  Thr His Phe Leu Ala  Gln Gly Arg Arg Leu  Pro Gln Pro
    1295                 1300                 1305

gag tat  tgc ccc aat tct ctg  tac caa gtg atg cag  caa tgc tgg        3969
Glu Tyr  Cys Pro Asn Ser Leu  Tyr Gln Val Met Gln  Gln Cys Trp
    1310                 1315                 1320

gag gcg  gac cca gca gca cga  ccc acc ttc gga gta  cta gtg ggg        4014
Glu Ala  Asp Pro Ala Ala Arg  Pro Thr Phe Gly Val  Leu Val Gly
    1325                 1330                 1335

gaa gtg  gag cag ata gtg tct  gca ctg ctt ggg gac  cat tat gtg        4059
Glu Val  Glu Gln Ile Val Ser  Ala Leu Leu Gly Asp  His Tyr Val
    1340                 1345                 1350

cag ctg  cca gca acc tac atg  aac ctg ggc ccc agc  acc tca cat        4104
Gln Leu  Pro Ala Thr Tyr Met  Asn Leu Gly Pro Ser  Thr Ser His
    1355                 1360                 1365

gag atg  aat gtg cat cca gaa  cag cag cag tcc tca  ccc atg cca        4149
Glu Met  Asn Val His Pro Glu  Gln Gln Gln Ser Ser  Pro Met Pro
    1370                 1375                 1380

ggg agt  gca cac cga ccc cgg  cca ctc tca gag cct  cct cgg ccc        4194
Gly Ser  Ala His Arg Pro Arg  Pro Leu Ser Glu Pro  Pro Arg Pro
    1385                 1390                 1395

act tga                                                             4200
Thr
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<210> 37
<211> 1399

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<212> PRT
<213> Macaca fascicularis

<400> 37

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Met Glu Leu Leu Pro Pro Leu Pro Gln Ser Phe Leu Leu Leu Leu Leu
1               5                  10                  15

Leu Pro Ala Lys Pro Ala Ala Ala Lys Glu Trp Gln Cys Pro Arg Thr
            20                  25                  30

Pro Tyr Ala Ala Ser Arg Asp Phe Asn Val Lys Tyr Met Val Pro Ser
        35                  40                  45

Phe Ser Ala Gly Gly Leu Val Gln Thr Met Val Thr Tyr Gln Gly Asp
    50                  55                  60

Lys Asn Glu Ser Ala Val Phe Val Ala Ile Arg Asn Arg Leu His Val
65                  70                  75                  80

Leu Gly Pro Asp Leu Lys Ser Val Gln Ser Leu Ala Thr Gly Pro Ala
                85                  90                  95

Gly Asp Pro Gly Cys Gln Thr Cys Ala Ala Cys Gly Pro Gly Pro His
            100                 105                 110

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

Leu Pro Ala Leu Val Ser Cys Gly Ser Ser Leu Gln Gly Arg Cys Phe
    130                 135                 140

Leu His Asp Leu Asp Pro Gln Gly Thr Ala Val His Leu Ala Ala Pro
145                 150                 155                 160

Ala Cys Leu Phe Ser Ala His His Asn Arg Pro Asp Asp Cys Pro Asp
                165                 170                 175

Cys Val Ala Ser Pro Leu Gly Thr Arg Val Thr Val Val Glu Gln Gly
            180                 185                 190

Gln Ala Ser Tyr Phe Tyr Val Ala Ser Ser Leu Asp Ala Ala Val Ala
        195                 200                 205

Ala Ser Phe Ser Pro Arg Ser Val Ser Ile Arg Arg Leu Lys Ala Asp
    210                 215                 220

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

His Leu Val Ser Tyr Ser Ile Glu Tyr Val His Ser Phe His Thr Gly
                245                 250                 255
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```
Ala Phe Val Tyr Phe Leu Thr Val Gln Pro Ala Ser Val Thr Asp Ala
            260                 265                 270

Pro Gly Ala Leu His Thr Arg Leu Ala Arg Leu Ser Ala Thr Glu Pro
        275                 280                 285

Glu Leu Gly Asp Tyr Arg Glu Leu Val Leu Asp Cys Arg Phe Ala Pro
    290                 295                 300

Lys Arg Arg Arg Arg Gly Ala Pro Glu Gly Gly Gln Pro Tyr Pro Val
305                 310                 315                 320

Leu Arg Val Ala His Ser Ala Pro Val Gly Ala Gln Leu Ala Thr Glu
                325                 330                 335

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

Gly Lys Asp Ser Gly Pro Gly Val Gly Pro Asn Ser Val Val Cys Ala
        355                 360                 365

Phe Pro Ile Asp Leu Leu Asp Thr Leu Ile Asp Glu Gly Val Glu Arg
    370                 375                 380

Cys Cys Glu Ser Pro Val His Pro Gly Leu Arg Arg Gly Leu Asp Phe
385                 390                 395                 400

Phe Gln Ser Pro Ser Phe Cys Pro Asn Pro Pro Gly Leu Glu Ala Pro
                405                 410                 415

Ser Pro Asn Thr Ser Cys Arg His Phe Pro Leu Leu Val Ser Ser Ser
            420                 425                 430

Phe Ser Arg Val Asp Leu Phe Asn Gly Leu Leu Gly Thr Val Glu Val
        435                 440                 445

Thr Ala Leu Tyr Val Thr Arg Leu Asp Asn Val Thr Val Ala His Met
    450                 455                 460

Gly Thr Ala Asp Gly Arg Ile Leu Gln Val Glu Leu Ala Arg Ser Leu
465                 470                 475                 480

Asn Tyr Leu Leu Tyr Val Ser Asn Phe Ser Leu Gly Asp Ser Gly Gln
                485                 490                 495

Pro Val Gln Arg Asp Val Ser Arg Leu Gly Asp His Leu Phe Phe Ala
            500                 505                 510

Ser Gly Asp Gln Val Phe Gln Val Pro Ile Gln Gly Pro Gly Cys Arg
        515                 520                 525
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```
His Phe Leu Thr Cys Gly Arg Cys Leu Arg Ala Gln Arg Phe Met Gly
    530                 535                 540

Cys Gly Trp Cys Gly Asn Met Cys Gly Arg Gln Lys Glu Cys Pro Gly
545                 550                 555                 560

Ser Trp Gln Gln Asp His Cys Pro Pro Lys Leu Thr Glu Phe His Pro
                565                 570                 575

His Ser Gly Pro Leu Arg Gly Ser Thr Arg Leu Thr Leu Cys Gly Ser
            580                 585                 590

Asn Phe Tyr Leu His Pro Ser Gly Leu Val Pro Glu Gly Thr His Gln
        595                 600                 605

Ile Thr Val Gly Gln Ser Pro Cys Arg Pro Leu Pro Lys Asp Ser Ser
    610                 615                 620

Lys Leu Arg Pro Val Pro Arg Lys Asp Phe Val Glu Glu Phe Glu Cys
625                 630                 635                 640

Glu Leu Glu Pro Leu Gly Thr Gln Ala Val Gly Pro Thr Asn Val Ser
                645                 650                 655

Leu Thr Val Thr Asn Met Pro Pro Gly Lys His Phe Arg Val Asp Gly
            660                 665                 670

Thr Ser Met Leu Arg Gly Phe Phe Phe Met Glu Pro Val Leu Ile Ala
        675                 680                 685

Val Gln Pro Leu Phe Gly Pro Arg Ala Gly Gly Thr Cys Leu Thr Leu
    690                 695                 700

Glu Gly Gln Ser Leu Ser Val Gly Thr Ser Arg Ala Val Leu Val Asn
705                 710                 715                 720

Gly Thr Glu Cys Leu Leu Ala Arg Val Ser Glu Gly Gln Leu Leu Cys
                725                 730                 735

Ala Thr Pro Pro Gly Ala Met Val Ala Ser Val Pro Leu Ser Leu Gln
            740                 745                 750

Val Gly Gly Ala Gln Val Pro Gly Ser Trp Thr Phe His Tyr Arg Glu
        755                 760                 765

Asp Pro Val Val Leu Ser Ile Ser Pro Asn Cys Gly Tyr Ser Asn Ser
    770                 775                 780

His Ile Thr Ile Cys Gly Gln His Leu Thr Ser Ala Trp His Leu Val
785                 790                 795                 800
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Leu Ser Phe His Asp Gly Leu Arg Ala Val Glu Ser Arg Cys Glu Arg
805 810 815

Gln Leu Pro Glu Gln Gln Leu Cys Arg Leu Pro Glu Tyr Val Val Gln
820 825 830

Asp Pro Gln Gly Trp Val Ala Gly Asn Leu Ser Ala Trp Gly Asp Gly
835 840 845

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

Pro Pro Ser Ala Asn Leu Ile Pro Leu Lys Pro Glu Glu His Ala Ile
865 870 875 880

Lys Phe Glu Tyr Ile Gly Leu Gly Ala Val Thr Asp Cys Val Gly Val
885 890 895

Asn Val Thr Val Gly Gly Glu Ser Cys Gln His Glu Phe Arg Gly Asp
900 905 910

Met Val Val Cys Pro Leu Pro Pro Ser Leu Gln Leu Gly Lys Asp Gly
915 920 925

Ala Pro Leu Gln Val Cys Val Asp Gly Glu Cys His Ile Leu Gly Arg
930 935 940

Val Val Trp Pro Gly Pro Asp Gly Val Pro Gln Ser Thr Leu Leu Gly
945 950 955 960

Ile Leu Leu Pro Leu Leu Leu Leu Val Ala Ala Leu Ala Thr Ala Leu
965 970 975

Val Phe Ser Tyr Trp Trp Gln Arg Lys Gln Leu Val Leu Pro Pro Asn
980 985 990

Leu Asp Asp Leu Ala Ser Leu Asp Gln Thr Thr Gly Ala Thr Pro Leu
995 1000 1005

Pro Ile Leu Tyr Ser Gly Ser Asp Tyr Arg Ser Gly Leu Ala Arg
1010 1015 1020

Pro Ala Thr Asp Gly Leu Asp Ser Thr Cys Val His Gly Ala Ser
1025 1030 1035

Phe Ser Asn Ser Glu Asp Glu Ser Cys Val Pro Leu Leu Arg Lys
1040 1045 1050

Glu Ser Ile Gln Leu Arg Asp Leu Asp Ser Ala Leu Leu Ala Glu
1055 1060 1065

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Val Lys Asp Val Leu Ile Pro His Glu Arg Val Val Ala His Ser
1070 1075 1080

Asp Arg Val Ile Gly Lys Gly His Phe Gly Val Val Tyr His Gly
1085 1090 1095

Glu Tyr Ile Asp Gln Ala Gln Asn Arg Ile Gln Cys Ala Ile Lys
1100 1105 1110

Ser Leu Ser Arg Ile Thr Glu Met Gln Gln Val Glu Ala Phe Leu
1115 1120 1125

Arg Glu Gly Leu Leu Met Arg Gly Leu Asn His Pro Asn Val Leu
1130 1135 1140

Ala Leu Ile Gly Ile Met Leu Pro Pro Glu Gly Leu Pro His Val
1145 1150 1155

Leu Leu Pro Tyr Met Cys His Gly Asp Leu Leu Gln Phe Ile Arg
1160 1165 1170

Ser Pro Gln Arg Asn Pro Thr Val Lys Asp Leu Ile Ser Phe Gly
1175 1180 1185

Leu Gln Val Ala His Gly Met Glu Tyr Leu Ala Glu Gln Lys Phe
1190 1195 1200

Val His Arg Asp Leu Ala Ala Arg Asn Cys Met Leu Asp Glu Ser
1205 1210 1215

Phe Thr Val Lys Val Ala Asp Phe Gly Leu Ala Arg Asp Ile Leu
1220 1225 1230

Asp Lys Glu Tyr Tyr Ser Val Gln Gln His Arg His Ala Arg Leu
1235 1240 1245

Pro Val Lys Trp Met Ala Leu Glu Ser Leu Gln Thr Tyr Arg Phe
1250 1255 1260

Thr Thr Lys Ser Asp Val Trp Ser Phe Gly Val Leu Leu Trp Glu
1265 1270 1275

Leu Leu Thr Arg Gly Ala Pro Pro Tyr Pro His Ile Asp Pro Phe
1280 1285 1290

Asp Leu Thr His Phe Leu Ala Gln Gly Arg Arg Leu Pro Gln Pro
1295 1300 1305

Glu Tyr Cys Pro Asn Ser Leu Tyr Gln Val Met Gln Gln Cys Trp
1310 1315 1320

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Glu Ala Asp Pro Ala Ala Arg Pro Thr Phe Gly Val Leu Val Gly
1325 1330 1335

Glu Val Glu Gln Ile Val Ser Ala Leu Leu Gly Asp His Tyr Val
1340 1345 1350

Gln Leu Pro Ala Thr Tyr Met Asn Leu Gly Pro Ser Thr Ser His
1355 1360 1365

Glu Met Asn Val His Pro Glu Gln Gln Gln Ser Ser Pro Met Pro
1370 1375 1380

Gly Ser Ala His Arg Pro Arg Pro Leu Ser Glu Pro Pro Arg Pro
1385 1390 1395

Thr

<210> 38
<211> 23
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR forward primer for human MST1R

<400> 38
gaggactggc agtgcccgcg cac 23

<210> 39
<211> 23
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR reverse primer for human MST1R

<400> 39
tcaagtgggc cgaggaggct ctg 23

<210> 40
<211> 24
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR forward primer for cynomolgus monkey MST1R

<400> 40
aaggaatggc agtgcccgcg cacc 24

<210> 41
<211> 25
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR forward primer for mouse MST1R

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<400> 41
tcaagtgggc cgaggaggct ctgag 25

<210> 42
<211> 23
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR forward primer for mouse MST1R

<400> 42
tccaccaacc tgaactggca gtg 23

<210> 43
<211> 23
<212> DNA
<213> Artificial Sequence

<220>
<223> PCR reverse primer for mouse MST1R

<400> 43
tcaagtgggc aggggtggct ctg 23

<210> 44
<211> 4785
<212> DNA
<213> Homo sapiens

<220>
<221> CDS
<222> (265)..(4467)

<400> 44
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gcaggcgtca ggtgctcaga cccgagggcc gggaagggat ttgggtttca caggaacctg 120

gggcgggggt ccgctatctt ggggctgtcg ggaccgctgc ttaaatttgg cccagtccag 180

acctcgagtc gggcccccag ccaggcccac gcccaggtcc aggcccaggc cggtagggat 240

cctctagggt cccagctcgc ctcg atg gag ctc ctc ccg ccg ctg cct cag 291
Met Glu Leu Leu Pro Pro Leu Pro Gln
1 5

tcc ttc ctg ttg ctg ctg ctg ttg cct gcc aag ccc gcg gcg ggc gag 339
Ser Phe Leu Leu Leu Leu Leu Leu Pro Ala Lys Pro Ala Ala Gly Glu
10 15 20 25

gac tgg cag tgc ccg cgc acc ccc tac gcg gcc tct cgc gac ttt gac 387
Asp Trp Gln Cys Pro Arg Thr Pro Tyr Ala Ala Ser Arg Asp Phe Asp
30 35 40

gtg aag tac gtg gtg ccc agc ttc tcc gcc gga ggc ctg gta cag gcc 435
Val Lys Tyr Val Val Pro Ser Phe Ser Ala Gly Gly Leu Val Gln Ala
45 50 55

atg gtg acc tac gag ggc gac aga aat gag agt gct gtg ttt gta gcc 483
Met Val Thr Tyr Glu Gly Asp Arg Asn Glu Ser Ala Val Phe Val Ala
60 65 70

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ata cgc aat cgc ctg cat gtg ctt ggg cct gac ctg aag tct gtc cag          531
Ile Arg Asn Arg Leu His Val Leu Gly Pro Asp Leu Lys Ser Val Gln
    75                  80                  85

agc ctg gcc acg ggc cct gct gga gac cct ggc tgc cag acg tgt gca          579
Ser Leu Ala Thr Gly Pro Ala Gly Asp Pro Gly Cys Gln Thr Cys Ala
90                  95                  100                 105

gcc tgt ggc cca gga ccc cac ggc cct ccc ggt gac aca gac aca aag          627
Ala Cys Gly Pro Gly Pro His Gly Pro Pro Gly Asp Thr Asp Thr Lys
                110                 115                 120

gtg ctg gtg ctg gat ccc gcg ctg cct gcg ctg gtc agt tgt ggc tcc          675
Val Leu Val Leu Asp Pro Ala Leu Pro Ala Leu Val Ser Cys Gly Ser
            125                 130                 135

agc ctg cag ggc cgc tgc ttc ctg cat gac cta gag ccc caa ggg aca          723
Ser Leu Gln Gly Arg Cys Phe Leu His Asp Leu Glu Pro Gln Gly Thr
        140                 145                 150

gcc gtg cat ctg gca gcg cca gcc tgc ctc ttc tca gcc cac cat aac          771
Ala Val His Leu Ala Ala Pro Ala Cys Leu Phe Ser Ala His His Asn
    155                 160                 165

cgg ccc gat gac tgc ccc gac tgt gtg gcc agc cca ttg ggc acc cgt          819
Arg Pro Asp Asp Cys Pro Asp Cys Val Ala Ser Pro Leu Gly Thr Arg
170                 175                 180                 185

gta act gtg gtt gag caa ggc cag gcc tcc tat ttc tac gtg gca tcc          867
Val Thr Val Val Glu Gln Gly Gln Ala Ser Tyr Phe Tyr Val Ala Ser
                190                 195                 200

tca ctg gac gca gcc gtg gct gcc agc ttc agc cca cgc tca gtg tct          915
Ser Leu Asp Ala Ala Val Ala Ala Ser Phe Ser Pro Arg Ser Val Ser
            205                 210                 215

atc agg cgt ctc aag gct gac gcc tcg gga ttc gca ccg ggc ttt gtg          963
Ile Arg Arg Leu Lys Ala Asp Ala Ser Gly Phe Ala Pro Gly Phe Val
        220                 225                 230

gcg ttg tca gtg ctg ccc aag cat ctt gtc tcc tac agt att gaa tac         1011
Ala Leu Ser Val Leu Pro Lys His Leu Val Ser Tyr Ser Ile Glu Tyr
    235                 240                 245

gtg cac agc ttc cac acg gga gcc ttc gta tac ttc ctg act gta cag         1059
Val His Ser Phe His Thr Gly Ala Phe Val Tyr Phe Leu Thr Val Gln
250                 255                 260                 265

ccg gcc agc gtg aca gat gat cct agt gcc ctg cac aca cgc ctg gca         1107
Pro Ala Ser Val Thr Asp Asp Pro Ser Ala Leu His Thr Arg Leu Ala
                270                 275                 280

cgg ctt agc gcc act gag cca gag ttg ggt gac tat cgg gag ctg gtc         1155
Arg Leu Ser Ala Thr Glu Pro Glu Leu Gly Asp Tyr Arg Glu Leu Val
            285                 290                 295

ctc gac tgc aga ttt gct cca aaa cgc agg cgc cgg ggg gcc cca gaa         1203
Leu Asp Cys Arg Phe Ala Pro Lys Arg Arg Arg Arg Gly Ala Pro Glu
        300                 305                 310

ggc gga cag ccc tac cct gtg ctg cgg gtg gcc cac tcc gct cca gtg         1251
Gly Gly Gln Pro Tyr Pro Val Leu Arg Val Ala His Ser Ala Pro Val
    315                 320                 325

ggt gcc caa ctt gcc act gag ctg agc atc gcc gag ggc cag gaa gta         1299
Gly Ala Gln Leu Ala Thr Glu Leu Ser Ile Ala Glu Gly Gln Glu Val
330                 335                 340                 345
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cta ttt ggg gtc ttt gtg act ggc aag gat ggt ggt cct ggc gtg ggc      1347
Leu Phe Gly Val Phe Val Thr Gly Lys Asp Gly Gly Pro Gly Val Gly
                350                 355                 360

ccc aac tct gtc gtc tgt gcc ttc ccc att gac ctg ctg gac aca cta      1395
Pro Asn Ser Val Val Cys Ala Phe Pro Ile Asp Leu Leu Asp Thr Leu
            365                 370                 375

att gat gag ggt gtg gag cgc tgt tgt gaa tcc cca gtc cat cca ggc      1443
Ile Asp Glu Gly Val Glu Arg Cys Cys Glu Ser Pro Val His Pro Gly
        380                 385                 390

ctc cgg cga ggc ctc gac ttc ttc cag tcg ccc agt ttt tgc ccc aac      1491
Leu Arg Arg Gly Leu Asp Phe Phe Gln Ser Pro Ser Phe Cys Pro Asn
    395                 400                 405

ccg cct ggc ctg gaa gcc ctc agc ccc aac acc agc tgc cgc cac ttc      1539
Pro Pro Gly Leu Glu Ala Leu Ser Pro Asn Thr Ser Cys Arg His Phe
410                 415                 420                 425

cct ctg ctg gtc agt agc agc ttc tca cgt gtg gac cta ttc aat ggg      1587
Pro Leu Leu Val Ser Ser Ser Phe Ser Arg Val Asp Leu Phe Asn Gly
                430                 435                 440

ctg ttg gga cca gta cag gtc act gca ttg tat gtg aca cgc ctt gac      1635
Leu Leu Gly Pro Val Gln Val Thr Ala Leu Tyr Val Thr Arg Leu Asp
            445                 450                 455

aac gtc aca gtg gca cac atg ggc aca atg gat ggg cgt atc ctg cag      1683
Asn Val Thr Val Ala His Met Gly Thr Met Asp Gly Arg Ile Leu Gln
        460                 465                 470

gtg gag ctg gtc agg tca cta aac tac ttg ctg tat gtg tcc aac ttc      1731
Val Glu Leu Val Arg Ser Leu Asn Tyr Leu Leu Tyr Val Ser Asn Phe
    475                 480                 485

tca ctg ggt gac agt ggg cag ccc gtg cag cgg gat gtc agt cgt ctt      1779
Ser Leu Gly Asp Ser Gly Gln Pro Val Gln Arg Asp Val Ser Arg Leu
490                 495                 500                 505

ggg gac cac cta ctc ttt gcc tct ggg gac cag gtt ttc cag gta cct      1827
Gly Asp His Leu Leu Phe Ala Ser Gly Asp Gln Val Phe Gln Val Pro
                510                 515                 520

atc caa ggc cct ggc tgc cgc cac ttc ctg acc tgt ggg cgt tgc cta      1875
Ile Gln Gly Pro Gly Cys Arg His Phe Leu Thr Cys Gly Arg Cys Leu
            525                 530                 535

agg gca tgg cat ttc atg ggc tgt ggc tgg tgt ggg aac atg tgc ggc      1923
Arg Ala Trp His Phe Met Gly Cys Gly Trp Cys Gly Asn Met Cys Gly
        540                 545                 550

cag cag aag gag tgt cct ggc tcc tgg caa cag gac cac tgc cca cct      1971
Gln Gln Lys Glu Cys Pro Gly Ser Trp Gln Gln Asp His Cys Pro Pro
    555                 560                 565

aag ctt act gag ttc cac ccc cac agt gga cct cta agg ggc agt aca      2019
Lys Leu Thr Glu Phe His Pro His Ser Gly Pro Leu Arg Gly Ser Thr
570                 575                 580                 585

agg ctg acc ctg tgt ggc tcc aac ttc tac ctt cac cct tct ggt ctg      2067
Arg Leu Thr Leu Cys Gly Ser Asn Phe Tyr Leu His Pro Ser Gly Leu
                590                 595                 600

gtg cct gag gga acc cat cag gtc act gtg ggc caa agt ccc tgc cgg      2115
Val Pro Glu Gly Thr His Gln Val Thr Val Gly Gln Ser Pro Cys Arg
            605                 610                 615
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cca ctg ccc aag gac agc tca aaa ctc aga cca gtg ccc cgg aaa gac      2163
Pro Leu Pro Lys Asp Ser Ser Lys Leu Arg Pro Val Pro Arg Lys Asp
        620                 625                 630

ttt gta gag gag ttt gag tgt gaa ctg gag ccc ttg ggc acc cag gca      2211
Phe Val Glu Glu Phe Glu Cys Glu Leu Glu Pro Leu Gly Thr Gln Ala
    635                 640                 645

gtg ggg cct acc aac gtc agc ctc acc gtg act aac atg cca ccg ggc      2259
Val Gly Pro Thr Asn Val Ser Leu Thr Val Thr Asn Met Pro Pro Gly
650                 655                 660                 665

aag cac ttc cgg gta gac ggc acc tcc gtg ctg aga ggc ttc tct ttc      2307
Lys His Phe Arg Val Asp Gly Thr Ser Val Leu Arg Gly Phe Ser Phe
                670                 675                 680

atg gag cca gtg ctg ata gca gtg caa ccc ctc ttt ggc cca cgg gca      2355
Met Glu Pro Val Leu Ile Ala Val Gln Pro Leu Phe Gly Pro Arg Ala
            685                 690                 695

gga ggc acc tgt ctc act ctt gaa ggc cag agt ctg tct gta ggc acc      2403
Gly Gly Thr Cys Leu Thr Leu Glu Gly Gln Ser Leu Ser Val Gly Thr
        700                 705                 710

agc cgg gct gtg ctg gtc aat ggg act gag tgt ctg cta gca cgg gtc      2451
Ser Arg Ala Val Leu Val Asn Gly Thr Glu Cys Leu Leu Ala Arg Val
    715                 720                 725

agt gag ggg cag ctt tta tgt gcc aca ccc cct ggg gcc acg gtg gcc      2499
Ser Glu Gly Gln Leu Leu Cys Ala Thr Pro Pro Gly Ala Thr Val Ala
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agt gtc ccc ctt agc ctg cag gtg ggg ggt gcc cag gta cct ggt tcc      2547
Ser Val Pro Leu Ser Leu Gln Val Gly Gly Ala Gln Val Pro Gly Ser
                750                 755                 760

tgg acc ttc cag tac aga gaa gac cct gtc gtg cta agc atc agc ccc      2595
Trp Thr Phe Gln Tyr Arg Glu Asp Pro Val Val Leu Ser Ile Ser Pro
            765                 770                 775

aac tgt ggc tac atc aac tcc cac atc acc atc tgt ggc cag cat cta      2643
Asn Cys Gly Tyr Ile Asn Ser His Ile Thr Ile Cys Gly Gln His Leu
        780                 785                 790

act tca gca tgg cac tta gtg ctg tca ttc cat gac ggg ctt agg gca      2691
Thr Ser Ala Trp His Leu Val Leu Ser Phe His Asp Gly Leu Arg Ala
    795                 800                 805

gtg gaa agc agg tgt gag agg cag ctt cca gag cag cag ctg tgc cgc      2739
Val Glu Ser Arg Cys Glu Arg Gln Leu Pro Glu Gln Gln Leu Cys Arg
810                 815                 820                 825

ctt cct gaa tat gtg gtc cga gac ccc cag gga tgg gtg gca ggg aat      2787
Leu Pro Glu Tyr Val Val Arg Asp Pro Gln Gly Trp Val Ala Gly Asn
                830                 835                 840

ctg agt gcc cga ggg gat gga gct gct ggc ttt aca ctg cct ggc ttt      2835
Leu Ser Ala Arg Gly Asp Gly Ala Ala Gly Phe Thr Leu Pro Gly Phe
            845                 850                 855

cgc ttc cta ccc cca ccc cat cca ccc agt gcc aac cta gtt cca ctg      2883
Arg Phe Leu Pro Pro Pro His Pro Pro Ser Ala Asn Leu Val Pro Leu
        860                 865                 870

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Lys Pro Glu Glu His Ala Ile Lys Phe Glu Tyr Ile Gly Leu Gly Ala
    875                 880                 885
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Val Ala Asp Cys Val Gly Ile Asn Val Thr Val Gly Gly Glu Ser Cys
890                 895                 900                 905

cag cac gag ttc cgg ggg gac atg gtt gtc tgc ccc ctg ccc cca tcc      3027
Gln His Glu Phe Arg Gly Asp Met Val Val Cys Pro Leu Pro Pro Ser
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ctg cag ctt ggc cag gat ggt gcc cca ttg cag gtc tgc gta gat ggt      3075
Leu Gln Leu Gly Gln Asp Gly Ala Pro Leu Gln Val Cys Val Asp Gly
            925                 930                 935

gaa tgt cat atc ctg ggt aga gtg gtg cgg cca ggg cca gat ggg gtc      3123
Glu Cys His Ile Leu Gly Arg Val Val Arg Pro Gly Pro Asp Gly Val
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cca cag agc acg ctc ctt ggt atc ctg ctg cct ttg ctg ctg ctt gtg      3171
Pro Gln Ser Thr Leu Leu Gly Ile Leu Leu Pro Leu Leu Leu Leu Val
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gct gca ctg gcg act gca ctg gtc ttc agc tac tgg tgg cgg agg aag      3219
Ala Ala Leu Ala Thr Ala Leu Val Phe Ser Tyr Trp Trp Arg Arg Lys
970                 975                 980                 985

cag cta gtt ctt cct ccc aac ctg aat gac ctg gca tcc ctg gac  cag      3267
Gln Leu Val Leu Pro Pro Asn Leu Asn Asp Leu Ala Ser Leu Asp  Gln
                990                 995                1000

act gct gga gcc  aca ccc ctg cct att  ctg tac tcg ggc tct  gac      3312
Thr Ala Gly Ala  Thr Pro Leu Pro Ile  Leu Tyr Ser Gly Ser  Asp
            1005                1010                1015

tac aga agt ggc  ctt gca ctc cct gcc  att gat ggt ctg gat  tcc      3357
Tyr Arg Ser Gly  Leu Ala Leu Pro Ala  Ile Asp Gly Leu Asp  Ser
        1020                1025                1030

acc act tgt gtc  cat gga gca tcc ttc  tcc gat agt gaa gat  gaa      3402
Thr Thr Cys Val  His Gly Ala Ser Phe  Ser Asp Ser Glu Asp  Glu
    1035                1040                1045

tcc tgt gtg cca  ctg ctg cgg aaa gag  tcc atc cag cta agg  gac      3447
Ser Cys Val Pro  Leu Leu Arg Lys Glu  Ser Ile Gln Leu Arg  Asp
1050                1055                1060

ctg gac tct gcg  ctc ttg gct gag gtc  aag gat gtg ctg att  ccc      3492
Leu Asp Ser Ala  Leu Leu Ala Glu Val  Lys Asp Val Leu Ile  Pro
    1065                1070                1075

cat gag cgg gtg  gtc acc cac agt gac  cga gtc att ggc aaa  ggc      3537
His Glu Arg Val  Val Thr His Ser Asp  Arg Val Ile Gly Lys  Gly
    1080                1085                1090

cac ttt gga gtt  gtc tac cac gga gaa  tac ata gac cag gcc  cag      3582
His Phe Gly Val  Val Tyr His Gly Glu  Tyr Ile Asp Gln Ala  Gln
    1095                1100                1105

aat cga atc caa  tgt gcc atc aag tca  cta agt cgc atc aca  gag      3627
Asn Arg Ile Gln  Cys Ala Ile Lys Ser  Leu Ser Arg Ile Thr  Glu
    1110                1115                1120

atg cag cag gtg  gag gcc ttc ctg cga  gag ggg ctg ctc atg  cgt      3672
Met Gln Gln Val  Glu Ala Phe Leu Arg  Glu Gly Leu Leu Met  Arg
    1125                1130                1135

ggc ctg aac cac  ccg aat gtg ctg gct  ctc att ggt atc atg  ttg      3717
Gly Leu Asn His  Pro Asn Val Leu Ala  Leu Ile Gly Ile Met  Leu
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Pro Pro Glu Gly  Leu Pro His Val Leu  Leu Pro Tyr Met Cys  His
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ggt gac ctg ctc  cag ttc atc cgc tca  cct cag cgg aac ccc  acc        3807
Gly Asp Leu Leu  Gln Phe Ile Arg Ser  Pro Gln Arg Asn Pro  Thr
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Val Lys Asp Leu  Ile Ser Phe Gly Leu  Gln Val Ala Arg Gly  Met
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gag tac ctg gca  gag cag aag ttt gtg  cac agg gac ctg gct  gcg        3897
Glu Tyr Leu Ala  Glu Gln Lys Phe Val  His Arg Asp Leu Ala  Ala
            1200                 1205                 1210

cgg aac tgc atg  ctg gac gag tca ttc  aca gtc aag gtg gct  gac        3942
Arg Asn Cys Met  Leu Asp Glu Ser Phe  Thr Val Lys Val Ala  Asp
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ttt ggt ttg gcc  cgc gac atc ctg gac  agg gag tac tat agt  gtt        3987
Phe Gly Leu Ala  Arg Asp Ile Leu Asp  Arg Glu Tyr Tyr Ser  Val
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caa cag cat cgc  cac gct cgc cta cct  gtg aag tgg atg gcg  ctg        4032
Gln Gln His Arg  His Ala Arg Leu Pro  Val Lys Trp Met Ala  Leu
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gag agc ctg cag  acc tat aga ttt acc  acc aag tct gat gtg  tgg        4077
Glu Ser Leu Gln  Thr Tyr Arg Phe Thr  Thr Lys Ser Asp Val  Trp
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tca ttt ggt gtg  ctg ctg tgg gaa ctg  ctg aca cgg ggt gcc  cca        4122
Ser Phe Gly Val  Leu Leu Trp Glu Leu  Leu Thr Arg Gly Ala  Pro
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cca tac cgc cac  att gac cct ttt gac  ctt acc cac ttc ctg  gcc        4167
Pro Tyr Arg His  Ile Asp Pro Phe Asp  Leu Thr His Phe Leu  Ala
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cag ggt cgg cgc  ctg ccc cag cct gag  tat tgc cct gat tct  ctg        4212
Gln Gly Arg Arg  Leu Pro Gln Pro Glu  Tyr Cys Pro Asp Ser  Leu
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tac caa gtg atg  cag caa tgc tgg gag  gca gac cca gca gtg  cga        4257
Tyr Gln Val Met  Gln Gln Cys Trp Glu  Ala Asp Pro Ala Val  Arg
            1320                 1325                 1330

ccc acc ttc aga  gta cta gtg ggg gag  gtg gag cag ata gtg  tct        4302
Pro Thr Phe Arg  Val Leu Val Gly Glu  Val Glu Gln Ile Val  Ser
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gca ctg ctt ggg  gac cat tat gtg cag  ctg cca gca acc tac  atg        4347
Ala Leu Leu Gly  Asp His Tyr Val Gln  Leu Pro Ala Thr Tyr  Met
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aac ttg ggc ccc  agc acc tcg cat gag  atg aat gtg cgt cca  gaa        4392
Asn Leu Gly Pro  Ser Thr Ser His Glu  Met Asn Val Arg Pro  Glu
            1365                 1370                 1375

cag ccg cag ttc  tca ccc atg cca ggg  aat gta cgc cgg ccc  cgg        4437
Gln Pro Gln Phe  Ser Pro Met Pro Gly  Asn Val Arg Arg Pro  Arg
            1380                 1385                 1390

cca ctc tca gag  cct cct cgg ccc act  tga cttagttctt gggctggacc      4487
Pro Leu Ser Glu  Pro Pro Arg Pro Thr
            1395                 1400
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tggccctcca ccttgttcct gccctttaac tttcagaggc aataggtaaa tggggcccat 4607

taggtccctc actccacaga gtgagccagt gagggcagtc ctgcaacatg tatttatgga 4667

gtgcctgctg tggaccctgt cttctgggca cagtggactc agcagtgacc acaccaacac 4727

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Pro Tyr Ala Ala Ser Arg Asp Phe Asp Val Lys Tyr Val Val Pro Ser
35 40 45

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

Gly Asp Pro Gly Cys Gln Thr Cys Ala Ala Cys Gly Pro Gly Pro His
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Gly Pro Pro Gly Asp Thr Asp Thr Lys Val Leu Val Leu Asp Pro Ala
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Leu Pro Ala Leu Val Ser Cys Gly Ser Ser Leu Gln Gly Arg Cys Phe
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Leu His Asp Leu Glu Pro Gln Gly Thr Ala Val His Leu Ala Ala Pro
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Ala Cys Leu Phe Ser Ala His His Asn Arg Pro Asp Asp Cys Pro Asp
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Cys Val Ala Ser Pro Leu Gly Thr Arg Val Thr Val Val Glu Gln Gly
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Gln Ala Ser Tyr Phe Tyr Val Ala Ser Ser Leu Asp Ala Ala Val Ala
195 200 205

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Ala Ser Phe Ser Pro Arg Ser Val Ser Ile Arg Arg Leu Lys Ala Asp
210 215 220

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

His Leu Val Ser Tyr Ser Ile Glu Tyr Val His Ser Phe His Thr Gly
245 250 255

Ala Phe Val Tyr Phe Leu Thr Val Gln Pro Ala Ser Val Thr Asp Asp
260 265 270

Pro Ser Ala Leu His Thr Arg Leu Ala Arg Leu Ser Ala Thr Glu Pro
275 280 285

Glu Leu Gly Asp Tyr Arg Glu Leu Val Leu Asp Cys Arg Phe Ala Pro
290 295 300

Lys Arg Arg Arg Arg Gly Ala Pro Glu Gly Gly Gln Pro Tyr Pro Val
305 310 315 320

Leu Arg Val Ala His Ser Ala Pro Val Gly Ala Gln Leu Ala Thr Glu
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Leu Ser Ile Ala Glu Gly Gln Glu Val Leu Phe Gly Val Phe Val Thr
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Gly Lys Asp Gly Gly Pro Gly Val Gly Pro Asn Ser Val Val Cys Ala
355 360 365

Phe Pro Ile Asp Leu Leu Asp Thr Leu Ile Asp Glu Gly Val Glu Arg
370 375 380

Cys Cys Glu Ser Pro Val His Pro Gly Leu Arg Arg Gly Leu Asp Phe
385 390 395 400

Phe Gln Ser Pro Ser Phe Cys Pro Asn Pro Pro Gly Leu Glu Ala Leu
405 410 415

Ser Pro Asn Thr Ser Cys Arg His Phe Pro Leu Leu Val Ser Ser Ser
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Phe Ser Arg Val Asp Leu Phe Asn Gly Leu Leu Gly Pro Val Gln Val
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Thr Ala Leu Tyr Val Thr Arg Leu Asp Asn Val Thr Val Ala His Met
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Gly Thr Met Asp Gly Arg Ile Leu Gln Val Glu Leu Val Arg Ser Leu
465 470 475 480

Asn Tyr Leu Leu Tyr Val Ser Asn Phe Ser Leu Gly Asp Ser Gly Gln
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Pro Val Gln Arg Asp Val Ser Arg Leu Gly Asp His Leu Leu Phe Ala
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Ser Gly Asp Gln Val Phe Gln Val Pro Ile Gln Gly Pro Gly Cys Arg
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His Phe Leu Thr Cys Gly Arg Cys Leu Arg Ala Trp His Phe Met Gly
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Cys Gly Trp Cys Gly Asn Met Cys Gly Gln Gln Lys Glu Cys Pro Gly
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Ser Trp Gln Gln Asp His Cys Pro Pro Lys Leu Thr Glu Phe His Pro
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His Ser Gly Pro Leu Arg Gly Ser Thr Arg Leu Thr Leu Cys Gly Ser
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Asn Phe Tyr Leu His Pro Ser Gly Leu Val Pro Glu Gly Thr His Gln
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Val Thr Val Gly Gln Ser Pro Cys Arg Pro Leu Pro Lys Asp Ser Ser
610 615 620

Lys Leu Arg Pro Val Pro Arg Lys Asp Phe Val Glu Glu Phe Glu Cys
625 630 635 640

Glu Leu Glu Pro Leu Gly Thr Gln Ala Val Gly Pro Thr Asn Val Ser
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Leu Thr Val Thr Asn Met Pro Pro Gly Lys His Phe Arg Val Asp Gly
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Thr Ser Val Leu Arg Gly Phe Ser Phe Met Glu Pro Val Leu Ile Ala
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Val Gln Pro Leu Phe Gly Pro Arg Ala Gly Gly Thr Cys Leu Thr Leu
690 695 700

Glu Gly Gln Ser Leu Ser Val Gly Thr Ser Arg Ala Val Leu Val Asn
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Gly Thr Glu Cys Leu Leu Ala Arg Val Ser Glu Gly Gln Leu Leu Cys
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Ala Thr Pro Pro Gly Ala Thr Val Ala Ser Val Pro Leu Ser Leu Gln
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Val Gly Gly Ala Gln Val Pro Gly Ser Trp Thr Phe Gln Tyr Arg Glu
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Asp Pro Val Val Leu Ser Ile Ser Pro Asn Cys Gly Tyr Ile Asn Ser
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His Ile Thr Ile Cys Gly Gln His Leu Thr Ser Ala Trp His Leu Val
785 790 795 800

Leu Ser Phe His Asp Gly Leu Arg Ala Val Glu Ser Arg Cys Glu Arg
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Gln Leu Pro Glu Gln Gln Leu Cys Arg Leu Pro Glu Tyr Val Val Arg
820 825 830

Asp Pro Gln Gly Trp Val Ala Gly Asn Leu Ser Ala Arg Gly Asp Gly
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Ala Ala Gly Phe Thr Leu Pro Gly Phe Arg Phe Leu Pro Pro Pro His
850 855 860

Pro Pro Ser Ala Asn Leu Val Pro Leu Lys Pro Glu Glu His Ala Ile
865 870 875 880

Lys Phe Glu Tyr Ile Gly Leu Gly Ala Val Ala Asp Cys Val Gly Ile
885 890 895

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Met Val Val Cys Pro Leu Pro Pro Ser Leu Gln Leu Gly Gln Asp Gly
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Ala Pro Leu Gln Val Cys Val Asp Gly Glu Cys His Ile Leu Gly Arg
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Val Val Arg Pro Gly Pro Asp Gly Val Pro Gln Ser Thr Leu Leu Gly
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Val Phe Ser Tyr Trp Trp Arg Arg Lys Gln Leu Val Leu Pro Pro Asn
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Leu Asn Asp Leu Ala Ser Leu Asp Gln Thr Ala Gly Ala Thr Pro Leu
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Pro Ala Ile Asp Gly Leu Asp Ser Thr Thr Cys Val His Gly Ala
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Ser Phe Ser Asp Ser Glu Asp Glu Ser Cys Val Pro Leu Leu Arg
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Glu Val Lys Asp Val Leu Ile Pro His Glu Arg Val Val Thr His
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Ser Asp Arg Val Ile Gly Lys Gly His Phe Gly Val Val Tyr His
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Gly Glu Tyr Ile Asp Gln Ala Gln Asn Arg Ile Gln Cys Ala Ile
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Lys Ser Leu Ser Arg Ile Thr Glu Met Gln Gln Val Glu Ala Phe
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Leu Arg Glu Gly Leu Leu Met Arg Gly Leu Asn His Pro Asn Val
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Leu Ala Leu Ile Gly Ile Met Leu Pro Pro Glu Gly Leu Pro His
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Val Leu Leu Pro Tyr Met Cys His Gly Asp Leu Leu Gln Phe Ile
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Gly Leu Gln Val Ala Arg Gly Met Glu Tyr Leu Ala Glu Gln Lys
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Ser Phe Thr Val Lys Val Ala Asp Phe Gly Leu Ala Arg Asp Ile
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Leu Asp Arg Glu Tyr Tyr Ser Val Gln Gln His Arg His Ala Arg
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Leu Pro Val Lys Trp Met Ala Leu Glu Ser Leu Gln Thr Tyr Arg
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Phe Thr Thr Lys Ser Asp Val Trp Ser Phe Gly Val Leu Leu Trp
1265 1270 1275

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Glu Leu Leu Thr Arg Gly Ala Pro Pro Tyr Arg His Ile Asp Pro
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Phe Asp Leu Thr His Phe Leu Ala Gln Gly Arg Arg Leu Pro Gln
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Pro Glu Tyr Cys Pro Asp Ser Leu Tyr Gln Val Met Gln Gln Cys
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Trp Glu Ala Asp Pro Ala Val Arg Pro Thr Phe Arg Val Leu Val
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Gly Glu Val Glu Gln Ile Val Ser Ala Leu Leu Gly Asp His Tyr
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Val Gln Leu Pro Ala Thr Tyr Met Asn Leu Gly Pro Ser Thr Ser
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His Glu Met Asn Val Arg Pro Glu Gln Pro Gln Phe Ser Pro Met
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Trp Gln Cys Pro Arg Ile Pro Tyr Ala Ala Ser Arg Asp Phe Ser Val
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Lys Tyr Val Val Pro Ser Phe Ser Ala Gly Gly Arg Val Gln Ala Thr
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gca gcc tac gag gac agt aca aat agt gcg gtg ttt gtg gcc aca cgc          484
Ala Ala Tyr Glu Asp Ser Thr Asn Ser Ala Val Phe Val Ala Thr Arg
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aat cac ctg cac gtg ctt ggg cct gac ctg cag ttc ata gag aac ctg          532
Asn His Leu His Val Leu Gly Pro Asp Leu Gln Phe Ile Glu Asn Leu
            80                  85                  90

acc act ggc cct atc ggg aac cct ggc tgc cag act tgt gcg agc tgt          580
Thr Thr Gly Pro Ile Gly Asn Pro Gly Cys Gln Thr Cys Ala Ser Cys
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ggt cca ggc cct cat gga cca cca aag gac aca gac aca ctg gtg cta          628
Gly Pro Gly Pro His Gly Pro Pro Lys Asp Thr Asp Thr Leu Val Leu
    110                 115                 120

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Val Met Glu Pro Gly Leu Pro Ala Leu Val Ser Cys Gly Ser Thr Leu
125                 130                 135                 140

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Gln Gly Arg Cys Phe Leu His Glu Leu Glu Pro Arg Gly Lys Ala Leu
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cac tta gca gct cca gcc tgc cta ttc tca gca aac aat aac aag cct          772
His Leu Ala Ala Pro Ala Cys Leu Phe Ser Ala Asn Asn Asn Lys Pro
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Glu Ala Cys Thr Asp Cys Val Ala Ser Pro Leu Gly Thr Arg Val Thr
        175                 180                 185

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Val Val Glu Gln Gly His Ala Ser Tyr Phe Tyr Val Ala Ser Ser Leu
    190                 195                 200

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Asp Pro Glu Leu Ala Ala Ser Phe Ser Pro Arg Ser Val Ser Ile Arg
205                 210                 215                 220

cgt cta aag tct gat act tct gga ttc caa cca ggt ttt ccg tcg ctg          964
Arg Leu Lys Ser Asp Thr Ser Gly Phe Gln Pro Gly Phe Pro Ser Leu
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Ser Val Leu Pro Lys Tyr Leu Ala Ser Tyr Leu Ile Lys Tyr Val Tyr
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Ser Val Thr Ser Pro Pro Ser Ala Leu His Thr Arg Leu Val Arg Leu
    270                 275                 280

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Asn Ala Val Glu Pro Glu Ile Gly Asp Tyr Arg Glu Leu Val Leu Asp
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Cys His Phe Ala Pro Lys Arg Arg Arg Arg Gly Ala Pro Glu Gly Thr
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Gln Pro Tyr Pro Val Leu Gln Ala Ala His Ser Ala Pro Val Asp Ala
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Lys Leu Ala Val Glu Leu Ser Ile Ser Glu Gly Gln Glu Val Leu Phe
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Gly Val Phe Val Thr Val Lys Asp Gly Gly Ser Gly Met Gly Pro Asn
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Ser Val Val Cys Ala Phe Pro Ile Tyr His Leu Asn Ile Leu Ile Glu
365                 370                 375                 380

gag ggt gtc gaa tat tgc tgt cac tct tca aat tct tct tcc ctg ttg      1444
Glu Gly Val Glu Tyr Cys Cys His Ser Ser Asn Ser Ser Ser Leu Leu
                385                 390                 395

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            400                 405                 410

cct ggt gga gag gcc tcc ggc ccc agc tcc cgt tgc cac tac ttc cct      1540
Pro Gly Gly Glu Ala Ser Gly Pro Ser Ser Arg Cys His Tyr Phe Pro
        415                 420                 425

ttg atg gtc cac gct agc ttc acc cgt gtg gac ctc ttc aat gga ctg      1588
Leu Met Val His Ala Ser Phe Thr Arg Val Asp Leu Phe Asn Gly Leu
    430                 435                 440

tta gga tca gtg aag gtc acc gca ctg cat gtg aca cgt ctt ggc aat      1636
Leu Gly Ser Val Lys Val Thr Ala Leu His Val Thr Arg Leu Gly Asn
445                 450                 455                 460

gtt aca gtg gcc cac atg ggc act gtg gat ggg cgt gtc cta cag gtg      1684
Val Thr Val Ala His Met Gly Thr Val Asp Gly Arg Val Leu Gln Val
                465                 470                 475

gag ata gcc agg tca ctc aac tac ctg ctg tat gtg tcc aac ttc tcc      1732
Glu Ile Ala Arg Ser Leu Asn Tyr Leu Leu Tyr Val Ser Asn Phe Ser
            480                 485                 490

ctg ggc agc agt gga cag cct gtt cat cgg gat gtc agc cgc ctc ggg      1780
Leu Gly Ser Ser Gly Gln Pro Val His Arg Asp Val Ser Arg Leu Gly
        495                 500                 505

aat gac cta ctc ttt gcc tct ggg gac cag gtc ttc aag gtg ccc atc      1828
Asn Asp Leu Leu Phe Ala Ser Gly Asp Gln Val Phe Lys Val Pro Ile
    510                 515                 520

cag ggc cct ggc tgt cgt cat ttt ctc acc tgt tgg cgt tgc ctg aga      1876
Gln Gly Pro Gly Cys Arg His Phe Leu Thr Cys Trp Arg Cys Leu Arg
525                 530                 535                 540

gca cag cgc ttc atg gga tgt ggc tgg tgt ggg gac cgg tgt gac cgg      1924
Ala Gln Arg Phe Met Gly Cys Gly Trp Cys Gly Asp Arg Cys Asp Arg
                545                 550                 555

cag aag gag tgt cct ggc tcc tgg caa cag gac cac tgt ccg cct gag      1972
Gln Lys Glu Cys Pro Gly Ser Trp Gln Gln Asp His Cys Pro Pro Glu
            560                 565                 570

atc agt gag ttc tat cct cac agc ggg cct cta agg ggc act acg agg      2020
Ile Ser Glu Phe Tyr Pro His Ser Gly Pro Leu Arg Gly Thr Thr Arg
        575                 580                 585
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ctc acc ctt tgt ggc tcc aac ttc tac ctg cga cct gat gat gtc gta      2068
Leu Thr Leu Cys Gly Ser Asn Phe Tyr Leu Arg Pro Asp Asp Val Val
    590                 595                 600

cct gag gga aca cac cag atc acc gtg ggc caa agt ccc tgc cga ctg      2116
Pro Glu Gly Thr His Gln Ile Thr Val Gly Gln Ser Pro Cys Arg Leu
605                 610                 615                 620

ctg cct aag gac tct tca agc cct agg cca ggg tcc ctc aag gaa ttc      2164
Leu Pro Lys Asp Ser Ser Ser Pro Arg Pro Gly Ser Leu Lys Glu Phe
                625                 630                 635

ata cag gaa ctt gaa tgt gag ctg gag ccc ctg gtc acc cag gca gtg      2212
Ile Gln Glu Leu Glu Cys Glu Leu Glu Pro Leu Val Thr Gln Ala Val
            640                 645                 650

ggg act aca aac atc agc ctt gtc atc acc aac atg cca gca ggc aag      2260
Gly Thr Thr Asn Ile Ser Leu Val Ile Thr Asn Met Pro Ala Gly Lys
        655                 660                 665

cac ttc cga gtg gaa ggc atc tct gta cag gaa ggc ttc tct ttc gtg      2308
His Phe Arg Val Glu Gly Ile Ser Val Gln Glu Gly Phe Ser Phe Val
    670                 675                 680

gag cca gtg ctg aca tca ata aaa cct gac ttt ggc ccg cgg gct ggg      2356
Glu Pro Val Leu Thr Ser Ile Lys Pro Asp Phe Gly Pro Arg Ala Gly
685                 690                 695                 700

ggt act tat ctc acc ctc gaa ggc cag agc ctg tct att gcc acc agc      2404
Gly Thr Tyr Leu Thr Leu Glu Gly Gln Ser Leu Ser Ile Ala Thr Ser
                705                 710                 715

cga gct gcg ctg gtc aat gga acc cag tgc cgg ctg gaa cag gtc aat      2452
Arg Ala Ala Leu Val Asn Gly Thr Gln Cys Arg Leu Glu Gln Val Asn
            720                 725                 730

gag gag cag atc tta tgt gtc acg cct cct gga gct ggc acg gcc agg      2500
Glu Glu Gln Ile Leu Cys Val Thr Pro Pro Gly Ala Gly Thr Ala Arg
        735                 740                 745

gtc ccc ctt cat ctg cag ata ggg ggt gct gag gtg cct ggc tcc tgg      2548
Val Pro Leu His Leu Gln Ile Gly Gly Ala Glu Val Pro Gly Ser Trp
    750                 755                 760

acc ttt cac tac aag gaa gac cct att gtg ttg gac atc agt ccc aag      2596
Thr Phe His Tyr Lys Glu Asp Pro Ile Val Leu Asp Ile Ser Pro Lys
765                 770                 775                 780

tgt ggc tac agt ggc tcc cac atc atg atc cat ggc cag cat ctg act      2644
Cys Gly Tyr Ser Gly Ser His Ile Met Ile His Gly Gln His Leu Thr
                785                 790                 795

tca gca tgg cac ttc acg cta tca ttc cat gat gga caa agt aca gtg      2692
Ser Ala Trp His Phe Thr Leu Ser Phe His Asp Gly Gln Ser Thr Val
            800                 805                 810

gag agc agg tgt gcg ggg cag ttt gtg gaa caa cag cag cgt cga tgt      2740
Glu Ser Arg Cys Ala Gly Gln Phe Val Glu Gln Gln Gln Arg Arg Cys
        815                 820                 825

cgc ctg cct gaa tat gtg gtc cga aac cct cag ggg tgg gca aca ggg      2788
Arg Leu Pro Glu Tyr Val Val Arg Asn Pro Gln Gly Trp Ala Thr Gly
    830                 835                 840

aat ctg agc gtc tgg ggt gat gga gca gct ggc ttc aca ctg cct ggt      2836
Asn Leu Ser Val Trp Gly Asp Gly Ala Ala Gly Phe Thr Leu Pro Gly
845                 850                 855                 860
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Phe Arg Phe Leu Pro Pro Pro Ser Pro Leu Arg Ala Gly Leu Val Glu
                865                 870                 875

ttg aaa cct gaa gaa cat tca gtt aaa gtt gag tat gtc ggg ctg ggc       2932
Leu Lys Pro Glu Glu His Ser Val Lys Val Glu Tyr Val Gly Leu Gly
            880                 885                 890

gct gtg gca gac tgt gtg act gtg aac atg acc gtg ggt ggt gag gtc       2980
Ala Val Ala Asp Cys Val Thr Val Asn Met Thr Val Gly Gly Glu Val
        895                 900                 905

tgc caa cat gag ctc cgg ggg gat gtg gtg atc tgc ccc ctg ccc cct       3028
Cys Gln His Glu Leu Arg Gly Asp Val Val Ile Cys Pro Leu Pro Pro
    910                 915                 920

tcc ctg caa ctt ggc aag gat ggt gtc cca ttg cag gtc tgt gta gac       3076
Ser Leu Gln Leu Gly Lys Asp Gly Val Pro Leu Gln Val Cys Val Asp
925                 930                 935                 940

ggt ggg tgt cac atc ctg agc caa gtg gtt cgc tca agc cca ggc agg       3124
Gly Gly Cys His Ile Leu Ser Gln Val Val Arg Ser Ser Pro Gly Arg
                945                 950                 955

gcc tca cag agg ata ctc ctt att gct ctt ctg gtc ttg atc ctg ctt       3172
Ala Ser Gln Arg Ile Leu Leu Ile Ala Leu Leu Val Leu Ile Leu Leu
            960                 965                 970

gtg gct gtg ctg gcc gtt gcc ctg atc ttt aac tcc cga aga cgg aaa       3220
Val Ala Val Leu Ala Val Ala Leu Ile Phe Asn Ser Arg Arg Arg Lys
        975                 980                 985

aag cag cta ggt gct cac tcc ctc tcc cca aca aca  ctc tct gac atc      3268
Lys Gln Leu Gly Ala His Ser Leu Ser Pro Thr Thr  Leu Ser Asp Ile
    990                 995                 1000

aac  gat aca gct tcc ggg  gct ccg aac cat gaa  gaa tcg tca gag        3313
Asn  Asp Thr Ala Ser Gly  Ala Pro Asn His Glu  Glu Ser Ser Glu
1005                1010                1015

agt  agg gat ggg aca agt  gtc cca ctg ctg cgg  aca gag tct atc       3358
Ser  Arg Asp Gly Thr Ser  Val Pro Leu Leu Arg  Thr Glu Ser Ile
1020                1025                1030

cgg  ctc cag gat ctg gac  agg atg ctc cta gct  gag gtc aag gat       3403
Arg  Leu Gln Asp Leu Asp  Arg Met Leu Leu Ala  Glu Val Lys Asp
1035                1040                1045

gta  ctg att ccc cat gaa  caa gtg gtc atc cat  act gac caa gtc       3448
Val  Leu Ile Pro His Glu  Gln Val Val Ile His  Thr Asp Gln Val
1050                1055                1060

att  ggc aaa ggc cac ttt  ggt gtt gtc tac cac  gga gaa tat aca       3493
Ile  Gly Lys Gly His Phe  Gly Val Val Tyr His  Gly Glu Tyr Thr
1065                1070                1075

gac  gga gca cag aat cag  acc cac tgt gcc atc  aag tct ctg agt       3538
Asp  Gly Ala Gln Asn Gln  Thr His Cys Ala Ile  Lys Ser Leu Ser
1080                1085                1090

cgc  att aca gag gtg cag  gag gtg gag gct ttc  ctg cgg gag ggg       3583
Arg  Ile Thr Glu Val Gln  Glu Val Glu Ala Phe  Leu Arg Glu Gly
1095                1100                1105

ctg  ctc atg cgt ggc cta  cat cac cca aac atc  ctg gct ctc atc       3628
Leu  Leu Met Arg Gly Leu  His His Pro Asn Ile  Leu Ala Leu Ile
1110                1115                1120
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ggt  atc atg ctg ccc ccg  gag ggg ctt ccc cgg  gtg ctg ttg ccc        3673
Gly  Ile Met Leu Pro Pro  Glu Gly Leu Pro Arg  Val Leu Leu Pro
1125                 1130                 1135

tat  atg cgc cac gga gac  ctg ctt cgt ttc att  cgc tcc cct cag        3718
Tyr  Met Arg His Gly Asp  Leu Leu Arg Phe Ile  Arg Ser Pro Gln
1140                 1145                 1150

agg  aac ccc act gtg aag  gat ctt gtc agc ttt  ggc ctg cag gta        3763
Arg  Asn Pro Thr Val Lys  Asp Leu Val Ser Phe  Gly Leu Gln Val
1155                 1160                 1165

gcc  tgt ggt atg gag tac  ctg gca gag cag aag  ttc gtg cac aga        3808
Ala  Cys Gly Met Glu Tyr  Leu Ala Glu Gln Lys  Phe Val His Arg
1170                 1175                 1180

gac  ctg gct gct agg aac  tgc atg ctg gac gag  tca ttc aca gtc        3853
Asp  Leu Ala Ala Arg Asn  Cys Met Leu Asp Glu  Ser Phe Thr Val
1185                 1190                 1195

aag  gtg gct gac ttt ggt  ctg gca cgg ggc gtc  cta gac aag gaa        3898
Lys  Val Ala Asp Phe Gly  Leu Ala Arg Gly Val  Leu Asp Lys Glu
1200                 1205                 1210

tac  tac agt gtt cgc cag  cat cgc cat gct cgc  ctg cca gtc aaa        3943
Tyr  Tyr Ser Val Arg Gln  His Arg His Ala Arg  Leu Pro Val Lys
1215                 1220                 1225

tgg  atg gca ctg gag agc  ctg cag acc tac agg  ttc acc acc aag        3988
Trp  Met Ala Leu Glu Ser  Leu Gln Thr Tyr Arg  Phe Thr Thr Lys
1230                 1235                 1240

tcc  gat gtg tgg tca ttc  ggg gtg ctg ctc tgg  gag cta cta aca        4033
Ser  Asp Val Trp Ser Phe  Gly Val Leu Leu Trp  Glu Leu Leu Thr
1245                 1250                 1255

cgg  ggt gct cca ccc tac  ccc cat atc gat ccc  ttc gac ctc tct        4078
Arg  Gly Ala Pro Pro Tyr  Pro His Ile Asp Pro  Phe Asp Leu Ser
1260                 1265                 1270

cac  ttc ctg gct cag ggc  cgc cgc ctg cct cag  cct gag tac tgt        4123
His  Phe Leu Ala Gln Gly  Arg Arg Leu Pro Gln  Pro Glu Tyr Cys
1275                 1280                 1285

cct  gat tca ctg tat cac  gtg atg ctt cga tgc  tgg gag gct gac        4168
Pro  Asp Ser Leu Tyr His  Val Met Leu Arg Cys  Trp Glu Ala Asp
1290                 1295                 1300

cca  gcg gca cga ccc acc  ttc aga gcc cta gtg  ctg gaa gta aag        4213
Pro  Ala Ala Arg Pro Thr  Phe Arg Ala Leu Val  Leu Glu Val Lys
1305                 1310                 1315

cag  gta gtg gcc tca ctg  ctt ggg gac cac tat  gtg cag ctg aca        4258
Gln  Val Val Ala Ser Leu  Leu Gly Asp His Tyr  Val Gln Leu Thr
1320                 1325                 1330

gca  gct tat gtg aac gta  ggc ccc aga gcg gtg  gat gat ggg agt        4303
Ala  Ala Tyr Val Asn Val  Gly Pro Arg Ala Val  Asp Asp Gly Ser
1335                 1340                 1345

gtg  cct ccg gag cag gta  cag ccc tcg cct cag  cat tgc agg agc        4348
Val  Pro Pro Glu Gln Val  Gln Pro Ser Pro Gln  His Cys Arg Ser
1350                 1355                 1360

acg  tca aag ccc cgg cct  ctc tca gag cca ccc  ctg ccc act tga        4393
Thr  Ser Lys Pro Arg Pro  Leu Ser Glu Pro Pro  Leu Pro Thr
1365                 1370                 1375
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ccaaagccct gagtaggcca caggcactag atctgctaag tggccttgag caaattacaa 4453

gctgcctctg ggcctaggac aagcctcagc atggaaaacc tccactcttt agctttctgg 4513

ggccactgaa ggtgggaaac cgggcccatt tgagcccctc gttccagcat gagccagtga 4573

cattttgta gcatgtattt atgtaatgtc tgttttgtac ctgtttcgga atacagagga 4633

tccagcagtg atcacagaga tactacagta taaaataata taaataaata aatgaatgaa 4693

tattcgagca caaaaaaaaa aaaaaaa 4720

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Arg Ile Pro Tyr Ala Ala Ser Arg Asp Phe Ser Val Lys Tyr Val Val
35 40 45

Pro Ser Phe Ser Ala Gly Gly Arg Val Gln Ala Thr Ala Ala Tyr Glu
50 55 60

Asp Ser Thr Asn Ser Ala Val Phe Val Ala Thr Arg Asn His Leu His
65 70 75 80

Val Leu Gly Pro Asp Leu Gln Phe Ile Glu Asn Leu Thr Thr Gly Pro
85 90 95

Ile Gly Asn Pro Gly Cys Gln Thr Cys Ala Ser Cys Gly Pro Gly Pro
100 105 110

His Gly Pro Pro Lys Asp Thr Asp Thr Leu Val Leu Val Met Glu Pro
115 120 125

Gly Leu Pro Ala Leu Val Ser Cys Gly Ser Thr Leu Gln Gly Arg Cys
130 135 140

Phe Leu His Glu Leu Glu Pro Arg Gly Lys Ala Leu His Leu Ala Ala
145 150 155 160

Pro Ala Cys Leu Phe Ser Ala Asn Asn Asn Lys Pro Glu Ala Cys Thr
165 170 175

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

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Gly His Ala Ser Tyr Phe Tyr Val Ala Ser Ser Leu Asp Pro Glu Leu
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Ala Ala Ser Phe Ser Pro Arg Ser Val Ser Ile Arg Arg Leu Lys Ser
    210                 215                 220

Asp Thr Ser Gly Phe Gln Pro Gly Phe Pro Ser Leu Ser Val Leu Pro
225                 230                 235                 240

Lys Tyr Leu Ala Ser Tyr Leu Ile Lys Tyr Val Tyr Ser Phe His Ser
                245                 250                 255

Gly Asp Phe Val Tyr Phe Leu Thr Val Gln Pro Ile Ser Val Thr Ser
            260                 265                 270

Pro Pro Ser Ala Leu His Thr Arg Leu Val Arg Leu Asn Ala Val Glu
        275                 280                 285

Pro Glu Ile Gly Asp Tyr Arg Glu Leu Val Leu Asp Cys His Phe Ala
    290                 295                 300

Pro Lys Arg Arg Arg Arg Gly Ala Pro Glu Gly Thr Gln Pro Tyr Pro
305                 310                 315                 320

Val Leu Gln Ala Ala His Ser Ala Pro Val Asp Ala Lys Leu Ala Val
                325                 330                 335

Glu Leu Ser Ile Ser Glu Gly Gln Glu Val Leu Phe Gly Val Phe Val
            340                 345                 350

Thr Val Lys Asp Gly Gly Ser Gly Met Gly Pro Asn Ser Val Val Cys
        355                 360                 365

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

Tyr Cys Cys His Ser Ser Asn Ser Ser Ser Leu Leu Ser Arg Gly Leu
385                 390                 395                 400

Asp Phe Phe Gln Thr Pro Ser Phe Cys Pro Asn Pro Pro Gly Gly Glu
                405                 410                 415

Ala Ser Gly Pro Ser Ser Arg Cys His Tyr Phe Pro Leu Met Val His
            420                 425                 430

Ala Ser Phe Thr Arg Val Asp Leu Phe Asn Gly Leu Leu Gly Ser Val
        435                 440                 445

Lys Val Thr Ala Leu His Val Thr Arg Leu Gly Asn Val Thr Val Ala
    450                 455                 460
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His Met Gly Thr Val Asp Gly Arg Val Leu Gln Val Glu Ile Ala Arg
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Ser Leu Asn Tyr Leu Leu Tyr Val Ser Asn Phe Ser Leu Gly Ser Ser
485 490 495

Gly Gln Pro Val His Arg Asp Val Ser Arg Leu Gly Asn Asp Leu Leu
500 505 510

Phe Ala Ser Gly Asp Gln Val Phe Lys Val Pro Ile Gln Gly Pro Gly
515 520 525

Cys Arg His Phe Leu Thr Cys Trp Arg Cys Leu Arg Ala Gln Arg Phe
530 535 540

Met Gly Cys Gly Trp Cys Gly Asp Arg Cys Asp Arg Gln Lys Glu Cys
545 550 555 560

Pro Gly Ser Trp Gln Gln Asp His Cys Pro Pro Glu Ile Ser Glu Phe
565 570 575

Tyr Pro His Ser Gly Pro Leu Arg Gly Thr Thr Arg Leu Thr Leu Cys
580 585 590

Gly Ser Asn Phe Tyr Leu Arg Pro Asp Asp Val Val Pro Glu Gly Thr
595 600 605

His Gln Ile Thr Val Gly Gln Ser Pro Cys Arg Leu Leu Pro Lys Asp
610 615 620

Ser Ser Ser Pro Arg Pro Gly Ser Leu Lys Glu Phe Ile Gln Glu Leu
625 630 635 640

Glu Cys Glu Leu Glu Pro Leu Val Thr Gln Ala Val Gly Thr Thr Asn
645 650 655

Ile Ser Leu Val Ile Thr Asn Met Pro Ala Gly Lys His Phe Arg Val
660 665 670

Glu Gly Ile Ser Val Gln Glu Gly Phe Ser Phe Val Glu Pro Val Leu
675 680 685

Thr Ser Ile Lys Pro Asp Phe Gly Pro Arg Ala Gly Gly Thr Tyr Leu
690 695 700

Thr Leu Glu Gly Gln Ser Leu Ser Ile Ala Thr Ser Arg Ala Ala Leu
705 710 715 720

Val Asn Gly Thr Gln Cys Arg Leu Glu Gln Val Asn Glu Glu Gln Ile
725 730 735

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Leu Cys Val Thr Pro Pro Gly Ala Gly Thr Ala Arg Val Pro Leu His
            740                 745                 750

Leu Gln Ile Gly Gly Ala Glu Val Pro Gly Ser Trp Thr Phe His Tyr
        755                 760                 765

Lys Glu Asp Pro Ile Val Leu Asp Ile Ser Pro Lys Cys Gly Tyr Ser
    770                 775                 780

Gly Ser His Ile Met Ile His Gly Gln His Leu Thr Ser Ala Trp His
785                 790                 795                 800

Phe Thr Leu Ser Phe His Asp Gly Gln Ser Thr Val Glu Ser Arg Cys
                805                 810                 815

Ala Gly Gln Phe Val Glu Gln Gln Gln Arg Arg Cys Arg Leu Pro Glu
            820                 825                 830

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

Trp Gly Asp Gly Ala Ala Gly Phe Thr Leu Pro Gly Phe Arg Phe Leu
    850                 855                 860

Pro Pro Pro Ser Pro Leu Arg Ala Gly Leu Val Glu Leu Lys Pro Glu
865                 870                 875                 880

Glu His Ser Val Lys Val Glu Tyr Val Gly Leu Gly Ala Val Ala Asp
                885                 890                 895

Cys Val Thr Val Asn Met Thr Val Gly Gly Glu Val Cys Gln His Glu
            900                 905                 910

Leu Arg Gly Asp Val Val Ile Cys Pro Leu Pro Pro Ser Leu Gln Leu
        915                 920                 925

Gly Lys Asp Gly Val Pro Leu Gln Val Cys Val Asp Gly Gly Cys His
    930                 935                 940

Ile Leu Ser Gln Val Val Arg Ser Ser Pro Gly Arg Ala Ser Gln Arg
945                 950                 955                 960

Ile Leu Leu Ile Ala Leu Leu Val Leu Ile Leu Leu Val Ala Val Leu
                965                 970                 975

Ala Val Ala Leu Ile Phe Asn Ser Arg Arg Arg Lys Lys Gln Leu Gly
            980                 985                 990

Ala His Ser Leu Ser Pro Thr Thr  Leu Ser Asp Ile Asn  Asp Thr Ala
        995                 1000                1005
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Ser Gly Ala Pro Asn His Glu Glu Ser Ser Glu Ser Arg Asp Gly
1010 1015 1020

Thr Ser Val Pro Leu Leu Arg Thr Glu Ser Ile Arg Leu Gln Asp
1025 1030 1035

Leu Asp Arg Met Leu Leu Ala Glu Val Lys Asp Val Leu Ile Pro
1040 1045 1050

His Glu Gln Val Val Ile His Thr Asp Gln Val Ile Gly Lys Gly
1055 1060 1065

His Phe Gly Val Val Tyr His Gly Glu Tyr Thr Asp Gly Ala Gln
1070 1075 1080

Asn Gln Thr His Cys Ala Ile Lys Ser Leu Ser Arg Ile Thr Glu
1085 1090 1095

Val Gln Glu Val Glu Ala Phe Leu Arg Glu Gly Leu Leu Met Arg
1100 1105 1110

Gly Leu His His Pro Asn Ile Leu Ala Leu Ile Gly Ile Met Leu
1115 1120 1125

Pro Pro Glu Gly Leu Pro Arg Val Leu Leu Pro Tyr Met Arg His
1130 1135 1140

Gly Asp Leu Leu Arg Phe Ile Arg Ser Pro Gln Arg Asn Pro Thr
1145 1150 1155

Val Lys Asp Leu Val Ser Phe Gly Leu Gln Val Ala Cys Gly Met
1160 1165 1170

Glu Tyr Leu Ala Glu Gln Lys Phe Val His Arg Asp Leu Ala Ala
1175 1180 1185

Arg Asn Cys Met Leu Asp Glu Ser Phe Thr Val Lys Val Ala Asp
1190 1195 1200

Phe Gly Leu Ala Arg Gly Val Leu Asp Lys Glu Tyr Tyr Ser Val
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Arg Gln His Arg His Ala Arg Leu Pro Val Lys Trp Met Ala Leu
1220 1225 1230

Glu Ser Leu Gln Thr Tyr Arg Phe Thr Thr Lys Ser Asp Val Trp
1235 1240 1245

Ser Phe Gly Val Leu Leu Trp Glu Leu Leu Thr Arg Gly Ala Pro
1250 1255 1260

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Pro Tyr Pro His Ile Asp Pro Phe Asp Leu Ser His Phe Leu Ala
1265 1270 1275

Gln Gly Arg Arg Leu Pro Gln Pro Glu Tyr Cys Pro Asp Ser Leu
1280 1285 1290

Tyr His Val Met Leu Arg Cys Trp Glu Ala Asp Pro Ala Ala Arg
1295 1300 1305

Pro Thr Phe Arg Ala Leu Val Leu Glu Val Lys Gln Val Val Ala
1310 1315 1320

Ser Leu Leu Gly Asp His Tyr Val Gln Leu Thr Ala Ala Tyr Val
1325 1330 1335

Asn Val Gly Pro Arg Ala Val Asp Asp Gly Ser Val Pro Pro Glu
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Gln Val Gln Pro Ser Pro Gln His Cys Arg Ser Thr Ser Lys Pro
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Arg Pro Leu Ser Glu Pro Pro Leu Pro Thr
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gtc ctg tcc cag gtg caa ttg gtt cag agc ggc gcg gaa gtg aaa aaa 96
Val Leu Ser Gln Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys
20 25 30

ccg ggc gaa agc ctg aaa att agc tgc aaa ggt tcc gga tat tcc ttt 144
Pro Gly Glu Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe
35 40 45

act aat tat tgg att tct tgg gtg cgc cag atg cct ggg aag ggt ctc 192
Thr Asn Tyr Trp Ile Ser Trp Val Arg Gln Met Pro Gly Lys Gly Leu
50 55 60

gag tgg atg ggc ttt atc tat ccg gat gat agc tat acc cgt tat tct 240
Glu Trp Met Gly Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser
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ccg agc ttt cag ggc cag gtg acc att agc gcg gat aaa agc att agc 288

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

acc gcg tat ctt caa tgg agc agc ctg aaa gcg agc gat acg gcc atg  336
Thr Ala Tyr Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met
            100                 105                 110

tat tat tgc gcg cgt ttt tct tat cgt cat tat ctt gat atg gat gat  384
Tyr Tyr Cys Ala Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp
        115                 120                 125

cat tgg ggc caa ggc acc ctg gtg acg gtt agc tca gcc tcc acc aag  432
His Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys
    130                 135                 140

ggt cca tcg gtc ttc ccc ctg gca ccc tcc tcc aag agc acc tct ggg  480
Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly
145                 150                 155                 160

ggc aca gcg gcc ctg ggc tgc ctg gtc aag gac tac ttc ccc gaa ccg  528
Gly Thr Ala Ala Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro
                165                 170                 175

gtg acg gtg tcg tgg aac tca ggc gcc ctg acc agc ggc gtg cac acc  576
Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
            180                 185                 190

ttc ccg gct gtc cta cag tcc tca gga ctc tac tcc ctc agc agc gtg  624
Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
        195                 200                 205

gtg acc gtg ccc tcc agc agc ttg ggc acc cag acc tac atc tgc aac  672
Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
    210                 215                 220

gtg aat cac aag ccc agc aac acc aag gtg gac aag aga gtt gag ccc  720
Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro
225                 230                 235                 240

aaa tct tgt gac aaa act cac aca tgc cca ccg tgc cca gca cct gaa  768
Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
                245                 250                 255

ctc ctg ggg gga ccg tca gtc ttc ctc ttc ccc cca aaa ccc aag gac  816
Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
            260                 265                 270

acc ctc atg atc tcc cgg acc cct gag gtc aca tgc gtg gtg gtg gac  864
Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
        275                 280                 285

gtg agc cac gaa gac cct gag gtc aag ttc aac tgg tac gtg gac ggc  912
Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
    290                 295                 300

gtg gag gtg cat aat gcc aag aca aag ccg cgg gag gag cag tac aac  960
Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
305                 310                 315                 320

agc acg tac cgg gtg gtc agc gtc ctc acc gtc ctg cac cag gac tgg 1008
Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
                325                 330                 335

ctg aat ggc aag gag tac aag tgc aag gtc tcc aac aaa gcc ctc cca 1056
Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro
            340                 345                 350

gcc ccc atc gag aaa acc atc tcc aaa gcc aaa ggg cag ccc cga gaa 1104
```

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```
Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
        355                 360                 365

cca cag gtg tac acc ctg ccc cca tcc cgg gag gag atg acc aag aac      1152
Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn
    370                 375                 380

cag gtc agc ctg acc tgc ctg gtc aaa ggc ttc tat ccc agc gac atc      1200
Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
385                 390                 395                 400

gcc gtg gag tgg gag agc aat ggg cag ccg gag aac aac tac aag acc      1248
Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
                405                 410                 415

acg cct ccc gtg ctg gac tcc gac ggc tcc ttc ttc ctc tac agc aag      1296
Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys
            420                 425                 430

ctc acc gtg gac aag agc agg tgg cag cag ggg aac gtc ttc tca tgc      1344
Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
        435                 440                 445

tcc gtg atg cat gag gct ctg cac aac cac tac acg cag aag agc ctc      1392
Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu
    450                 455                 460

tcc ctg tct ccg ggt aaa tga                                         1413
Ser Leu Ser Pro Gly Lys
465                 470
```

<210> 49
<211> 470
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 49

```
Met Lys His Leu Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp
1               5                   10                  15

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

Pro Gly Glu Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe
        35                  40                  45

Thr Asn Tyr Trp Ile Ser Trp Val Arg Gln Met Pro Gly Lys Gly Leu
    50                  55                  60

Glu Trp Met Gly Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser
65                  70                  75                  80

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

Thr Ala Tyr Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met
            100                 105                 110
```

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```
Tyr Tyr Cys Ala Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp
        115                 120                 125

His Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys
    130                 135                 140

Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly
145                 150                 155                 160

Gly Thr Ala Ala Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro
                165                 170                 175

Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
            180                 185                 190

Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
        195                 200                 205

Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
    210                 215                 220

Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro
225                 230                 235                 240

Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
                245                 250                 255

Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
            260                 265                 270

Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
        275                 280                 285

Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
    290                 295                 300

Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
305                 310                 315                 320

Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
                325                 330                 335

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

Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
        355                 360                 365

Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn
    370                 375                 380
```

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20102143O1

```
Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
385                 390                 395                 400

Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
                405                 410                 415

Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys
            420                 425                 430

Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
        435                 440                 445

Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu
    450                 455                 460

Ser Leu Ser Pro Gly Lys
465                 470


<210>  50
<211>  1401
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07692, MOR07923, MOR07924, MOR07925 and MOR07926 heavy chain


<220>
<221>  CDS
<222>  (1)..(1401)

<400>  50
atg aaa cac ctg tgg ttc ttc ctc ctg ctg gtg gca gct ccc aga tgg          48
Met Lys His Leu Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp
1               5                   10                  15

gtc ctg tcc cag gtg caa ttg gtg gaa agc ggc ggc ggc ctg gtg caa          96
Val Leu Ser Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln
            20                  25                  30

ccg ggc ggc agc ctg cgt ctg agc tgc gcg gcc tcc gga ttt acc ttt         144
Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe
        35                  40                  45

aat tct tat tct atg tct tgg gtg cgc caa gcc cct ggg aag ggt ctc         192
Asn Ser Tyr Ser Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu
    50                  55                  60

gag tgg gtg agc tat atc tct tct cgt tct agc act acc tat tat gcg         240
Glu Trp Val Ser Tyr Ile Ser Ser Arg Ser Ser Thr Thr Tyr Tyr Ala
65                  70                  75                  80

gat agc gtg aaa ggc cgt ttt acc att tca cgt gat aat tcg aaa aac         288
Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn
                85                  90                  95

acc ctg tat ctg caa atg aac agc ctg cgt gcg gaa gat acg gcc gtg         336
Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val
            100                 105                 110
```

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```
tat tat tgc gcg cgt ggt tat ttt cat ggt atg gat tat tgg ggc caa      384
Tyr Tyr Cys Ala Arg Gly Tyr Phe His Gly Met Asp Tyr Trp Gly Gln
        115                 120                 125

ggc acc ctg gtg acg gtt agc tca gcc tcc acc aag ggt cca tcg gtc      432
Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val
    130                 135                 140

ttc ccc ctg gca ccc tcc tcc aag agc acc tct ggg ggc aca gcg gcc      480
Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala
145                 150                 155                 160

ctg ggc tgc ctg gtc aag gac tac ttc ccc gaa ccg gtg acg gtg tcg      528
Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser
                165                 170                 175

tgg aac tca ggc gcc ctg acc agc ggc gtg cac acc ttc ccg gct gtc      576
Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val
            180                 185                 190

cta cag tcc tca gga ctc tac tcc ctc agc agc gtg gtg acc gtg ccc      624
Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro
        195                 200                 205

tcc agc agc ttg ggc acc cag acc tac atc tgc aac gtg aat cac aag      672
Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys
    210                 215                 220

ccc agc aac acc aag gtg gac aag aga gtt gag ccc aaa tct tgt gac      720
Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp
225                 230                 235                 240

aaa act cac aca tgc cca ccg tgc cca gca cct gaa ctc ctg ggg gga      768
Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly
                245                 250                 255

ccg tca gtc ttc ctc ttc ccc cca aaa ccc aag gac acc ctc atg atc      816
Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile
            260                 265                 270

tcc cgg acc cct gag gtc aca tgc gtg gtg gtg gac gtg agc cac gaa      864
Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu
        275                 280                 285

gac cct gag gtc aag ttc aac tgg tac gtg gac ggc gtg gag gtg cat      912
Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His
    290                 295                 300

aat gcc aag aca aag ccg cgg gag gag cag tac aac agc acg tac cgg      960
Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg
305                 310                 315                 320

gtg gtc agc gtc ctc acc gtc ctg cac cag gac tgg ctg aat ggc aag     1008
Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys
                325                 330                 335

gag tac aag tgc aag gtc tcc aac aaa gcc ctc cca gcc ccc atc gag     1056
Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu
            340                 345                 350

aaa acc atc tcc aaa gcc aaa ggg cag ccc cga gaa cca cag gtg tac     1104
Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr
        355                 360                 365

acc ctg ccc cca tcc cgg gag gag atg acc aag aac cag gtc agc ctg     1152
Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu
    370                 375                 380
```

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```
acc tgc ctg gtc aaa ggc ttc tat ccc agc gac atc gcc gtg gag tgg      1200
Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp
385                 390                 395                 400

gag agc aat ggg cag ccg gag aac aac tac aag acc acg cct ccc gtg      1248
Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val
                405                 410                 415

ctg gac tcc gac ggc tcc ttc ttc ctc tac agc aag ctc acc gtg gac      1296
Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp
            420                 425                 430

aag agc agg tgg cag cag ggg aac gtc ttc tca tgc tcc gtg atg cat      1344
Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His
        435                 440                 445

gag gct ctg cac aac cac tac acg cag aag agc ctc tcc ctg tct ccg      1392
Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro
    450                 455                 460

ggt aaa tga                                                         1401
Gly Lys
465
```

<210> 51
<211> 466
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 51

```
Met Lys His Leu Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp
1               5                   10                  15

Val Leu Ser Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln
            20                  25                  30

Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe
        35                  40                  45

Asn Ser Tyr Ser Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu
    50                  55                  60

Glu Trp Val Ser Tyr Ile Ser Ser Arg Ser Ser Thr Thr Tyr Tyr Ala
65                  70                  75                  80

Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn
                85                  90                  95

Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val
            100                 105                 110

Tyr Tyr Cys Ala Arg Gly Tyr Phe His Gly Met Asp Tyr Trp Gly Gln
        115                 120                 125

Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val
```

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130 135 140

Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala
145 150 155 160

Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser
165 170 175

Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val
180 185 190

Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro
195 200 205

Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys
210 215 220

Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp
225 230 235 240

Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly
245 250 255

Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile
260 265 270

Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu
275 280 285

Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His
290 295 300

Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg
305 310 315 320

Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys
325 330 335

Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu
340 345 350

Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr
355 360 365

Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu
370 375 380

Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp
385 390 395 400

Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val

405 410 415

Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp
420 425 430

Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His
435 440 445

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

Gly Lys
465

<210> 52
<211> 702
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07919 light chain

<220>
<221> CDS
<222> (1)..(702)

<400> 52

```
atg gcc tgg gct ctg ctg ctc ctc acc ctc ctc act cag ggc aca gga          48
Met Ala Trp Ala Leu Leu Leu Leu Thr Leu Leu Thr Gln Gly Thr Gly
1               5                   10                  15

tcc tgg gct gat atc gaa ctg acc cag ccg cct tca gtg agc gtt gca          96
Ser Trp Ala Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala
            20                  25                  30

cca ggt cag acc gcg cgt atc tcg tgt agc ggc gat tct ctt ggt tct         144
Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser
        35                  40                  45

aag tat gtt cat tgg tac cag cag aaa ccc ggg cag gcg cca gtt ctt         192
Lys Tyr Val His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu
    50                  55                  60

gtg att tat cgt gat aat aag cgt ccc tca ggc atc ccg gaa cgc ttt         240
Val Ile Tyr Arg Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe
65                  70                  75                  80

agc gga tcc aac agc ggc aac acc gcg acc ctg acc att agc ggc act         288
Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr
                85                  90                  95

cag gcg gaa gac gaa gcg gat tat tat tgc cag tct tat gat gct act         336
Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr
            100                 105                 110

gag ttt act tat gtg ttt ggc ggc ggc acg aag tta acc gtc cta ggt         384
Glu Phe Thr Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly
        115                 120                 125

cag ccc aag gct gcc ccc tcg gtc act ctg ttc ccg ccc tcc tct gag         432
Gln Pro Lys Ala Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser Glu
    130                 135                 140
```

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```
gag ctt caa gcc aac aag gcc aca ctg gtg tgt ctc ata agt gac ttc        480
Glu Leu Gln Ala Asn Lys Ala Thr Leu Val Cys Leu Ile Ser Asp Phe
145                 150                 155                 160

tac ccg gga gcc gtg aca gtg gcc tgg aag gca gat agc agc ccc gtc        528
Tyr Pro Gly Ala Val Thr Val Ala Trp Lys Ala Asp Ser Ser Pro Val
                165                 170                 175

aag gcg gga gtg gag acc acc aca ccc tcc aaa caa agc aac aac aag        576
Lys Ala Gly Val Glu Thr Thr Thr Pro Ser Lys Gln Ser Asn Asn Lys
            180                 185                 190

tac gcg gcc agc agc tat ctg agc ctg acg cct gag cag tgg aag tcc        624
Tyr Ala Ala Ser Ser Tyr Leu Ser Leu Thr Pro Glu Gln Trp Lys Ser
        195                 200                 205

cac aga agc tac agc tgc cag gtc acg cat gaa ggg agc acc gtg gag        672
His Arg Ser Tyr Ser Cys Gln Val Thr His Glu Gly Ser Thr Val Glu
    210                 215                 220

aag aca gtg gcc cct aca gaa tgt tca tag                                702
Lys Thr Val Ala Pro Thr Glu Cys Ser
225                 230
```

<210> 53
<211> 233
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 53

```
Met Ala Trp Ala Leu Leu Leu Leu Thr Leu Leu Thr Gln Gly Thr Gly
1               5                   10                  15

Ser Trp Ala Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala
            20                  25                  30

Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser
        35                  40                  45

Lys Tyr Val His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu
    50                  55                  60

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

Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr
                85                  90                  95

Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr
            100                 105                 110

Glu Phe Thr Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly
        115                 120                 125
```

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Gln Pro Lys Ala Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser Glu
130 135 140

Glu Leu Gln Ala Asn Lys Ala Thr Leu Val Cys Leu Ile Ser Asp Phe
145 150 155 160

Tyr Pro Gly Ala Val Thr Val Ala Trp Lys Ala Asp Ser Ser Pro Val
165 170 175

Lys Ala Gly Val Glu Thr Thr Thr Pro Ser Lys Gln Ser Asn Asn Lys
180 185 190

Tyr Ala Ala Ser Ser Tyr Leu Ser Leu Thr Pro Glu Gln Trp Lys Ser
195 200 205

His Arg Ser Tyr Ser Cys Gln Val Thr His Glu Gly Ser Thr Val Glu
210 215 220

Lys Thr Val Ala Pro Thr Glu Cys Ser
225 230

<210> 54
<211> 708
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07692 light chain

<220>
<221> CDS
<222> (1)..(708)

<400> 54

```
atg gtg ttg cag acc cag gtc ttc att tct ctg ttg ctc tgg atc tct          48
Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

ggt gcc tac ggg gat atc gtg ctg acc cag agc ccg gcg acc ctg agc          96
Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

ctg tct ccg ggc gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct         144
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

gtt tct ttt gat tat ctg ggt tgg tac cag cag aaa cca ggt caa gca         192
Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

ccg cgt cta tta att tat ggt gct tct aat cgt gca act ggg gtc ccg         240
Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

gcg cgt ttt agc ggc tct gga tcc ggc acg gat ttt acc ctg acc att         288
Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

agc agc ctg gaa cct gaa gac ttt gcg act tat tat tgc cag cag tat         336
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
```

```
            100                 105                 110

tat aat atg cct tat acc ttt ggc cag ggt acg aaa gtt gaa att aaa        384
Tyr Asn Met Pro Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

cgt acg gtg gct gca cca tct gtc ttc atc ttc ccg cca tct gat gag        432
Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

cag ttg aaa tct gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc        480
Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

tat ccc aga gag gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa        528
Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

tcg ggt aac tcc cag gag agt gtc aca gag cag gac agc aag gac agc        576
Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

acc tac agc ctc agc agc acc ctg acg ctg agc aaa gca gac tac gag        624
Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

aaa cac aaa gtc tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg        672
Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

ccc gtc aca aag agc ttc aac agg gga gag tgt tag                       708
Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  55
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  55

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15


Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30


Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45


Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60


Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80


Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95
```

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```
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
            100                 105                 110

Tyr Asn Met Pro Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  56
<211>  708
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07923 light chain


<220>
<221>  CDS
<222>  (1)..(708)

<400>  56
atg gtg ttg cag acc cag gtc ttc att tct ctg ttg ctc tgg atc tct          48
Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

ggt gcc tac ggg gat atc gtg ctg acc cag agc ccg gcg acc ctg agc          96
Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

ctg tct ccg ggc gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct         144
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

gtt tct ttt gat tat ctg ggt tgg tac cag cag aaa cca ggt caa gca         192
Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

ccg cgt cta tta att tat ggt gct tct aat cgt gca act ggg gtc ccg         240
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```
Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

gcg cgt ttt agc ggc tct gga tcc ggc acg gat ttt acc ctg acc att        288
Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

agc agc ctg gaa cct gaa gac ttt gcg acc tat tat tgc ttt cag tat        336
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr
            100                 105                 110

ctt att gtt cct ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa        384
Leu Ile Val Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

cgt acg gtg gct gca cca tct gtc ttc atc ttc ccg cca tct gat gag        432
Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

cag ttg aaa tct gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc        480
Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

tat ccc aga gag gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa        528
Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

tcg ggt aac tcc cag gag agt gtc aca gag cag gac agc aag gac agc        576
Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

acc tac agc ctc agc agc acc ctg acg ctg agc aaa gca gac tac gag        624
Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

aaa cac aaa gtc tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg        672
Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

ccc gtc aca aag agc ttc aac agg gga gag tgt tag                        708
Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  57
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  57

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15


Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30


Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45


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

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr
            100                 105                 110

Leu Ile Val Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  58
<211>  708
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07924 light chain


<220>
<221>  CDS
<222>  (1)..(708)

<400>  58
atg gtg ttg cag acc cag gtc ttc att tct ctg ttg ctc tgg atc tct        48
Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

ggt gcc tac ggg gat atc gtg ctg acc cag agc ccg gcg acc ctg agc        96
Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30
```

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2010214301

```
ctg tct ccg ggc gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct        144
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

gtt tct ttt gat tat ctg ggt tgg tac cag cag aaa cca ggt caa gca        192
Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

ccg cgt cta tta att tat ggt gct tct aat cgt gca act ggg gtc ccg        240
Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

gcg cgt ttt agc ggc tct gga tcc ggc acg gat ttt acc ctg acc att        288
Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

agc agc ctg gaa cct gaa gac ttt gcg acc tat tat tgc cag cag tat        336
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
            100                 105                 110

aat att aat cct ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa        384
Asn Ile Asn Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

cgt acg gtg gct gca cca tct gtc ttc atc ttc ccg cca tct gat gag        432
Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

cag ttg aaa tct gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc        480
Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

tat ccc aga gag gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa        528
Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

tcg ggt aac tcc cag gag agt gtc aca gag cag gac agc aag gac agc        576
Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

acc tac agc ctc agc agc acc ctg acg ctg agc aaa gca gac tac gag        624
Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

aaa cac aaa gtc tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg        672
Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

ccc gtc aca aag agc ttc aac agg gga gag tgt tag                       708
Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  59
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  59

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15


Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
```

20 25 30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
35 40 45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
50 55 60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65 70 75 80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
85 90 95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
100 105 110

Asn Ile Asn Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225 230 235

<210> 60
<211> 708
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07925 light chain

<220>
<221> CDS
<222> (1)..(708)

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<400> 60

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atg gtg ttg cag acc cag gtc ttc att tct ctg ttg ctc tgg atc tct        48
Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                  10                  15

ggt gcc tac ggg gat atc gtg ctg acc cag agc ccg gcg acc ctg agc        96
Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

ctg tct ccg ggc gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct       144
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

gtt tct ttt gat tat ctg ggt tgg tac cag cag aaa cca ggt caa gca       192
Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

ccg cgt cta tta att tat ggt gct tct aat cgt gca act ggg gtc ccg       240
Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

gcg cgt ttt agc ggc tct gga tcc ggc acg gat ttt acc ctg acc att       288
Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

agc agc ctg gaa cct gaa gac ttt gcg acc tat tat tgc ctt cag tat       336
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr
            100                 105                 110

ttt aat cct cct cat acc ttt ggc cag ggt acg aaa gtt gaa att aaa       384
Phe Asn Pro Pro His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

cgt acg gtg gct gca cca tct gtc ttc atc ttc ccg cca tct gat gag       432
Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

cag ttg aaa tct gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc       480
Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

tat ccc aga gag gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa       528
Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

tcg ggt aac tcc cag gag agt gtc aca gag cag gac agc aag gac agc       576
Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

acc tac agc ctc agc agc acc ctg acg ctg agc aaa gca gac tac gag       624
Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

aaa cac aaa gtc tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg       672
Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

ccc gtc aca aag agc ttc aac agg gga gag tgt tag                       708
Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235
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<210> 61
<211> 235
<212> PRT
<213> Artificial Sequence

<220>

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<223> Synthetic Construct

<400> 61

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1 5 10 15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
20 25 30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
35 40 45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
50 55 60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65 70 75 80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
85 90 95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr
100 105 110

Phe Asn Pro Pro His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225 230 235

<210> 62
<211> 708
<212> DNA
<213> Artificial Sequence

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<220>
<223> MOR07926 light chain

<220>
<221> CDS
<222> (1)..(708)

<400> 62

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atg gtg ttg cag acc cag gtc ttc att tct ctg ttg ctc tgg atc tct          48
Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                  10                  15

ggt gcc tac ggg gat atc gtg ctg acc cag agc ccg gcg acc ctg agc          96
Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

ctg tct ccg ggc gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct         144
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

gtt tct ttt gat tat ctg ggt tgg tac cag cag aaa cca ggt caa gca         192
Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

ccg cgt cta tta att tat ggt gct tct aat cgt gca act ggg gtc ccg         240
Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

gcg cgt ttt agc ggc tct gga tcc ggc acg gat ttt acc ctg acc att         288
Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

agc agc ctg gaa cct gaa gac ttt gcg acc tat tat tgc ttt cag gct         336
Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala
            100                 105                 110

ctt att atg cct ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa         384
Leu Ile Met Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

cgt acg gtg gct gca cca tct gtc ttc atc ttc ccg cca tct gat gag         432
Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

cag ttg aaa tct gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc         480
Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

tat ccc aga gag gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa         528
Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

tcg ggt aac tcc cag gag agt gtc aca gag cag gac agc aag gac agc         576
Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

acc tac agc ctc agc agc acc ctg acg ctg agc aaa gca gac tac gag         624
Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

aaa cac aaa gtc tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg         672
Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

ccc gtc aca aag agc ttc aac agg gga gag tgt tag                         708
Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
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225 230 235

<210> 63
<211> 235
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 63

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1 5 10 15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
20 25 30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
35 40 45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
50 55 60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65 70 75 80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
85 90 95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala
100 105 110

Leu Ile Met Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

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Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  64
<211>  1551
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR7919 heavy chain


<220>
<221>  CDS
<222>  (45)..(1457)

<400>  64
taatacgact cactataggg agacccaagc tggctagcgc cacc atg aaa cac ctg        56
                                                 Met Lys His Leu
                                                 1

tgg ttc ttc ctc ctg ctg gtg gca gct ccc aga tgg gtc ctg tcc cag        104
Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp Val Leu Ser Gln
5                   10                  15                  20

gtg caa ttg gtt cag agc ggc gcg gaa gtg aaa aaa ccg ggc gaa agc        152
Val Gln Leu Val Gln Ser Gly Ala Glu Val Lys Lys Pro Gly Glu Ser
                25                  30                  35

ctg aaa att agc tgc aaa ggt tcc gga tat tcc ttt act aat tat tgg        200
Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe Thr Asn Tyr Trp
            40                  45                  50

att tct tgg gtg cgc cag atg cct ggg aag ggt ctc gag tgg atg ggc        248
Ile Ser Trp Val Arg Gln Met Pro Gly Lys Gly Leu Glu Trp Met Gly
        55                  60                  65

ttt atc tat ccg gat gat agc tat acc cgt tat tct ccg agc ttt cag        296
Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser Pro Ser Phe Gln
    70                  75                  80

ggc cag gtg acc att agc gcg gat aaa agc att agc acc gcg tat ctt        344
Gly Gln Val Thr Ile Ser Ala Asp Lys Ser Ile Ser Thr Ala Tyr Leu
85                  90                  95                  100

caa tgg agc agc ctg aaa gcg agc gat acg gcc atg tat tat tgc gcg        392
Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met Tyr Tyr Cys Ala
                105                 110                 115

cgt ttt tct tat cgt cat tat ctt gat atg gat gat cat tgg ggc caa        440
Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp His Trp Gly Gln
            120                 125                 130

ggc acc ctg gtg acg gtt agc tca gcc tcc acc aag ggt cca tcg gtc        488
Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val
        135                 140                 145

ttc ccc ctg gca ccc tcc tcc aag agc acc tct ggg ggc aca gcg gcc        536
Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala
    150                 155                 160

ctg ggc tgc ctg gtc aag gac tac ttc ccc gaa ccg gtg acg gtg tcg        584
Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser
165                 170                 175                 180

tgg aac tca ggc gcc ctg acc agc ggc gtg cac acc ttc ccg gct gtc        632
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Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val
                185                 190                 195

cta cag tcc tca gga ctc tac tcc ctc agc agc gtg gtg acc gtg ccc      680
Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro
            200                 205                 210

tcc agc agc ttg ggc acc cag acc tac atc tgc aac gtg aat cac aag      728
Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys
        215                 220                 225

ccc agc aac acc aag gtg gac aag aga gtt gag ccc aaa tct tgt gac      776
Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp
    230                 235                 240

aaa act cac aca tgc cca ccg tgc cca gca cct gaa ctc ctg ggg gga      824
Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly
245                 250                 255                 260

ccg tca gtc ttc ctc ttc ccc cca aaa ccc aag gac acc ctc atg atc      872
Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile
                265                 270                 275

tcc cgg acc cct gag gtc aca tgc gtg gtg gtg gac gtg agc cac gaa      920
Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu
            280                 285                 290

gac cct gag gtc aag ttc aac tgg tac gtg gac ggc gtg gag gtg cat      968
Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His
        295                 300                 305

aat gcc aag aca aag ccg cgg gag gag cag tac aac agc acg tac cgg     1016
Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg
    310                 315                 320

gtg gtc agc gtc ctc acc gtc ctg cac cag gac tgg ctg aat ggc aag     1064
Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys
325                 330                 335                 340

gag tac aag tgc aag gtc tcc aac aaa gcc ctc cca gcc ccc atc gag     1112
Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu
                345                 350                 355

aaa acc atc tcc aaa gcc aaa ggg cag ccc cga gaa cca cag gtg tac     1160
Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr
            360                 365                 370

acc ctg ccc cca tcc cgg gag gag atg acc aag aac cag gtc agc ctg     1208
Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu
        375                 380                 385

acc tgc ctg gtc aaa ggc ttc tat ccc agc gac atc gcc gtg gag tgg     1256
Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp
    390                 395                 400

gag agc aat ggg cag ccg gag aac aac tac aag acc acg cct ccc gtg     1304
Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val
405                 410                 415                 420

ctg gac tcc gac ggc tcc ttc ttc ctc tac agc aag ctc acc gtg gac     1352
Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp
                425                 430                 435

aag agc agg tgg cag cag ggg aac gtc ttc tca tgc tcc gtg atg cat     1400
Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His
            440                 445                 450

gag gct ctg cac aac cac tac acg cag aag agc ctc tcc ctg tct ccg     1448
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Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro
455 460 465

ggt aaa tga gggcccgttt aaacgggtgg catccctgtg acccctcccc 1497
Gly Lys
470

agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg tcct 1551

<210> 65
<211> 470
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 65

Met Lys His Leu Trp Phe Phe Leu Leu Val Ala Ala Pro Arg Trp
1 5 10 15

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

Pro Gly Glu Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe
35 40 45

Thr Asn Tyr Trp Ile Ser Trp Val Arg Gln Met Pro Gly Lys Gly Leu
50 55 60

Glu Trp Met Gly Phe Ile Tyr Pro Asp Asp Ser Tyr Thr Arg Tyr Ser
65 70 75 80

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

Thr Ala Tyr Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met
100 105 110

Tyr Tyr Cys Ala Arg Phe Ser Tyr Arg His Tyr Leu Asp Met Asp Asp
115 120 125

His Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys
130 135 140

Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly
145 150 155 160

Gly Thr Ala Ala Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro
165 170 175

Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr
180 185 190

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Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val
        195                 200                 205

Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn
    210                 215                 220

Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro
225                 230                 235                 240

Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu
                245                 250                 255

Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp
            260                 265                 270

Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp
        275                 280                 285

Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly
    290                 295                 300

Val Glu Val His Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn
305                 310                 315                 320

Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp
                325                 330                 335

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

Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu
        355                 360                 365

Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn
    370                 375                 380

Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile
385                 390                 395                 400

Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr
                405                 410                 415

Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys
            420                 425                 430

Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys
        435                 440                 445

Ser Val Met His Glu Ala Leu His Asn His Tyr Thr Gln Lys Ser Leu
    450                 455                 460
```

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Ser Leu Ser Pro Gly Lys
465                 470


<210>  66
<211>  1539
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR7692, MOR7923, MOR7924, MOR7925 and MOR7926 heavy chain


<220>
<221>  CDS
<222>  (45)..(1445)

<400>  66
taatacgact cactataggg agacccaagc tggctagcgc cacc atg aaa cac ctg        56
                                                Met Lys His Leu
                                                1

tgg ttc ttc ctc ctg ctg gtg gca gct ccc aga tgg gtc ctg tcc cag        104
Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp Val Leu Ser Gln
5                   10                  15                  20

gtg caa ttg gtg gaa agc ggc ggc ggc ctg gtg caa ccg ggc ggc agc        152
Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln Pro Gly Gly Ser
                25                  30                  35

ctg cgt ctg agc tgc gcg gcc tcc gga ttt acc ttt aat tct tat tct        200
Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe Asn Ser Tyr Ser
            40                  45                  50

atg tct tgg gtg cgc caa gcc cct ggg aag ggt ctc gag tgg gtg agc        248
Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu Glu Trp Val Ser
        55                  60                  65

tat atc tct tct cgt tct agc act acc tat tat gcg gat agc gtg aaa        296
Tyr Ile Ser Ser Arg Ser Ser Thr Thr Tyr Tyr Ala Asp Ser Val Lys
    70                  75                  80

ggc cgt ttt acc att tca cgt gat aat tcg aaa aac acc ctg tat ctg        344
Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn Thr Leu Tyr Leu
85                  90                  95                  100

caa atg aac agc ctg cgt gcg gaa gat acg gcc gtg tat tat tgc gcg        392
Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val Tyr Tyr Cys Ala
                105                 110                 115

cgt ggt tat ttt cat ggt atg gat tat tgg ggc caa ggc acc ctg gtg        440
Arg Gly Tyr Phe His Gly Met Asp Tyr Trp Gly Gln Gly Thr Leu Val
            120                 125                 130

acg gtt agc tca gcc tcc acc aag ggt cca tcg gtc ttc ccc ctg gca        488
Thr Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val Phe Pro Leu Ala
        135                 140                 145

ccc tcc tcc aag agc acc tct ggg ggc aca gcg gcc ctg ggc tgc ctg        536
Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala Leu Gly Cys Leu
    150                 155                 160

gtc aag gac tac ttc ccc gaa ccg gtg acg gtg tcg tgg aac tca ggc        584
Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Trp Asn Ser Gly
165                 170                 175                 180

gcc ctg acc agc ggc gtg cac acc ttc ccg gct gtc cta cag tcc tca        632
Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val Leu Gln Ser Ser
```

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                185                 190                 195

gga ctc tac tcc ctc agc agc gtg gtg acc gtg ccc tcc agc agc ttg        680
Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro Ser Ser Ser Leu
            200                 205                 210

ggc acc cag acc tac atc tgc aac gtg aat cac aag ccc agc aac acc        728
Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys Pro Ser Asn Thr
        215                 220                 225

aag gtg gac aag aga gtt gag ccc aaa tct tgt gac aaa act cac aca        776
Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp Lys Thr His Thr
    230                 235                 240

tgc cca ccg tgc cca gca cct gaa ctc ctg ggg gga ccg tca gtc ttc        824
Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser Val Phe
245                 250                 255                 260

ctc ttc ccc cca aaa ccc aag gac acc ctc atg atc tcc cgg acc cct        872
Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro
                265                 270                 275

gag gtc aca tgc gtg gtg gtg gac gtg agc cac gaa gac cct gag gtc        920
Glu Val Thr Cys Val Val Val Asp Val Ser His Glu Asp Pro Glu Val
            280                 285                 290

aag ttc aac tgg tac gtg gac ggc gtg gag gtg cat aat gcc aag aca        968
Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His Asn Ala Lys Thr
        295                 300                 305

aag ccg cgg gag gag cag tac aac agc acg tac cgg gtg gtc agc gtc       1016
Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val
    310                 315                 320

ctc acc gtc ctg cac cag gac tgg ctg aat ggc aag gag tac aag tgc       1064
Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys
325                 330                 335                 340

aag gtc tcc aac aaa gcc ctc cca gcc ccc atc gag aaa acc atc tcc       1112
Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser
                345                 350                 355

aaa gcc aaa ggg cag ccc cga gaa cca cag gtg tac acc ctg ccc cca       1160
Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro
            360                 365                 370

tcc cgg gag gag atg acc aag aac cag gtc agc ctg acc tgc ctg gtc       1208
Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val
        375                 380                 385

aaa ggc ttc tat ccc agc gac atc gcc gtg gag tgg gag agc aat ggg       1256
Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly
    390                 395                 400

cag ccg gag aac aac tac aag acc acg cct ccc gtg ctg gac tcc gac       1304
Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp
405                 410                 415                 420

ggc tcc ttc ttc ctc tac agc aag ctc acc gtg gac aag agc agg tgg       1352
Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp
                425                 430                 435

cag cag ggg aac gtc ttc tca tgc tcc gtg atg cat gag gct ctg cac       1400
Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala Leu His
            440                 445                 450

aac cac tac acg cag aag agc ctc tcc ctg tct ccg ggt aaa tga           1445
Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly Lys
```

455 460 465

gggcccgttt aaacgggtgg catccctgtg acccctcccc agtgcctctc ctggccctgg 1505

aagttgccac tccagtgccc accagccttg tcct 1539

<210> 67
<211> 466
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 67

Met Lys His Leu Trp Phe Phe Leu Leu Leu Val Ala Ala Pro Arg Trp
1 5 10 15

Val Leu Ser Gln Val Gln Leu Val Glu Ser Gly Gly Gly Leu Val Gln
20 25 30

Pro Gly Gly Ser Leu Arg Leu Ser Cys Ala Ala Ser Gly Phe Thr Phe
35 40 45

Asn Ser Tyr Ser Met Ser Trp Val Arg Gln Ala Pro Gly Lys Gly Leu
50 55 60

Glu Trp Val Ser Tyr Ile Ser Ser Arg Ser Ser Thr Thr Tyr Tyr Ala
65 70 75 80

Asp Ser Val Lys Gly Arg Phe Thr Ile Ser Arg Asp Asn Ser Lys Asn
85 90 95

Thr Leu Tyr Leu Gln Met Asn Ser Leu Arg Ala Glu Asp Thr Ala Val
100 105 110

Tyr Tyr Cys Ala Arg Gly Tyr Phe His Gly Met Asp Tyr Trp Gly Gln
115 120 125

Gly Thr Leu Val Thr Val Ser Ser Ala Ser Thr Lys Gly Pro Ser Val
130 135 140

Phe Pro Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala
145 150 155 160

Leu Gly Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser
165 170 175

Trp Asn Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val
180 185 190

Leu Gln Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro
195 200 205

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```
Ser Ser Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys
    210                 215                 220

Pro Ser Asn Thr Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp
225                 230                 235                 240

Lys Thr His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly
                245                 250                 255

Pro Ser Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile
            260                 265                 270

Ser Arg Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu
        275                 280                 285

Asp Pro Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His
    290                 295                 300

Asn Ala Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg
305                 310                 315                 320

Val Val Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys
                325                 330                 335

Glu Tyr Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu
            340                 345                 350

Lys Thr Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr
        355                 360                 365

Thr Leu Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu
    370                 375                 380

Thr Cys Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp
385                 390                 395                 400

Glu Ser Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val
                405                 410                 415

Leu Asp Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp
            420                 425                 430

Lys Ser Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His
        435                 440                 445

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

Gly Lys
465
```

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<210> 68
<211> 840
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07919 lambda light chain

<220>
<221> CDS
<222> (45)..(746)

<400> 68
taatacgact cactataggg agacccaagc tggctagcgc cacc atg gcc tgg gct 56
Met Ala Trp Ala
1

ctg ctg ctc ctc acc ctc ctc act cag ggc aca gga tcc tgg gct gat 104
Leu Leu Leu Leu Thr Leu Leu Thr Gln Gly Thr Gly Ser Trp Ala Asp
5 10 15 20

atc gaa ctg acc cag ccg cct tca gtg agc gtt gca cca ggt cag acc 152
Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala Pro Gly Gln Thr
25 30 35

gcg cgt atc tcg tgt agc ggc gat tct ctt ggt tct aag tat gtt cat 200
Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser Lys Tyr Val His
40 45 50

tgg tac cag cag aaa ccc ggg cag gcg cca gtt ctt gtg att tat cgt 248
Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu Val Ile Tyr Arg
55 60 65

gat aat aag cgt ccc tca ggc atc ccg gaa cgc ttt agc gga tcc aac 296
Asp Asn Lys Arg Pro Ser Gly Ile Pro Glu Arg Phe Ser Gly Ser Asn
70 75 80

agc ggc aac acc gcg acc ctg acc att agc ggc act cag gcg gaa gac 344
Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu Asp
85 90 95 100

gaa gcg gat tat tat tgc cag tct tat gat gct act gag ttt act tat 392
Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr Glu Phe Thr Tyr
105 110 115

gtg ttt ggc ggc ggc acg aag tta acc gtc cta ggt cag ccc aag gct 440
Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln Pro Lys Ala
120 125 130

gcc ccc tcg gtc act ctg ttc ccg ccc tcc tct gag gag ctt caa gcc 488
Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser Glu Glu Leu Gln Ala
135 140 145

aac aag gcc aca ctg gtg tgt ctc ata agt gac ttc tac ccg gga gcc 536
Asn Lys Ala Thr Leu Val Cys Leu Ile Ser Asp Phe Tyr Pro Gly Ala
150 155 160

gtg aca gtg gcc tgg aag gca gat agc agc ccc gtc aag gcg gga gtg 584
Val Thr Val Ala Trp Lys Ala Asp Ser Ser Pro Val Lys Ala Gly Val
165 170 175 180

gag acc acc aca ccc tcc aaa caa agc aac aac aag tac gcg gcc agc 632
Glu Thr Thr Thr Pro Ser Lys Gln Ser Asn Asn Lys Tyr Ala Ala Ser
185 190 195

agc tat ctg agc ctg acg cct gag cag tgg aag tcc cac aga agc tac 680

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```
Ser Tyr Leu Ser Leu Thr Pro Glu Gln Trp Lys Ser His Arg Ser Tyr
            200                 205                 210

agc tgc cag gtc acg cat gaa ggg agc acc gtg gag aag aca gtg gcc      728
Ser Cys Gln Val Thr His Glu Gly Ser Thr Val Glu Lys Thr Val Ala
        215                 220                 225

cct aca gaa tgt tca tag gggcccgttt aaacgggtgg catccctgtg             776
Pro Thr Glu Cys Ser
    230

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg    836

tcct                                                                 840


<210>  69
<211>  233
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  69

Met Ala Trp Ala Leu Leu Leu Leu Thr Leu Leu Thr Gln Gly Thr Gly
1               5                   10                  15

Ser Trp Ala Asp Ile Glu Leu Thr Gln Pro Pro Ser Val Ser Val Ala
            20                  25                  30

Pro Gly Gln Thr Ala Arg Ile Ser Cys Ser Gly Asp Ser Leu Gly Ser
        35                  40                  45

Lys Tyr Val His Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Leu
    50                  55                  60

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

Ser Gly Ser Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr
                85                  90                  95

Gln Ala Glu Asp Glu Ala Asp Tyr Tyr Cys Gln Ser Tyr Asp Ala Thr
            100                 105                 110

Glu Phe Thr Tyr Val Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly
        115                 120                 125

Gln Pro Lys Ala Ala Pro Ser Val Thr Leu Phe Pro Pro Ser Ser Glu
    130                 135                 140

Glu Leu Gln Ala Asn Lys Ala Thr Leu Val Cys Leu Ile Ser Asp Phe
145                 150                 155                 160

Tyr Pro Gly Ala Val Thr Val Ala Trp Lys Ala Asp Ser Ser Pro Val
                165                 170                 175
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Lys Ala Gly Val Glu Thr Thr Thr Pro Ser Lys Gln Ser Asn Asn Lys
            180                 185                 190

Tyr Ala Ala Ser Ser Tyr Leu Ser Leu Thr Pro Glu Gln Trp Lys Ser
        195                 200                 205

His Arg Ser Tyr Ser Cys Gln Val Thr His Glu Gly Ser Thr Val Glu
    210                 215                 220

Lys Thr Val Ala Pro Thr Glu Cys Ser
225                 230

<210>  70
<211>  846
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07692 kappa light chain

<220>
<221>  CDS
<222>  (45)..(752)

<400>  70
taatacgact cactataggg agacccaagc tggctagcgc cacc atg gtg ttg cag        56
                                                Met Val Leu Gln
                                                1

acc cag gtc ttc att tct ctg ttg ctc tgg atc tct ggt gcc tac ggg        104
Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser Gly Ala Tyr Gly
5                   10                  15                  20

gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc        152
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
                25                  30                  35

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat        200
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            40                  45                  50

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta        248
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        55                  60                  65

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc        296
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    70                  75                  80

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa        344
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
85                  90                  95                  100

cct gaa gac ttt gcg act tat tat tgc cag cag tat tat aat atg cct        392
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Tyr Asn Met Pro
                105                 110                 115

tat acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg gtg gct        440
Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
            120                 125                 130
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gca cca tct gtc ttc atc ttc ccg cca tct gat gag cag ttg aaa tct        488
Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
        135                 140                 145

gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc tat ccc aga gag        536
Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
    150                 155                 160

gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa tcg ggt aac tcc        584
Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
165                 170                 175                 180

cag gag agt gtc aca gag cag gac agc aag gac agc acc tac agc ctc        632
Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
                185                 190                 195

agc agc acc ctg acg ctg agc aaa gca gac tac gag aaa cac aaa gtc        680
Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
            200                 205                 210

tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg ccc gtc aca aag        728
Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
        215                 220                 225

agc ttc aac agg gga gag tgt tag gggcccgttt aaacgggtgg catccctgtg        782
Ser Phe Asn Arg Gly Glu Cys
    230                 235

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg        842

tcct                                                                    846


<210>  71
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  71

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
```

100 105 110

Tyr Asn Met Pro Tyr Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225 230 235

<210> 72
<211> 846
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07923 kappa light chain

<220>
<221> CDS
<222> (45)..(752)

<400> 72
taatacgact cactataggg agacccaagc tggctagcgc cacc atg gtg ttg cag 56
Met Val Leu Gln
1

acc cag gtc ttc att tct ctg ttg ctc tgg atc tct ggt gcc tac ggg 104
Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser Gly Ala Tyr Gly
5 10 15 20

gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc 152
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
25 30 35

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat 200
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
40 45 50

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta 248
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
55 60 65

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att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc        296
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    70                  75                  80

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa        344
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
85                  90                  95                  100

cct gaa gac ttt gcg acc tat tat tgc ttt cag tat ctt att gtt cct        392
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr Leu Ile Val Pro
                105                 110                 115

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg gtg gct        440
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
            120                 125                 130

gca cca tct gtc ttc atc ttc ccg cca tct gat gag cag ttg aaa tct        488
Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
        135                 140                 145

gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc tat ccc aga gag        536
Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
    150                 155                 160

gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa tcg ggt aac tcc        584
Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
165                 170                 175                 180

cag gag agt gtc aca gag cag gac agc aag gac agc acc tac agc ctc        632
Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
                185                 190                 195

agc agc acc ctg acg ctg agc aaa gca gac tac gag aaa cac aaa gtc        680
Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
            200                 205                 210

tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg ccc gtc aca aag        728
Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
        215                 220                 225

agc ttc aac agg gga gag tgt tag gggcccgttt aaacgggtgg catccctgtg        782
Ser Phe Asn Arg Gly Glu Cys
    230                 235

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg       842

tcct                                                                   846


<210>  73
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  73

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15


Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30
```

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```
Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Tyr
            100                 105                 110

Leu Ile Val Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  74
<211>  846
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07924 kappa light chain


<220>
<221>  CDS
<222>  (45)..(752)

<400>  74
taatacgact cactataggg agacccaagc tggctagcgc cacc atg gtg ttg cag        56
                                                Met Val Leu Gln
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acc cag gtc ttc att tct ctg ttg ctc tgg atc tct ggt gcc tac ggg        104
Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser Gly Ala Tyr Gly
5                   10                  15                  20

gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc        152
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
                25                  30                  35

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat        200
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            40                  45                  50

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta        248
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        55                  60                  65

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc        296
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    70                  75                  80

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa        344
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
85                  90                  95                  100

cct gaa gac ttt gcg acc tat tat tgc cag cag tat aat att aat cct        392
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr Asn Ile Asn Pro
                105                 110                 115

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg gtg gct        440
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
            120                 125                 130

gca cca tct gtc ttc atc ttc ccg cca tct gat gag cag ttg aaa tct        488
Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
        135                 140                 145

gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc tat ccc aga gag        536
Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
    150                 155                 160

gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa tcg ggt aac tcc        584
Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
165                 170                 175                 180

cag gag agt gtc aca gag cag gac agc aag gac agc acc tac agc ctc        632
Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
                185                 190                 195

agc agc acc ctg acg ctg agc aaa gca gac tac gag aaa cac aaa gtc        680
Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
            200                 205                 210

tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg ccc gtc aca aag        728
Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
        215                 220                 225

agc ttc aac agg gga gag tgt tag gggcccgttt aaacgggtgg catccctgtg        782
Ser Phe Asn Arg Gly Glu Cys
    230                 235

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg        842

tcct                                                                    846
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<210> 75

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<211> 235
<212> PRT
<213> Artificial Sequence

<220>
<223> Synthetic Construct

<400> 75

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1 5 10 15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
20 25 30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
35 40 45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
50 55 60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65 70 75 80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
85 90 95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Gln Gln Tyr
100 105 110

Asn Ile Asn Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225 230 235

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<210> 76
<211> 846
<212> DNA
<213> Artificial Sequence

<220>
<223> MOR07925 kappa light chain

<220>
<221> CDS
<222> (45)..(752)

<400> 76
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Met Val Leu Gln
1

acc cag gtc ttc att tct ctg ttg ctc tgg atc tct ggt gcc tac ggg 104
Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser Gly Ala Tyr Gly
5 10 15 20

gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc 152
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
25 30 35

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat 200
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
40 45 50

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta 248
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
55 60 65

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc 296
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
70 75 80

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa 344
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
85 90 95 100

cct gaa gac ttt gcg acc tat tat tgc ctt cag tat ttt aat cct cct 392
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr Phe Asn Pro Pro
105 110 115

cat acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg gtg gct 440
His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
120 125 130

gca cca tct gtc ttc atc ttc ccg cca tct gat gag cag ttg aaa tct 488
Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
135 140 145

gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc tat ccc aga gag 536
Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
150 155 160

gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa tcg ggt aac tcc 584
Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
165 170 175 180

cag gag agt gtc aca gag cag gac agc aag gac agc acc tac agc ctc 632
Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
185 190 195

agc agc acc ctg acg ctg agc aaa gca gac tac gag aaa cac aaa gtc 680

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Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
            200                 205                 210

tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg ccc gtc aca aag        728
Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
        215                 220                 225

agc ttc aac agg gga gag tgt tag gggcccgttt aaacgggtgg catccctgtg       782
Ser Phe Asn Arg Gly Glu Cys
    230                 235

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg      842

tcct                                                                   846


<210>  77
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  77

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Leu Gln Tyr
            100                 105                 110

Phe Asn Pro Pro His Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
        115                 120                 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
    130                 135                 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145                 150                 155                 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
                165                 170                 175
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Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
            180                 185                 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
        195                 200                 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
    210                 215                 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225                 230                 235


<210>  78
<211>  846
<212>  DNA
<213>  Artificial Sequence

<220>
<223>  MOR07926 kappa light chain


<220>
<221>  CDS
<222>  (45)..(752)

<400>  78
taatacgact cactataggg agacccaagc tggctagcgc cacc atg gtg ttg cag        56
                                                Met Val Leu Gln
                                                1

acc cag gtc ttc att tct ctg ttg ctc tgg atc tct ggt gcc tac ggg        104
Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser Gly Ala Tyr Gly
5                   10                  15                  20

gat atc gtg ctg acc cag agc ccg gcg acc ctg agc ctg tct ccg ggc        152
Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser Leu Ser Pro Gly
                25                  30                  35

gaa cgt gcg acc ctg agc tgc aga gcg agc cag tct gtt tct ttt gat        200
Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Ser Phe Asp
            40                  45                  50

tat ctg ggt tgg tac cag cag aaa cca ggt caa gca ccg cgt cta tta        248
Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
        55                  60                  65

att tat ggt gct tct aat cgt gca act ggg gtc ccg gcg cgt ttt agc        296
Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro Ala Arg Phe Ser
    70                  75                  80

ggc tct gga tcc ggc acg gat ttt acc ctg acc att agc agc ctg gaa        344
Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
85                  90                  95                  100

cct gaa gac ttt gcg acc tat tat tgc ttt cag gct ctt att atg cct        392
Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala Leu Ile Met Pro
                105                 110                 115

ttt acc ttt ggc cag ggt acg aaa gtt gaa att aaa cgt acg gtg gct        440
Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
            120                 125                 130
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gca cca tct gtc ttc atc ttc ccg cca tct gat gag cag ttg aaa tct        488
Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
        135                 140                 145

gga act gcc tct gtt gtg tgc ctg ctg aat aac ttc tat ccc aga gag        536
Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
    150                 155                 160

gcc aaa gta cag tgg aag gtg gat aac gcc ctc caa tcg ggt aac tcc        584
Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
165                 170                 175                 180

cag gag agt gtc aca gag cag gac agc aag gac agc acc tac agc ctc        632
Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
                185                 190                 195

agc agc acc ctg acg ctg agc aaa gca gac tac gag aaa cac aaa gtc        680
Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
            200                 205                 210

tac gcc tgc gaa gtc acc cat cag ggc ctg agc tcg ccc gtc aca aag        728
Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
        215                 220                 225

agc ttc aac agg gga gag tgt tag gggcccgttt aaacgggtgg catccctgtg        782
Ser Phe Asn Arg Gly Glu Cys
    230                 235

acccctcccc agtgcctctc ctggccctgg aagttgccac tccagtgccc accagccttg        842

tcct                                                                    846


<210>  79
<211>  235
<212>  PRT
<213>  Artificial Sequence

<220>
<223>  Synthetic Construct

<400>  79

Met Val Leu Gln Thr Gln Val Phe Ile Ser Leu Leu Leu Trp Ile Ser
1               5                   10                  15

Gly Ala Tyr Gly Asp Ile Val Leu Thr Gln Ser Pro Ala Thr Leu Ser
            20                  25                  30

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
        35                  40                  45

Val Ser Phe Asp Tyr Leu Gly Trp Tyr Gln Gln Lys Pro Gly Gln Ala
    50                  55                  60

Pro Arg Leu Leu Ile Tyr Gly Ala Ser Asn Arg Ala Thr Gly Val Pro
65                  70                  75                  80

Ala Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
                85                  90                  95

Ser Ser Leu Glu Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Phe Gln Ala
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100 105 110

Leu Ile Met Pro Phe Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
115 120 125

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
130 135 140

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
145 150 155 160

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
165 170 175

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
180 185 190

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
195 200 205

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
210 215 220

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
225 230 235

<210> 80
<211> 127
<212> PRT
<213> Artificial Sequence

<220>
<223> Variable heavy chain 3

<220>
<221> misc_feature
<222> (30)..(35)
<223> Xaa can be any naturally occurring amino acid

<220>
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<222> (50)..(66)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
<222> (99)..(116)
<223> Xaa can be any naturally occurring amino acid

<400> 80

Gln 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 Xaa Xaa Xaa
20 25 30

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

Ser Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
50 55 60

Xaa Xaa 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 Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
100 105 110

Xaa Xaa Xaa Xaa Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
115 120 125

<210> 81
<211> 127
<212> PRT
<213> Artificial Sequence

<220>
<223> Variable heavy chain 5

<220>
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<222> (30)..(35)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
<222> (50)..(66)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
<222> (99)..(116)
<223> Xaa can be any naturally occurring amino acid

<400> 81

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

Ser Leu Lys Ile Ser Cys Lys Gly Ser Gly Tyr Ser Phe Xaa Xaa Xaa
20 25 30

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

Gly Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
50 55 60

Xaa Xaa Gln Val Thr Ile Ser Ala Asp Lys Ser Ile Ser Thr Ala Tyr

65 70 75 80

Leu Gln Trp Ser Ser Leu Lys Ala Ser Asp Thr Ala Met Tyr Tyr Cys
85 90 95

Ala Arg Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
100 105 110

Xaa Xaa Xaa Xaa Trp Gly Gln Gly Thr Leu Val Thr Val Ser Ser
115 120 125

<210> 82
<211> 110
<212> PRT
<213> Artificial Sequence

<220>
<223> Variable light chain keppa 3

<220>
<221> misc_feature
<222> (24)..(35)
<223> Xaa can be any naturally occurring amino acid

<220>
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<222> (50)..(57)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> VARIANT
<222> (86)..(86)
<223> T

<220>
<221> misc_feature
<222> (90)..(98)
<223> Xaa can be any naturally occurring amino acid

<400> 82

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

Glu Arg Ala Thr Leu Ser Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
20 25 30

Xaa Xaa Xaa Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
35 40 45

Ile Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Gly Val Pro Ala Arg Phe Ser
50 55 60

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Ser Leu Glu
65 70 75 80

Pro Glu Asp Phe Ala Thr Tyr Tyr Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa
85 90 95

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Xaa Xaa Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
100 105 110

<210> 83
<211> 110
<212> PRT
<213> Artificial Sequence

<220>
<223> Variable light chain lambda

<220>
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<222> (23)..(33)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
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<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
<222> (48)..(55)
<223> Xaa can be any naturally occurring amino acid

<220>
<221> misc_feature
<222> (88)..(98)
<223> Xaa can be any naturally occurring amino acid

<400> 83

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

Thr Ala Arg Ile Ser Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
20 25 30

Xaa Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Val Xaa Val Ile Xaa
35 40 45

Xaa Xaa Xaa Xaa Xaa Xaa Xaa Gly Ile Pro Glu Arg Phe Ser Gly Ser
50 55 60

Asn Ser Gly Asn Thr Ala Thr Leu Thr Ile Ser Gly Thr Gln Ala Glu
65 70 75 80

Asp Glu Ala Asp Tyr Tyr Cys Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa Xaa
85 90 95

Xaa Xaa Phe Gly Gly Gly Thr Lys Leu Thr Val Leu Gly Gln
100 105 110