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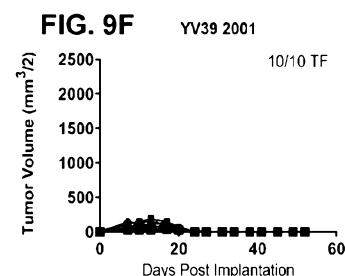
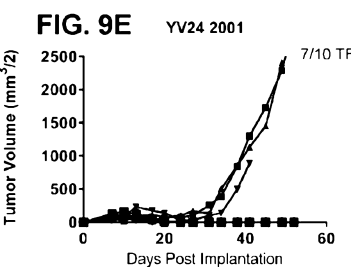
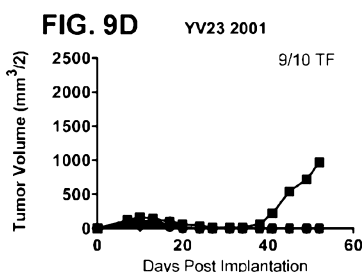
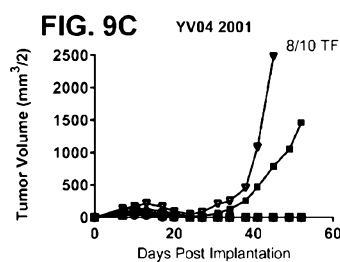
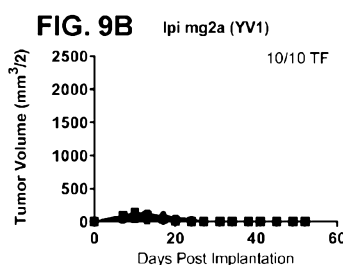
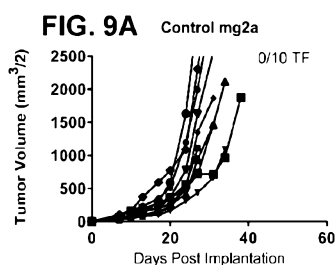
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(54) Title: ACTIVATABLE ANTI-CTLA-4 ANTIBODIES AND USES THEREOF



(57) **Abstract:** Provided herein are activatable anti-human CTLA-4 antibodies comprising a heavy chain comprising a VH domain and a light chain comprising a masking moiety (MM), a cleavable moiety (CM), and a VL domain. Such activatable anti-human CTLA-4 antibodies have CTLA-4 binding activity in the tumor microenvironment, where the masking moiety is removed by proteolytic cleavage of the cleavable moiety by tumor-specific proteases, but exhibit greatly reduced binding to CTLA-4 outside the tumor. In this way, the activatable anti-human CTLA-4 antibodies of the present invention retain anti-tumor activity while reducing the side effects associated with anti-CTLA-4 activity outside the tumor.

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ACTIVATABLE ANTI-CTLA-4 ANTIBODIES AND USES THEREOF

CROSS REFERENCE TO RELATED APPLICATIONS

- [0001] This application claims the priority benefit of U.S. Provisional Application No. 62/417,212, filed November 3, 2016, which is hereby incorporated by reference in its entirety.

REFERENCE TO SEQUENCE LISTING
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- [0002] The content of the electronically submitted sequence listing (Name: 3338_059PC02_SeqListing.txt; Size: 527,968 bytes; and Date of Creation: October 27, 2017) is herein incorporated by reference in its entirety.

BACKGROUND OF THE INVENTION

- [0003] The immune system is capable of controlling tumor development and mediating tumor regression. This requires the generation and activation of tumor antigen-specific T cells. Multiple T-cell co-stimulatory receptors and T-cell negative regulators, or co-inhibitory receptors, act in concert to control T-cell activation, proliferation, and gain or loss of effector function. Among the earliest and best-characterized T-cell co-stimulatory and co-inhibitory molecules are CD28 and CTLA-4. Rudd *et al.* (2009) *Immunol. Rev.* 229: 12. CD28 provides co-stimulatory signals to T-cell receptor engagement by binding to B7-1 and B7-2 ligands on antigen-presenting cells, while CTLA-4 provides a negative signal down-regulating T-cell proliferation and function. CTLA-4, which also binds the B7-1 (CD80) and B7-2 (CD86) ligands but with higher affinity than CD28, acts as a negative regulator of T-cell function through both cell autonomous (or intrinsic) and cell non-autonomous (or extrinsic) pathways. Intrinsic control of CD8 and CD4 T effector (T_{eff}) function is mediated by the inducible surface expression of CTLA-4 as a result of T-cell activation, and inhibition of T-cell proliferation and cytokine proliferation by multivalent engagement of B7 ligands on opposing cells. Peggs *et al.* (2008) *Immunol. Rev.* 224:141.

[0004] Anti-CTLA-4 antibodies, when cross-linked, suppress T cell function *in vitro*. Krummel & Allison (1995) *J. Exp. Med.* 182:459; Walunas *et al.* (1994) *Immunity* 1:405. Regulatory T cells (T_{regs}), which express CTLA-4 constitutively, control effector T cell (T_{eff}) function in a non-cell autonomous fashion. T_{regs} that are deficient for CTLA-4 have impaired suppressive ability (Wing *et al.* (2008) *Science* 322:271) and antibodies that block CTLA-4 interaction with B7 can inhibit T_{reg} function (Read *et al.* (2000) *J. Exp. Med.* 192:295; Quezada *et al.* (2006) *J. Clin. Invest.* 116:1935). More recently, T_{effs} have also been shown to control T cell function through extrinsic pathways (Corse & Allison (2012) *J. Immunol.* 189:1123; Wang *et al.* (2012) *J. Immunol.* 189:1118). Extrinsic control of T cell function by T_{regs} and T_{effs} occurs through the ability of CTLA-4-positive cells to remove B7 ligands on antigen-presenting cells, thereby limiting their co-stimulatory potential. Qureshi *et al.* (2011) *Science* 332: 600; Onishi *et al.* (2008) *Proc. Nat'l Acad. Sci. (USA)* 105:10113. Antibody blockade of CTLA-4/B7 interactions is thought to promote T_{eff} activation by interfering with negative signals transmitted by CTLA-4 engagement; this intrinsic control of T-cell activation and proliferation can promote both T_{eff} and T_{reg} proliferation (Krummel & Allison (1995) *J. Exp. Med.* 182:459; Quezada *et al.* (2006) *J. Clin. Invest.* 116:1935). In early studies with animal models, antibody blockade of CTLA-4 was shown to exacerbate autoimmunity. Perrin *et al.* (1996) *J. Immunol.* 157:1333; Hurwitz *et al.* (1997) *J. Neuroimmunol.* 73:57. By extension to tumor immunity, the ability of anti-CTLA-4 to cause regression of established tumors provided a dramatic example of the therapeutic potential of CTLA-4 blockade. Leach *et al.* (1996) *Science* 271:1734.

[0005] Human antibodies to human CTLA-4, ipilimumab and tremelimumab, were selected to inhibit CTLA-4-B7 interactions (Keler *et al.* (2003) *J. Immunol.* 171:6251; Ribas *et al.* (2007) *Oncologist* 12:873) and have been tested in a variety of clinical trials for multiple malignancies. Hoos *et al.* (2010) *Semin. Oncol.* 37:533; Ascierto *et al.* (2011) *J. Transl. Med.* 9:196. Tumor regressions and disease stabilization were frequently observed, and treatment with these antibodies has been accompanied by adverse events with inflammatory infiltrates capable of affecting a variety of organ systems. In 2011, ipilimumab, which has an IgG1 constant region, was approved in the US and EU for the treatment of unresectable or metastatic melanoma based on an improvement in overall

survival in a phase III trial of previously treated patients with advanced melanoma. Hodi *et al.* (2010) *N. Engl. J. Med.* 363:711.

[0006] Treatment with ipilimumab has, however, been hampered by dose limiting toxicities, such as colitis. Di Giacomo *et al.* (2010) *Seminars in Oncology* 37:499. Accordingly, the need exists for improved anti-CTLA-4 antibodies, such as modified forms of ipilimumab, with reduced toxicity but with comparable anti-tumor efficacy. Such improved anti-CTLA-4 antibodies may be more effective anti-tumor agents than current antibodies.

SUMMARY OF THE INVENTION

[0007] Provided herein are activatable anti-human CTLA-4 antibodies comprising a heavy chain comprising a VH domain and a light chain comprising a masking moiety (MM), a cleavable moiety (CM), and a VL domain. Such activatable anti-human CTLA-4 antibodies have CTLA-4 binding activity in the tumor microenvironment, where the masking moiety is removed by proteolytic cleavage of the cleavable moiety by tumor-specific proteases, but exhibit greatly reduced binding to CTLA-4 outside the tumor. In this way, the activatable anti-human CTLA-4 antibodies of the present invention retain anti-tumor activity while reducing the side effects associated with anti-CTLA-4 activity outside the tumor.

[0008] Provided herein are improved anti-CTLA-4 antibodies, such as an improved ipilimumab, in particular an activatable antibody that when activated binds Cytotoxic T-Lymphocyte Antigen 4 (CTLA-4). In some embodiments, the activatable anti-human CTLA-4 antibody comprises:

(i) a heavy chain comprising a heavy chain variable domain (VH) comprising complementarity determining regions (CDRs) CDRH1: SYTMH (SEQ ID NO: 557); CDRH2: FISYDGNKYYADSVKG (SEQ ID NO: 558); and CDRH3: TGWLGPFDY (SEQ ID NO: 559); and

(ii) a light chain comprising:

- (a) a light chain variable domain (VL) comprising CDRL1: RASQSVGSSYLA (SEQ ID NO: 560); CDRL2: GAFSRAT (SEQ ID NO: 561); and CDRL3: QQYGSSPWT (SEQ ID NO: 562);
- (b) a cleavable moiety (CM); and

(c) a masking moiety (MM),

wherein the light chain has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-VL.

[0009] In some embodiments, an activatable anti-human CTLA-4 antibody comprises:

(i) a heavy chain comprising a heavy chain variable domain (VH) comprising CDRH1: SYTMH (SEQ ID NO: 557); CDRH2: FISYDGNNKYYADSVKG (SEQ ID NO: 558); and CDRH3: TGWLGPFDY (SEQ ID NO: 559); and

(ii) a light chain comprising, from N-terminus to C-Terminus:

(a) a masking moiety (MM);

(b) a cleavable moiety (CM); and

(c) a light chain variable domain (VL) comprising CDRL1: RASQSVGSSYLA (SEQ ID NO: 560); CDRL2: GAFSRAT (SEQ ID NO: 561); and CDRL3: QQYGSSPWT (SEQ ID NO: 562).

[0010] In some embodiments, the activatable antibody comprises a heavy chain and a light chain such that the light chain has the structural arrangement, from N-terminus to C-terminus of the light chain, MM-CM-VL. As used herein, the N-terminal fragment that is joined to the VL domain is referred to as the prodomain and comprises MM and CM.

[0011] In some embodiments, the activatable antibody comprises a complete antibody, i.e., an antibody comprising two mature full-length heavy chains and two mature full-length light chains. In some embodiments, the activatable antibody comprises a Fab fragment, a F(ab')₂ fragment, an scFv, or a scAb. In some embodiments, the activatable antibody comprises a monoclonal antibody.

[0012] In some embodiments, the CM functions as a substrate for a protease. In some embodiments, the CM is selected from the group of CMs provided in Table 3. In some embodiments, the CM is selected from the group consisting of 2001 (SEQ ID NO: 297), 2003 (SEQ ID NO: 298), 2005 (SEQ ID NO: 299), 2006 (SEQ ID NO: 300), 2007 (SEQ ID NO: 301), 2008 (SEQ ID NO: 302), 2009 (SEQ ID NO: 303), 2011 (SEQ ID NO: 304), 2012 (SEQ ID NO: 305), 3001 (SEQ ID NO: 306), 3006 (SEQ ID NO: 307), 3007 (SEQ ID NO: 308), 3008 (SEQ ID NO: 309), 3009 (SEQ ID NO: 310), 3011 (SEQ ID NO: 311), and 3012 (SEQ ID NO: 312). In some embodiments, the CM is 2001 (SEQ ID NO: 297). In some embodiments, the CM is 2011 (SEQ ID NO: 304). In some embodiments, the CM is 2012 (SEQ ID NO: 305).

- [0013]** In some embodiments, the MM is selected from the group consisting of the MMs provided in Tables 4-6. In some embodiments, the MM is selected from the group consisting of YV01 (SEQ ID NO: 1), YV02 (SEQ ID NO: 2), YV03, (SEQ ID NO: 3), YV04 (SEQ ID NO: 4), YV09, (SEQ ID NO: 9), YV23 (SEQ ID NO: 23), YV24 (SEQ ID NO: 24), YV35 (SEQ ID NO: 35), YV39 (SEQ ID NO: 39), YV51 (SEQ ID NO: 51), YV61 (SEQ ID NO: 60), YV62 (SEQ ID NO: 61), YV63 (SEQ ID NO: 62), YV64 (SEQ ID NO: 63), YV65 (SEQ ID NO: 64), and YV66 (SEQ ID NO: 65); and the CM is selected from the group consisting of 2001, 2006, 2007, 2008, 2009, 2011, and 2012. In some embodiments, the MM is YV39 and the CM is 2011. In some embodiments, the MM is YV39 and the CM is 2012. In some embodiments, the MM is YV39 and the CM is 2001.
- [0014]** In some embodiments, the activatable antibody comprises a heavy chain comprising the amino acid sequence of SEQ ID NO: 353 and a light chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 356 to 529. In some embodiments, the activatable anti-CTLA-4 antibodies comprise a light chain having a prodomain and VL corresponding to the prodomain and VL of SEQ ID NOs: 356 to 529. In some embodiments, the activatable anti-CTLA-4 antibodies comprise a light chain having a prodomain and VL of SEQ ID NOs: 564, 565, or 563. In one embodiment, the activatable anti-CTLA-4 antibody comprises a light chain having a prodomain and VL of SEQ ID NO: 564.
- [0015]** In some embodiments, the activatable anti-CTLA-4 antibodies comprise a heavy chain variable domain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to SEQ ID NO: 345. In some embodiments, the activatable anti-CTLA-4 antibodies comprise a light chain variable domain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 564, 565, and 563.
- [0016]** In some embodiments, the activatable antibody comprises a combination of heavy chain sequence SEQ ID NO: 353 and light chain sequence SEQ ID NO: 449, 473, or 383. In some embodiments, the activatable antibody comprises a combination of heavy chain sequence SEQ ID NO: 349 and light chain sequence SEQ ID NO: 448, 472, or 382.

- [0017] Provided herein is an activatable anti-CTLA-4 antibody that, when activated, specifically binds to human CTLA-4 and is referred to as an activated activatable anti-CTLA-4 antibody. In some embodiments, the activated activatable anti-CTLA-4 antibody binds to CTLA-4 with the same binding affinity as ipilimumab. Also provided herein is an activatable anti-CTLA-4 antibody that does not bind to CTLA-4 as effectively as ipilimumab since the activatable anti-CTLA-4 antibody comprises a heavy chain and a light chain comprising a prodomain comprising a MM and CM linked to the ipilimumab light chain such that the prodomain reduces the ability of the ipilimumab to bind to CTLA-4.
- [0018] In some embodiments, the activatable antibody binds to human CTLA-4 with an EC_{50} of 1 $\mu\text{g/mL}$ or higher as measured by flow cytometry. In some embodiments, the activatable anti-CTLA-4 antibodies bind to CTLA-4 with an EC_{50} of 5 $\mu\text{g/mL}$ or higher, 10 $\mu\text{g/mL}$ or higher, 20 $\mu\text{g/mL}$ or higher, or 40 $\mu\text{g/mL}$ or higher.
- [0019] In some embodiments, the MM is a polypeptide of no more than 40 amino acids in length. In some embodiments, the MM is a polypeptide that is no more than 50% identical to any natural binding partner of the antibody. In some embodiments, the MM does not comprise more than 25% amino acid sequence identity to CTLA-4. In some embodiments, the MM does not comprise more than 10% amino acid sequence identity to CTLA-4.
- [0020] Activatable anti-CTLA-4 antibodies of the disclosure are activated when the cleavable moiety is cleaved by a protease. In some embodiments, the protease is produced by a tumor that is in proximity to T cells that express CTLA-4. In some embodiments, the protease is produced by a tumor that is co-localized with T cells that express CTLA-4. In some embodiments, the protease is selected from the group of proteases provided in Table 1 provided below. In some embodiments, the protease is selected from the group consisting of a matrix metalloprotease (MMP), a thrombin, a neutrophil elastase, a cysteine protease, a legumain, and a serine protease, such as a matriptase or a urokinase (uPA). In some embodiments, the protease is selected from the group consisting of MMP1, MMP2, MMP3, MMP8, MMP9, MMP11, MMP13, MMP14, MMP17, legumain, matriptase, and uPA, or a combination of one or more of such proteases. In some embodiments, the CM is cleaved by a matrix metalloprotease (MMP) and a serine

protease. In some embodiments, the CM is cleaved by a matrix metalloprotease (MMP), a serine protease and a legumain.

Table 1: Exemplary Proteases and/or Enzymes

ADAMS, ADAMTS, <i>e.g.</i> ADAM8 ADAM9 ADAM10 ADAM12 ADAM15 ADAM17/TACE	Cysteine proteinases, <i>e.g.</i> , Cruzipain Legumain Otubain-2	Serine proteases, <i>e.g.</i> , activated protein C Cathepsin A Cathepsin G Chymase coagulation factor proteases (<i>e.g.</i> , FVIIa, FIXa, FXa, FXIa, FXIIa)
ADAMDEC1 ADAMTS1 ADAMTS4 ADAMTS5	KLKs, <i>e.g.</i> , KLK4	
	KLK5	Elastase
	KLK6	Granzyme B
	KLK7	Guanidinobenzoatase
	KLK8	HtrA1
	KLK10	Human Neutrophil Elastase
Aspartate proteases, <i>e.g.</i> , BACE Renin	KLK11 KLK13 KLK14	Lactoferrin
		Marapsin
Aspartic cathepsins, <i>e.g.</i> , Cathepsin D Cathepsin E	Metallo proteinases, <i>e.g.</i> , Meprin Neprilysin PSMA BMP-1	NS3/4A PACE4 Plasmin PSA tPA Thrombin Trypsin uPA
		Type II Transmembrane Serine Proteases (TTSPs), <i>e.g.</i> , DESC1 DPP-4 FAP Hepsin Matriptase-2 MT-SP1/Matriptase TMPRSS2 TMPRSS3 TMPRSS4
Caspases, <i>e.g.</i> , Caspase 1 Caspase 2 Caspase 3 Caspase 4 Caspase 5	MMPs, <i>e.g.</i> , MMP1 MMP2 MMP3	
Caspase 6 Caspase 7 Caspase 8 Caspase 9 Caspase 10 Caspase 14	MMP7 MMP8 MMP9 MMP10 MMP11 MMP12 MMP13	
Cysteine cathepsins, <i>e.g.</i> , Cathepsin B Cathepsin C Cathepsin K Cathepsin L Cathepsin S Cathepsin V/L2	MMP14 MMP15 MMP16 MMP17 MMP19 MMP20 MMP23	

Cathepsin X/Z/P	MMP24
	MMP26
	MMP27

[0021] Provided herein are activatable anti-CTLA-4 antibodies that further comprise one or more linker peptides. In some embodiments, the linker peptide is between the MM and the CM. In some embodiments, the linker peptide is between the CM and the VL. In some embodiments, the activatable antibody comprises a first linker peptide (LP1) and a second linker peptide (LP2). In some embodiments, the activatable antibody comprises a heavy chain and a light chain such that the light chain has the structural arrangement, from N-terminus to C-terminus of the light chain, MM-LP1-CM-LP2-VL. In some embodiments, the LP1 and the LP2 are not identical to each other. In some embodiments, the LP1 and the LP2 are identical to each other. In some embodiments, the prodomain comprises MM-LP1-CM-LP2.

[0022] In some embodiments, the LP1 and/or the LP2 comprise a glycine-serine polymer. In some embodiments, the LP1 and/or the LP2 comprise an amino acid sequence selected from the group consisting of (GS)_n (SEQ ID NO: 532), (GGS)_n (SEQ ID NO: 533), (GSGGS)_n (SEQ ID NO: 534), and (GGGS)_n (SEQ ID NO: 535), where n is an integer of at least one. In some embodiments, the LP1 comprises the amino acid sequence GGGSSGGS (SEQ ID NO: 542). In some embodiments, the LP2 comprises the amino acid sequence GGGS (SEQ ID NO: 543).

[0023] Provided herein are activatable anti-CTLA-4 antibodies that also comprise a spacer. In some embodiments, the spacer is joined directly to the MM and has the structural arrangement from N-terminus to C-terminus as follows: spacer-MM-CM-VL. In some embodiments, the spacer comprises an amino acid sequence selected from the group consisting of QGQSGQG (SEQ ID NO: 544), GQSGQG (SEQ ID NO: 545), QGQSGS (SEQ ID NO: 546), QGQSGQ (SEQ ID NO: 547), QSGQG (SEQ ID NO: 548), GQSGS (SEQ ID NO: 549), QGQSG (SEQ ID NO: 550), SGQG (SEQ ID NO: 551), QSGS (SEQ ID NO: 552), QGQS (SEQ ID NO: 553), QQG, SGS, QQG, QG, GS, G, S, and Q. In some embodiments, the spacer and the MM comprise the amino acid sequence QGQSGSCRTQLYGYNLCPY (SEQ ID NO: 556).

[0024] Also provided herein are activatable antibodies that comprise a toxic agent, such as a dolastatin, an auristatin, an auristatin E, a monomethyl auristatin E (MMAE), a maytansinoid, a duocarmycin, a calicheamicin, a pyrrolobenzodiazepine, or a derivative

thereof. In some embodiments, the toxic agent is conjugated to the activatable antibody via a linker. In some embodiments, the linker is a cleavable linker. In some embodiments, the linker is a non-cleavable linker.

[0025] Provided herein are activatable anti-CTLA-4 antibodies that comprises a detectable moiety. In some embodiments, the detectable moiety is a diagnostic agent.

[0026] Provided herein are pharmaceutical compositions comprising an activatable anti-CTLA-4 antibody described herein. In some embodiments, the pharmaceutical composition comprises an additional therapeutic agent.

[0027] Also provided herein are isolated nucleic acid molecules encoding the heavy and/or light chains of the activatable anti-CTLA-4 antibodies described herein, vectors that comprise one or more of the isolated nucleic acid molecules, and methods of producing an activatable antibody by culturing a cell comprising the vector or vectors under conditions that lead to expression of the activatable antibody.

[0028] Provided herein are methods of manufacturing an activatable antibody, the methods comprising: (a) culturing a cell comprising a nucleic acid construct that encodes the activatable antibody described herein under conditions that lead to expression of the activatable antibody, and (b) recovering the activatable antibody.

[0029] Provided herein are methods of reducing CTLA-4 activity comprising administering an effective amount of the activatable antibody described herein or pharmaceutical compositions comprising an activatable anti-CTLA-4 antibody described herein to a subject in need thereof.

[0030] Provided herein are methods of blocking binding of a natural ligand to CTLA-4 comprising administering an effective amount of the activatable antibodies described herein or pharmaceutical compositions comprising an activatable anti-CTLA-4 antibody described herein to a subject in need thereof.

[0031] Provided herein are methods of treating, alleviating a symptom of, or delaying the progression of a CTLA-4-related disorder comprising administering a therapeutically effective amount of the activatable antibodies described herein or the pharmaceutical compositions comprising an activatable anti-CTLA-4 antibody described herein to a subject in need thereof. In some embodiments, the CTLA-4 related disorder is a cancer. In some embodiments, the cancer is a melanoma, such as unresectable or metastatic melanoma, breast cancer, colorectal cancer, gastric cancer, glioblastoma, head and neck

cancer, lung cancer, ovarian cancer, endometrial cancer, pancreatic cancer, prostate cancer, renal cancer, sarcoma, or skin cancer. In some embodiments, the CTLA-4 related disorder is a disorder known to be treatable with ipilimumab.

[0032] Where aspects or embodiments of the invention are described in terms of a Markush group or other grouping of alternatives, the present invention encompasses not only the entire group listed as a whole, but also each member of the group individually and all possible subgroups of the main group, and also the main group absent one or more of the group members. The present invention also envisages the explicit exclusion of one or more of any of the group members in the claimed invention.

BRIEF DESCRIPTION OF THE DRAWINGS/FIGURES

[0033] FIGs. 1A to 1C show tumor volumes as a function of days post tumor implantation in mice (n = 10) treated with (i) an unrelated mouse IgG2a antibody (FIG. 1A), (ii) a mouse anti-CTLA-4 (9D9) IgG2a antibody (FIG. 2B), or (iii) an activatable 9D9 antibody (FIG. 1C). All antibodies and activatable antibodies were dosed at 25 µg/mouse. The activatable 9D9 antibody comprises MY11 (SEQ ID NO: 294) as the masking moiety and 2001 (SEQ ID NO: 297) as the cleavable moiety. "TF" indicates the number of tumor free mice at the end of each experiment. The unrelated mouse IgG2a antibody and the mouse anti-CTLA-4 (9D9) IgG2a antibody were used as controls.

[0034] FIGs. 2A to 2C show the frequency of regulatory T cells in the tumor (FIG. 2A) and proliferation and activation of regulatory T cells in the spleen (FIGs. 2B and 2C) of mice treated with different activatable mouse anti-CTLA-4 (9D9) IgG2a antibodies. The different activatable 9D9 antibodies comprise (i) either MY03 (SEQ ID NO: 293) or MY11 (SEQ ID NO: 294) as the masking moiety and (2) 0003 (SEQ ID NO: 320), 1004 (SEQ ID NO: 323), or 2001 (SEQ ID NO: 297) as the cleavable moiety. The unrelated mouse IgG2a antibody ("DT 1D12 mg2a") and the mouse anti-CTLA-4 (9D9) IgG2a antibody ("9D9 mg2a") were used as controls. In FIG. 2A, the frequency of regulatory T cells is shown as a percentage of total CD4⁺ T cells that are Foxp3⁺ in the tumor. FIGs. 2B and 2C show the frequency of proliferating (Ki-67⁺) and activated (ICOS⁺) regulatory T cells, as a percentage of Foxp3⁺ T cells, in the spleen, respectively.

[0035] FIGs. 3A to 3E show the ability of different anti-CTLA-4 activatable antibodies (human IgG1 isotype) to bind to human CTLA-4, as measured *in vitro* with an ELISA

binding assay. Ipilimumab ("YV1") was used as a control in all experiments. In FIG. 3A, the anti-CTLA-4 activatable antibodies comprise YV04 (SEQ ID NO: 4), YV06 (SEQ ID NO: 6), YV09 (SEQ ID NO: 9), or YV23 (SEQ ID NO: 23) as the masking moiety. In FIG. 3B, the anti-CTLA-4 activatable antibodies comprise YV27 (SEQ ID NO: 27), YV29 (SEQ ID NO: 29), YV32 (SEQ ID NO: 32), or YV33 (SEQ ID NO: 33) as the masking moiety. In FIG. 3C, the anti-CTLA-4 activatable antibodies comprise YV35 (SEQ ID NO: 35) or YV41 (SEQ ID NO: 41) as the masking moiety. In FIG. 3D, the anti-CTLA-4 activatable antibodies comprise YV24 (SEQ ID NO: 24), YV39 (SEQ ID NO: 39), YV51 (SEQ ID NO: 51), YV52 (SEQ ID NO: 52), or YV53 (SEQ ID NO: 53) as the masking moiety. In FIG. 3E, the anti-CTLA-4 activatable antibodies comprise YV54 (SEQ ID NO: 54), YV55 (SEQ ID NO: 55), YV56 (SEQ ID NO: 56), YV57 (SEQ ID NO: 57), or YV58 (SEQ ID NO: 58) as the masking moiety. In FIGS. 3A to 3E, all the anti-CTLA-4 activatable antibodies comprise 2001 (SEQ ID NO: 297) as the cleavable moiety.

[0036] FIGS. 4A to 4D show the ability of additional anti-CTLA-4 activatable antibodies (human IgG1 isotype) to bind to human CTLA-4, as measured *in vitro* with an ELISA binding assay. Ipilimumab ("YV1") was used as a control in all experiments. In FIG. 4A, the anti-CTLA-4 activatable antibodies comprise YV04, YV06, YV09, YV23, YV27, or YV29 as the masking moiety. In FIG. 4B, the anti-CTLA-4 activatable antibodies comprise YV32, YV33, YV35, or YV41 as the masking moiety. In FIG. 4C, the anti-CTLA-4 activatable antibodies comprise YV24, YV39, YV51, YV52, or YV53 as the masking moiety. In FIG. 4D, the anti-CTLA-4 activatable antibodies comprise YV54, YV55, YV56, YV57, or YV58 as the masking moiety. In FIGS. 4A to 4D, all the anti-CTLA-4 activatable antibodies comprise 3001 as the cleavable moiety.

[0037] FIGS. 5A to 5F show the ability of several anti-CTLA-4 activatable antibodies (mouse IgG2a isotype) to bind to human CTLA-4, as measured *in vitro* with an ELISA binding assay. Ipilimumab ("YV1") was used as a control. In FIG. 5A, the anti-CTLA-4 activatable antibodies comprise YV04 as the masking moiety and 2001 (SEQ ID NO: 297), 2006 (SEQ ID NO: 300), 2007 (SEQ ID NO: 301), 2008 (SEQ ID NO: 302), or 2009 (SEQ ID NO: 303) as the cleavable moiety. In FIG. 5B, the anti-CTLA-4 activatable antibodies comprise YV04 or YV23 as the masking moiety, and 2001, 2006, 2007, 2008, or 2009 as the cleavable moiety. In FIG. 5C, the anti-CTLA-4 activatable

antibodies comprise YV39 as the masking moiety and 2001, 2006, 2008, or 2009 as the cleavable moiety. In FIG. 5D, the anti-CTLA-4 activatable antibodies comprise YV61 (SEQ ID NO: 60), YV62 (SEQ ID NO: 61), YV63 (SEQ ID NO: 62), YV64 (SEQ ID NO: 63), or YV39 (SEQ ID NO: 39) as the masking moiety and 2001 or 2012 as the cleavable moiety. In FIG. 5E, the anti-CTLA-4 activatable antibodies comprise YV65 (SEQ ID NO: 64), YV66 (SEQ ID NO: 65), YV01 (SEQ ID NO: 1), YV02 (SEQ ID NO: 2), or YV39 (SEQ ID NO: 39) as the masking moiety and 2001 or 2012 as the cleavable moiety. In FIG. 5F, the anti-CTLA-4 activatable antibodies comprise YV39 or YV03 (SEQ ID NO: 3) as the masking moiety and 2001 or 2012 as the cleavable moiety.

[0038] FIGs. 6A and 6B compares the ability of anti-CTLA-4 activatable antibodies having either a mouse IgG2a isotype (FIG. 6A) or human IgG1 isotype (FIG. 6B) to bind to human CTLA-4, as measured *in vitro* with an ELISA binding assay. Ipilimumab ("YV1") was used as a control. In both FIGs. 6A and 6B, the anti-CTLA-4 activatable antibodies comprise YV39 as the masking moiety and 2001, 2008, 2011, or 2012 as the cleavable moiety. In a modified antibody of the disclosure (YV39-NSUB), the cleavable moiety was replaced with a protease resistant linker ("NSUB") comprising the amino acid sequence GSGSGSGGGSGGGS (SEQ ID NO: 570).

[0039] FIGs. 7A to 7D show the ability of different anti-CTLA-4 activatable antibodies to bind 58 $\alpha\beta^+$ cells overexpressing human CTLA-4, as measured via flow cytometry. Binding is presented as arbitrary fluorescence units (mean fluorescence intensity, MFI, or geometric mean fluorescence intensity, gMFI) as a function of the concentration of anti-CTLA-4 antibody added. In FIG. 7A, the anti-CTLA-4 activatable antibodies comprise YV04, YV23, YV24, or YV39 as the masking moiety and 2001 as the cleavable moiety. In FIG. 7B, the anti-CTLA-4 activatable antibodies comprise YV61, YV62, YV64, or YV39 as the masking moiety and 2001 or 2011 as the cleavable moiety. In FIG. 7C, the anti-CTLA-4 activatable antibodies comprise YV39 as the masking moiety and for the cleavable moiety, 2011 ("Ipi YV39 2011") or three variants of Ipi YV39 2011: (i) mono-clipped ("Ipi YV39 MMP monoclipped"), (ii) fully clipped by MMP ("Ipi YV39 MMP"), or (iii) fully clipped by uPA ("Ipi YV39 2011 uPA"). FIG. 7D provides the EC₅₀ values for the different activatable antibodies shown in FIG. 7C. Ipilimumab was used as a control for FIGs. 7A to 7D.

- [0040] FIG. 8 shows the activity of the anti-CTLA-4 activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011") (square) at different concentrations, as measured *in vitro* with an SEB (Staphylococcal enterotoxin B) assay. Antibody activity is shown via IL-2 production by the human PBMCs after SEB stimulation. An unrelated human IgG1 isotype (triangle), ipilimumab (circle), and SEB only stimulation (x-mark) were used as controls.
- [0041] FIGs. 9A to 9F show tumor volume as a function of days post tumor implantation in human CTLA-4 knock-in mice (n = 10) treated with different anti-human CTLA-4 activatable antibodies (mouse IgG2a isotype) dosed once at 10 mg/kg. An unrelated mouse IgG2a antibody (FIG. 9A) and ipilimumab with a mouse IgG2a isotype (FIG. 9B) were used as controls. In FIGs. 9C to 9F, the activatable antibodies comprise YV04, YV23, YV24, and YV39, respectively, as the masking moiety and 2001 as the cleavable moiety.
- [0042] FIGs. 10A to 10F show tumor volume as a function of days post tumor implantation in human CTLA-4 knock-in mice (n = 10) treated with different anti-human CTLA-4 activatable antibodies (human IgG1 isotype). The antibodies were dosed once at 200 µg/mouse on day 7 post-implantation. An unrelated human IgG1 antibody (FIG. 10A) and ipilimumab with a human IgG1 isotype (FIG. 10B) were used as controls. In FIGs. 10C to 10F, the activatable antibodies comprise YV39 as the masking moiety and 2001, 2012, 2011, or 2008 as the cleavable moiety. Cleavable moieties 2012, 2011, and 2008 have been modified to overcome a deamidation site in 2001.
- [0043] FIGs. 11A to 11G show tumor volume as a function of days post tumor implantation in human CTLA-4 knock-in mice (n = 16) treated with different doses of an anti-CTLA activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011") (FIGs. 11E to 11G). The antibody was dosed once at 10 mg/kg (FIG. 11E), 3 mg/kg (FIG. 11F), or 1 mg/kg (FIG. 11G) on day 7 post tumor implantation. Control animals were treated with ipilimumab (10 mg/kg, 3 mg/kg, or 1 mg/kg; FIGs. 11B to 11D, respectively) or an unrelated human IgG1 antibody (FIG. 11A).
- [0044] FIGs. 12A to 12D show the frequency of regulatory T cells in the tumor (FIGs. 12A and 12B) or the spleen (FIGs. 12C and 12D) in human CTLA-4 knock-in mice (n = 10) treated with different anti-human CTLA-4 activatable antibodies with a mouse IgG2a

isotype. All antibodies were dosed once at 10 mg/kg. The activatable antibodies comprise YV04, YV23, YV24, or YV39 as the masking moiety and 2001 as the cleavable moiety. The labels on the abscissas of FIGs. 12C and 12D also apply to FIGs. 12A and 12B, respectively. An unrelated human IgG1 antibody and ipilimumab with a mouse IgG2a isotype were used as controls. In FIGs. 12A and 12C, the frequency of regulatory T cells is shown as a percentage of total CD4⁺ T cells that are Foxp3⁺. In FIGs. 12B and 12D, the frequency of regulatory T cells is shown as a percentage of total CD45⁺ T cells that are Foxp3⁺. FIGs. 12E and 12F show the frequency of activated (ICOS⁺) cells and proliferating (Ki-67⁺) cells is shown as a percentage of regulatory T cells in the spleen.

[0045] FIGs. 13A to 13C show the frequency of regulatory T cells in the tumor (FIGs. 13A and 13B) or the spleen (FIG. 13C) in human CTLA-4 knock-in mice treated with anti-CTLA-4 activatable antibody. The activatable antibody used comprises YV39 as the masking moiety and were either a mouse IgG2a isotype or human IgG1 isotype. An unrelated human IgG1 antibody and ipilimumab with a human IgG1 isotype were used as controls. In FIGs. 13A and 13C, the frequency of regulatory T cells is shown as a percentage of total CD4⁺ T cells that are Foxp3⁺. In FIG. 13B, the frequency of regulatory T cells is shown as a percentage of total CD45⁺ T cells that are Foxp3⁺. FIGs. 13D and 13E show the frequency of proliferating (Ki-67⁺) and activated (ICOS⁺) cells as a percentage of regulatory T cells in the spleen.

[0046] FIGs. 14A to 14C show the frequency of regulatory T cells (FIGs. 14A and 14B) or CD4⁺ effector T cells (FIG. 14C) in the tumors of mice treated with different anti-CTLA-4 activatable antibodies. FIGs. 14D and 14E show the regulatory T cells in the spleen. The anti-CTLA-4 activatable antibodies comprise YV39 as the masking moiety and 2012, 2011, 2008, or 2001 as the cleavable moiety. An unrelated human IgG1 antibody and ipilimumab with a human IgG1 isotype were used as controls. In FIGs. 14A and 14D, the frequency of regulatory T cells is shown as a percentage of total CD4⁺ T cells that are Foxp3⁺. In FIGs. 14B and 14E, the frequency of regulatory T cells is shown as a percentage of total CD45⁺ T cells that are Foxp3⁺. FIG. 14C shows the frequency of CD4⁺ effector T cells as a percentage of the total CD45⁺ T cells in the tumor. FIGs. 14F and 14G show the percentages of proliferating (Ki-67⁺) and activated (ICOS⁺) regulatory T cells in the spleen

- [0047] FIG. 15 shows the frequency of regulatory T cells in the tumors of human CTLA-4 knock-in mice ($n = 8$) treated with different doses of either ipilimumab or an anti-CTLA-4 activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011"). The antibodies were dosed once at 10 mg/kg, 3 mg/kg, or 1 mg/kg on day 7 post tumor implantation. An unrelated human IgG1 antibody was used as a control.
- [0048] FIGs. 16A and 16B show the percentages of activated (ICOS+) and proliferating (Ki-67+) regulatory T cells in the spleen of human CTLA-4 knock-in mice ($n = 8$) treated with different doses of either ipilimumab or an anti-CTLA-4 activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011"). The antibodies were dosed once at 10 mg/kg, 3 mg/kg, or 1 mg/kg on day 7 post tumor implantation. An unrelated human IgG1 antibody was used as a control.
- [0049] FIGs. 17A to 17D show tumor volume as a function of days post tumor implantation in human CTLA-4 knock-in mice ($n = 10$) treated with different doses of ipilimumab ("Ipi") (FIG. 17B), a nonfucosylated version of ipilimumab ("Ipi NF") (FIG. 17C), or a nonfucosylated version of an anti-CTLA-4 activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011 NF") (FIG. 17D). The antibodies were dosed once at 10 mg/kg, 3 mg/kg, or 1 mg/kg (left panel, middle panel, and right panel, respectively, in FIGs. 17B to 17D). Control animals received an unrelated human IgG1 antibody (FIG. 17A).
- [0050] FIG. 18 shows the frequency of regulatory T cells in the tumors of human CTLA-4 knock-in mice ($n = 5$) treated with either the nonfucosylated version of ipilimumab ("Ipi NF") or a nonfucosylated version of the anti-CTLA-4 activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("NF Ipi YV39 2011"). The antibodies were dosed once at 200 μ g/mouse on day 7 post tumor implantation. An unrelated human IgG1 antibody was used as a control.
- [0051] FIG. 19 shows the binding affinities (K_d) for both ipilimumab ("Ipi") and a nonfucosylated version of ipilimumab ("Ipi NF") to various human, cyno, and mouse Fc receptors.
- [0052] FIG. 20 shows the median percentage of Ki67+ CD4+ T cells in the blood of cynomolgus monkeys after treatment with an anti-CTLA-4 activatable antibody. The anti-

CTLA-4 activatable antibody comprises YV39 as the masking moiety and 2001 as the cleavable moiety. Vehicle and ipilimumab were used as controls.

DETAILED DESCRIPTION OF INVENTION

- [0053] In order that the present description can be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description.
- [0054] It is to be noted that the term "a" or "an" entity refers to one or more of that entity; for example, "a nucleotide sequence," is understood to represent one or more nucleotide sequences. As such, the terms "a" (or "an"), "one or more," and "at least one" can be used interchangeably herein.
- [0055] Furthermore, "and/or" where used herein is to be taken as specific disclosure of each of the two specified features or components with or without the other. Thus, the term "and/or" as used in a phrase such as "A and/or B" herein is intended to include "A and B," "A or B," "A" (alone), and "B" (alone). Likewise, the term "and/or" as used in a phrase such as "A, B, and/or C" is intended to encompass each of the following aspects: A, B, and C; A, B, or C; A or C; A or B; B or C; A and C; A and B; B and C; A (alone); B (alone); and C (alone).
- [0056] It is understood that wherever aspects are described herein with the language "comprising," otherwise analogous aspects described in terms of "consisting of" and/or "consisting essentially of" are also provided.
- [0057] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this disclosure is related. For example, the Concise Dictionary of Biomedicine and Molecular Biology, Juo, Pei-Show, 2nd ed., 2002, CRC Press; The Dictionary of Cell and Molecular Biology, 3rd ed., 1999, Academic Press; and the Oxford Dictionary Of Biochemistry And Molecular Biology, Revised, 2000, Oxford University Press, provide one of skill with a general dictionary of many of the terms used in this disclosure.
- [0058] Units, prefixes, and symbols are denoted in their Système International de Unites (SI) accepted form. Numeric ranges are inclusive of the numbers defining the range. Unless otherwise indicated, nucleotide sequences are written left to right in 5' to 3' orientation. Amino acid sequences are written left to right in amino to carboxy orientation. The headings provided herein are not limitations of the various aspects of the

disclosure, which can be had by reference to the specification as a whole. Accordingly, the terms defined immediately below are more fully defined by reference to the specification in its entirety.

- [0059]** The term "cytotoxic T-lymphocyte antigen 4" or "CTLA-4" as used herein refers to a receptor that is a member of the immunoglobulin superfamily that is expressed by activated T cells and transmits an inhibitory signal to T cells. CTLA-4 is homologous to the T-cell co-stimulatory protein, CD28, and both molecules bind to CD80 and CD86, also called B7-1 and B7-2 respectively, on antigen-presenting cells. CTLA4 is also found in regulatory T cells and contributes to its inhibitory function. CTLA-4 is also referred to as cytotoxic T-lymphocyte-associated protein 4, CD152, Insulin-dependent Diabetes Mellitus 12 (IDDM12), Celiac Disease 3 (CELIAC3), GRD4, and GSE. The term "CTLA-4" includes any variants or isoforms of CTLA-4 which are naturally expressed by cells.
- [0060]** The term "T cell" as used herein is defined as a thymus-derived lymphocyte that participates in a variety of cell-mediated immune reactions. The term "regulatory T cell" as used herein refers to a $CD4^+CD25^+FoxP3^+$ T cell with suppressive properties. "Treg" is the abbreviation used herein for a regulatory T cell.
- [0061]** The term "helper T cell" as used herein refers to a $CD4^+$ T cell; helper T cells recognize antigen bound to MHC Class II molecules. There are at least two types of helper T cells, Th1 and Th2, which produce different cytokines. Helper T cells become $CD25^+$ when activated, but only transiently become $FoxP3^+$.
- [0062]** The term "cytotoxic T cell" as used herein refers to a $CD8^+$ T cell; cytotoxic T cells recognize antigen bound to MHC Class I molecules.
- [0063]** The term "antibody" refers to immunoglobulin molecules and immunologically active portions of immunoglobulin (Ig) molecules, i.e., molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. By "specifically bind" or "immunoreacts with" or "immunospecifically bind" is meant that the antibody reacts with one or more antigenic determinants of the desired antigen and does not react with other polypeptides or binds at much lower affinity ($K_d > 10^{-6}$). Antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, domain antibody, single chain, Fab, and F(ab')₂ fragments, scFvs, and a Fab expression library.

[0064] The basic antibody structural unit is known to comprise a tetramer. Each tetramer is composed of two identical pairs of polypeptide chains, each pair having one "light" (about 25 kDa) and one "heavy" chain (about 50-70 kDa). The amino-terminal portion of each chain includes a variable region of about 100 to 110 or more amino acids primarily responsible for antigen recognition. The carboxy-terminal portion of each chain defines a constant region primarily responsible for effector function. In general, antibody molecules obtained from humans relate to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG1, IgG2, and others. Furthermore, in humans, the light chain may be a kappa chain or a lambda chain.

[0065] As used herein, the term "activatable antibody" refers to an antibody that also comprises a masking moiety (MM) and a cleavable moiety (CM), wherein the MM is joined to the VL of the antibody via the CM, which is cleavable by a protease. As used herein, a "prodomain" comprises the N-terminal fragment that is joined to the VL domain of the anti-human CTLA-4 activatable antibodies and, as such, comprises the MM and CM. In some embodiments, the light chain of the activatable antibody has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-VL. In some embodiments, the prodomain is joined to the VH domain of the anti-human CTLA-4 antibody. An activatable antibody is designed to be cleaved by upregulated proteolytic activity present in most if not all cancers. Such proteolytic cleavage, or activation, removes the prodomain and releases an active antibody, i.e., an activated activatable antibody. Protease activation of activatable antibodies in normal tissue is significantly reduced due to the tight control of proteolytic activity in normal tissues. As such, activatable antibodies remain largely inert in circulation and in normal tissues.

[0066] An activatable antibody, in view of its prodomain masking the antigen binding domain thereby inhibiting the ability of the antigen binding domain to bind to its target, has a lower affinity for binding to the target than does an activated activatable antibody, in which the MM has been removed by proteolytic cleavage of the CM thereby releasing an active antibody. Such released antibody exhibits higher affinity for binding to its target. In some embodiments, the MM interacts specifically with the antigen binding domain of ipilimumab to reduce the antibody's ability to bind to its target. When the MM

is removed by proteolytic cleavage of the activatable antibody, the released antibody binds to its target with an affinity similar to the parental ipilimumab.

[0067] Schematic representations of activatable antibodies of the present invention, *e.g.*, MM-CM-VL, are not intended to be exclusive. Other sequence elements, such as linkers, spacers and signal sequences, may be present before, after, or between the listed sequence elements in such schematic representations. It is also to be appreciated that a prodomain comprising a MM and a CM can be joined to a VH of an antibody instead of to a VL of an antibody such that the heavy chain has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-VH.

[0068] The term "monoclonal antibody" (mAb) or "monoclonal antibody composition", as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs contain an antigen binding site, or domain, capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it. Monoclonal antibody molecules will typically comprise two heavy chains and two light chains.

[0069] The term "antigen binding domain" refers to the part of the immunoglobulin molecule that participates in antigen binding. The antigen binding site is formed by amino acid residues of the N-terminal variable ("V") regions of the heavy ("H") and light ("L") chains. Three highly divergent stretches within the V regions of the heavy and light chains, referred to as "hypervariable regions," are interposed between more conserved flanking stretches known as "framework regions," or "FRs". Thus, the term "FR" refers to amino acid sequences which are naturally found between, and adjacent to, hypervariable regions in immunoglobulins. In an antibody molecule, the three hypervariable regions of a light chain and the three hypervariable regions of a heavy chain are disposed relative to each other in three dimensional space to form an antigen-binding surface. The antigen-binding surface is complementary to the three-dimensional surface of a bound antigen, and the three hypervariable regions of each of the heavy and light chains are referred to as "complementarity-determining regions," or "CDRs." The assignment of amino acids to each domain is in accordance with the definitions of Kabat Sequences of Proteins of

Immunological Interest (National Institutes of Health, Bethesda, Md. (1987 and 1991)), or Chothia & Lesk *J. Mol. Biol.* 196:901-917 (1987), Chothia *et al. Nature* 342:878-883 (1989).

[0070] As used herein, the term "epitope" includes any protein determinant capable of specific binding to an immunoglobulin, an scFv, or a T-cell receptor. The term "epitope" includes any protein determinant capable of specific binding to an immunoglobulin or T-cell receptor. Epitopic determinants usually consist of chemically active surface groupings of molecules such as amino acids or sugar side chains and usually have specific three dimensional structural characteristics, as well as specific charge characteristics. For example, antibodies may be raised against N-terminal or C-terminal peptides of a polypeptide. An antibody is said to specifically bind an antigen when the dissociation constant is $\leq 1 \mu\text{M}$; preferably $\leq 100 \text{ nM}$ and most preferably $\leq 10 \text{ nM}$.

[0071] As used herein, the terms "specific binding," "immunological binding," and "immunological binding properties" refer to the non-covalent interactions of the type which occur between an immunoglobulin molecule and an antigen for which the immunoglobulin is specific. The strength, or affinity of immunological binding interactions can be expressed in terms of the dissociation constant (K_d) of the interaction, wherein a smaller K_d represents a greater affinity. Immunological binding properties of selected polypeptides can be quantified using methods well known in the art. One such method entails measuring the rates of antigen-binding site/antigen complex formation and dissociation, wherein those rates depend on the concentrations of the complex partners, the affinity of the interaction, and geometric parameters that equally influence the rate in both directions. Thus, both the "on rate constant" (k_{on}) and the "off rate constant" (k_{off}) can be determined by calculation of the concentrations and the actual rates of association and dissociation. (See *Nature* 361:186-87 (1993)). The ratio of k_{off}/k_{on} enables the cancellation of all parameters not related to affinity, and is equal to the dissociation constant K_d . (See, generally, Davies *et al.* (1990) *Annual Rev Biochem* 59:439-473). An antibody of the present invention is said to specifically bind to CTLA-4, when the equilibrium binding constant (K_d) is $\leq 1 \mu\text{M}$, preferably $\leq 100 \text{ nM}$, more preferably $\leq 10 \text{ nM}$, and most preferably $\leq 100 \text{ pM}$ to about 1 pM , as measured by assays such as radioligand binding assays or similar assays known to those skilled in the art.

- [0072] The term "isolated polynucleotide" as used herein refers to a polynucleotide of genomic, cDNA, or synthetic origin or some combination thereof, which by virtue of its origin the "isolated polynucleotide" (1) is not associated with all or a portion of a polynucleotide in which the "isolated polynucleotide" is found in nature, (2) is operably linked to a polynucleotide which it is not linked to in nature, or (3) does not occur in nature as part of a larger sequence. Polynucleotides in accordance with the invention include the nucleic acid molecules encoding the heavy chain immunoglobulin molecules shown herein, and nucleic acid molecules encoding the light chain immunoglobulin molecules shown herein.
- [0073] The term "isolated protein" referred to herein means a protein of cDNA, recombinant RNA, or synthetic origin or some combination thereof, which by virtue of its origin, or source of derivation, the "isolated protein" (1) is not associated with proteins found in nature, (2) is free of other proteins from the same source, e.g., free of murine proteins, (3) is expressed by a cell from a different species, or (4) does not occur in nature.
- [0074] The term "polypeptide" is used herein as a generic term to refer to native protein, fragments, or analogs of a polypeptide sequence. Hence, native protein fragments, and analogs are species of the polypeptide genus. Polypeptides in accordance with the invention comprise the heavy chain immunoglobulin molecules shown herein, and the light chain immunoglobulin molecules shown herein, as well as antibody molecules formed by combinations comprising the heavy chain immunoglobulin molecules with light chain immunoglobulin molecules, such as kappa light chain immunoglobulin molecules, and vice versa, as well as fragments and analogs thereof.
- [0075] The term "naturally-occurring" as used herein as applied to an object refers to the fact that an object can be found in nature. For example, a polypeptide or polynucleotide sequence that is present in an organism (including viruses) that can be isolated from a source in nature and which has not been intentionally modified by man in the laboratory or otherwise is naturally-occurring.
- [0076] The term "operably linked" as used herein refers to positions of components so described are in a relationship permitting them to function in their intended manner. A control sequence "operably linked" to a coding sequence is ligated in such a way that

expression of the coding sequence is achieved under conditions compatible with the control sequences.

[0077] The term "control sequence" as used herein refers to polynucleotide sequences which are necessary to effect the expression and processing of coding sequences to which they are ligated. The nature of such control sequences differs depending upon the host organism in prokaryotes, such control sequences generally include promoter, ribosomal binding site, and transcription termination sequence in eukaryotes, generally, such control sequences include promoters and transcription termination sequence. The term "control sequences" is intended to include, at a minimum, all components whose presence is essential for expression and processing, and can also include additional components whose presence is advantageous, for example, leader sequences and fusion partner sequences. The term "polynucleotide" as referred to herein means nucleotides of at least 10 bases in length, either ribonucleotides or deoxynucleotides or a modified form of either type of nucleotide. The term includes single and double stranded forms of DNA.

[0078] The term "oligonucleotide" referred to herein includes naturally occurring, and modified nucleotides linked together by naturally occurring, and non-naturally occurring oligonucleotide linkages. Oligonucleotides are a polynucleotide subset generally comprising a length of 200 bases or fewer. Preferably oligonucleotides are 10 to 60 bases in length and most preferably 12, 13, 14, 15, 16, 17, 18, 19, or 20 to 40 bases in length. Oligonucleotides are usually single stranded, e.g., for probes, although oligonucleotides may be double stranded, e.g., for use in the construction of a gene mutant. Oligonucleotides of the invention are either sense or antisense oligonucleotides.

[0079] The term "naturally occurring nucleotides" referred to herein includes deoxyribonucleotides and ribonucleotides. The term "modified nucleotides" referred to herein includes nucleotides with modified or substituted sugar groups and the like. The term "oligonucleotide linkages" referred to herein includes oligonucleotide linkages such as phosphorothioate, phosphorodithioate, phosphoroselenoate, phosphorodiselenoate, phosphoroanilothioate, phosphoraniladate, phosphoronmidate, and the like. *See e.g., LaPlanche et al. Nucl. Acids Res.* 14:9081 (1986); *Stec et al. J. Am. Chem. Soc.* 106:6077 (1984); *Stein et al. Nucl. Acids Res.* 16:3209 (1988); *Zon et al. Anti Cancer Drug Design* 6:539 (1991); *Zon et al. Oligonucleotides and Analogues: A Practical Approach*, pp. 87-108 (F. Eckstein, Ed., Oxford University Press, Oxford England (1991)); *Stec et al. U.S.*

Pat. No. 5,151,510; Uhlmann and Peyman *Chemical Reviews* 90:543 (1990). An oligonucleotide can include a label for detection, if desired.

[0080] As used herein, the twenty conventional amino acids and their abbreviations follow conventional usage. See Immunology—A Synthesis (2nd Edition, E. S. Golub and D. R. Gren, Eds., Sinauer Associates, Sunderland Mass. (1991)). Stereoisomers (e.g., D-amino acids) of the twenty conventional amino acids, unnatural amino acids such as α -, α -disubstituted amino acids, N-alkyl amino acids, lactic acid, and other unconventional amino acids may also be suitable components for polypeptides of the present invention. Examples of unconventional amino acids include: 4 hydroxyproline, γ -carboxyglutamate, ϵ -N,N,N-trimethyllysine, ϵ -N-acetyllysine, O-phosphoserine, N-acetylserine, N-formylmethionine, 3-methylhistidine, 5-hydroxylysine, σ -N-methylarginine, and other similar amino acids and imino acids (e.g., 4-hydroxyproline). In the polypeptide notation used herein, the left-hand direction is the amino terminal direction and the right-hand direction is the carboxy-terminal direction, in accordance with standard usage and convention.

[0081] As applied to polypeptides, the term "substantial identity" means that two peptide sequences, when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights, share at least 80 percent sequence identity, preferably at least 90 percent sequence identity, more preferably at least 95 percent sequence identity, and most preferably at least 99 percent sequence identity.

[0082] As discussed herein, minor variations in the amino acid sequences of antibodies or immunoglobulin molecules are contemplated as being encompassed by the present invention, providing that the variations in the amino acid sequence maintain at least 75%, more preferably at least 80%, 90%, 95%, and most preferably 99% sequence identity. In particular, conservative amino acid replacements are contemplated. Conservative replacements are those that take place within a family of amino acids that are related in their side chains. Genetically encoded amino acids are generally divided into families: (1) acidic amino acids are aspartate, glutamate; (2) basic amino acids are lysine, arginine, histidine; (3) non-polar amino acids are alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan, and (4) uncharged polar amino acids are glycine, asparagine, glutamine, cysteine, serine, threonine, tyrosine. The hydrophilic amino acids include arginine, asparagine, aspartate, glutamine, glutamate, histidine, lysine, serine, and

threonine. The hydrophobic amino acids include alanine, cysteine, isoleucine, leucine, methionine, phenylalanine, proline, tryptophan, tyrosine and valine. Other families of amino acids include (i) serine and threonine, which are the aliphatic-hydroxy family; (ii) asparagine and glutamine, which are the amide containing family; (iii) alanine, valine, leucine and isoleucine, which are the aliphatic family; and (iv) phenylalanine, tryptophan, and tyrosine, which are the aromatic family. In the case of an antibody, it is reasonable to expect that an isolated replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a similar replacement of an amino acid with a structurally related amino acid will not have a major effect on the binding or properties of the resulting molecule, especially if the replacement does not involve an amino acid within a CDR or framework region. Whether an amino acid change results in a functional peptide can readily be determined by assaying the specific activity of the polypeptide derivative. Assays are described in detail herein. Fragments or analogs of antibodies or immunoglobulin molecules can be readily prepared by those of ordinary skill in the art. Preferred amino- and carboxy-termini of fragments or analogs occur near boundaries of functional domains. Structural and functional domains can be identified by comparison of the nucleotide and/or amino acid sequence data to public or proprietary sequence databases. Preferably, computerized comparison methods are used to identify sequence motifs or predicted protein conformation domains that occur in other proteins of known structure and/or function. Methods to identify protein sequences that fold into a known three-dimensional structure are known. Bowie *et al. Science* 253:164 (1991). Thus, the foregoing examples demonstrate that those of skill in the art can recognize sequence motifs and structural conformations that may be used to define structural and functional domains in accordance with the invention.

[0083] Preferred amino acid substitutions are those which: (1) reduce susceptibility to proteolysis in regions of the activatable antibody other than in the cleavable linker comprising the CM, (2) reduce susceptibility to oxidation, (3) alter binding affinity for forming protein complexes, (4) alter binding affinities, and (4) confer or modify other physicochemical or functional properties of such analogs. Analogs can include various muteins of a sequence other than the naturally-occurring peptide sequence. For example, single or multiple amino acid substitutions (preferably conservative amino acid substitutions) may be made in the naturally-occurring sequence (preferably in the portion

of the polypeptide outside the domain(s) forming intermolecular contacts). A conservative amino acid substitution should not substantially change the structural characteristics of the parent sequence (e.g., a replacement amino acid should not tend to break a helix that occurs in the parent sequence, or disrupt other types of secondary structure that characterizes the parent sequence). Examples of art-recognized polypeptide secondary and tertiary structures are described in *Proteins, Structures and Molecular Principles* (Creighton, Ed., W. H. Freeman and Company, New York (1984)); *Introduction to Protein Structure* (C. Branden and J. Tooze, eds., Garland Publishing, New York, N.Y. (1991)); and Thornton *et al. Nature* 354:105 (1991).

[0084] The term "polypeptide fragment" as used herein refers to a polypeptide that has an amino terminal and/or carboxy-terminal deletion and/or one or more internal deletion(s), but where the remaining amino acid sequence is identical to the corresponding positions in the naturally-occurring sequence deduced, for example, from a full length cDNA sequence. Fragments typically are at least 5, 6, 8 or 10 amino acids long, preferably at least 14 amino acids long, more preferably at least 20 amino acids long, usually at least 50 amino acids long, and even more preferably at least 70 amino acids long. The term "analog" as used herein refers to polypeptides which comprise a segment of at least 25 amino acids that has substantial identity to a portion of a deduced amino acid sequence and which has specific binding to CTLA-4, under suitable binding conditions. Typically, polypeptide analogs comprise a conservative amino acid substitution (or addition or deletion) with respect to the naturally-occurring sequence. Analogs typically are at least 20 amino acids long, preferably at least 50 amino acids long or longer, and can often be as long as a full-length naturally-occurring polypeptide.

[0085] The term "agent" is used herein to denote a chemical compound, a mixture of chemical compounds, a biological macromolecule, or an extract made from biological materials.

[0086] As used herein, the terms "label" or "labeled" refers to incorporation of a detectable marker, e.g., by incorporation of a radiolabeled amino acid or attachment to a polypeptide of biotinyl moieties that can be detected by marked avidin (e.g., streptavidin containing a fluorescent marker or enzymatic activity that can be detected by optical or calorimetric methods). In certain situations, the label or marker can also be therapeutic. Various methods of labeling polypeptides and glycoproteins are known in the art and may

be used. Examples of labels for polypeptides include, but are not limited to, the following: radioisotopes or radionuclides (e.g., ^3H , ^{14}C , ^{15}N , ^{35}S , ^{90}Y , ^{99}Tc , ^{111}In , ^{125}I , ^{131}I) fluorescent labels (e.g., FITC, rhodamine, lanthanide phosphors), enzymatic labels (e.g., horseradish peroxidase, p-galactosidase, luciferase, alkaline phosphatase), chemiluminescent, biotinyl groups, predetermined polypeptide epitopes recognized by a secondary reporter (e.g., leucine zipper pair sequences, binding sites for secondary antibodies, metal binding domains, epitope tags). In some embodiments, labels are attached by spacer arms of various lengths to reduce potential steric hindrance.

[0087] Other chemistry terms herein are used according to conventional usage in the art, as exemplified by The McGraw-Hill Dictionary of Chemical Terms (Parker, S., Ed., McGraw-Hill, San Francisco (1985)).

[0088] As used herein, "substantially pure" means an object species is the predominant species present (i.e., on a molar basis it is more abundant than any other individual species in the composition), and preferably a substantially purified fraction is a composition wherein the object species comprises at least about 50 percent (on a molar basis) of all macromolecular species present. Generally, a substantially pure composition will comprise more than about 80 percent of all macromolecular species present in the composition, more preferably more than about 85%, 90%, 95%, and 99%. Most preferably, the object species is purified to essential homogeneity (contaminant species cannot be detected in the composition by conventional detection methods) wherein the composition consists essentially of a single macromolecular species.

[0089] As used herein, "treatment" is an approach for obtaining beneficial or desired clinical results. Beneficial or desired clinical results may include, but are not limited to, any one or more of: alleviation of one or more symptoms, diminishment of extent of disease, stabilized (i.e., not worsening) state of disease, preventing or delaying spread (e.g., metastasis) of disease, preventing or delaying occurrence or recurrence of disease, delay or slowing of disease progression, amelioration of the disease state, and remission (whether partial or total). Also encompassed by "treatment" is a reduction of pathological consequence of a proliferative disease such as cancer. The methods provided herein contemplate any one or more of these aspects of treatment.

[0090] The term "effective amount" used herein refers to an amount of a compound or composition, when used alone or in combination with a second therapy, is sufficient to

treat a specified disorder, condition or disease such as ameliorate, palliate, lessen, and/or delay one or more of its symptoms. In reference to cancers or other unwanted cell proliferation, an effective amount comprises an amount sufficient to cause a tumor to shrink and/or to decrease the growth rate of the tumor (such as to suppress tumor growth) or to prevent or delay other unwanted cell proliferation. An effective amount can be administered in one or more administrations.

[0091] As used herein, by "combination therapy" is meant that a first agent be administered in conjunction with another agent. "In conjunction with" refers to administration of one treatment modality in addition to another treatment modality. As such, "in conjunction with" refers to administration of one treatment modality before, during, or after delivery of the other treatment modality to the individual.

[0092] The term "pharmaceutical agent or drug" as used herein refers to a chemical compound or composition capable of inducing a desired therapeutic effect when properly administered to a subject.

[0093] As used herein, by "pharmaceutically acceptable" or "pharmacologically compatible" is meant a material that is not biologically or otherwise undesirable, e.g., the material may be incorporated into a pharmaceutical composition administered to an individual or subject without causing any significant undesirable biological effects or interacting in a deleterious manner with any of the other components of the composition in which it is contained. Pharmaceutically acceptable carriers or excipients have for example met the required standards of toxicological and manufacturing testing and/or are included on the Inactive Ingredient Guide prepared by the U.S. Food and Drug administration.

[0094] The terms "cancer", "cancerous", or "malignant" refer to or describe the physiological condition in mammals that is typically characterized by unregulated cell growth. Examples of cancer include, for example, melanoma, such as unresectable or metastatic melanoma, leukemia, lymphoma, blastoma, carcinoma and sarcoma. More particular examples of such cancers include chronic myeloid leukemia, acute lymphoblastic leukemia, Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL), squamous cell carcinoma, small-cell lung cancer, non-small cell lung cancer, glioma, gastrointestinal cancer, renal cancer, ovarian cancer, liver cancer, colorectal cancer, endometrial cancer, kidney cancer, prostate cancer, thyroid cancer,

neuroblastoma, pancreatic cancer, glioblastoma multiforme, cervical cancer, stomach cancer, bladder cancer, hepatoma, breast cancer, colon carcinoma, and head and neck cancer, gastric cancer, germ cell tumor, pediatric sarcoma, sinonasal natural killer, multiple myeloma, acute myelogenous leukemia (AML), and chronic lymphocytic leukemia (CML).

[0095] "Leukemia" refers to progressive, malignant diseases of the blood-forming organs and is generally characterized by a distorted proliferation and development of leukocytes and their precursors in the blood and bone marrow. Leukemia is generally clinically classified on the basis of (1) the duration and character of the disease--acute or chronic; (2) the type of cell involved; myeloid (myelogenous), lymphoid (lymphogenous), or monocytic; and (3) the increase or non-increase in the number of abnormal cells in the blood--leukemic or aleukemic (subleukemic). Leukemia includes, for example, acute nonlymphocytic leukemia, chronic lymphocytic leukemia, acute granulocytic leukemia, chronic granulocytic leukemia, acute promyelocytic leukemia, adult T-cell leukemia, aleukemic leukemia, a leukocythemiac leukemia, basophilic leukemia, blast cell leukemia, bovine leukemia, chronic myelocytic leukemia, leukemia cutis, embryonal leukemia, eosinophilic leukemia, Gross' leukemia, hairy-cell leukemia, hemoblastic leukemia, hemocytoblastic leukemia, histiocytic leukemia, stem cell leukemia, acute monocytic leukemia, leukopenic leukemia, lymphatic leukemia, lymphoblastic leukemia, lymphocytic leukemia, lymphogenous leukemia, lymphoid leukemia, lymphosarcoma cell leukemia, mast cell leukemia, megakaryocytic leukemia, micromyeloblastic leukemia, monocytic leukemia, myeloblastic leukemia, myelocytic leukemia, myeloid granulocytic leukemia, myelomonocytic leukemia, Naegeli leukemia, plasma cell leukemia, plasmacytic leukemia, promyelocytic leukemia, Rieder cell leukemia, Schilling's leukemia, stem cell leukemia, subleukemic leukemia, and undifferentiated cell leukemia. In certain aspects, the present invention provides treatment for chronic myeloid leukemia, acute lymphoblastic leukemia, and/or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL).

I. Anti-CTLA-4 Activatable Antibodies

[0096] The present invention provides improved anti-CTLA-4 antibodies that are as efficacious as the traditional anti-CTLA-4 antibodies (e.g., ipilimumab) but with a

greater, i.e., improved, safety profile. Specifically, the improved anti-CTLA-4 antibodies are activatable monoclonal antibodies (mAbs) that specifically bind human CTLA-4 when activated. These improved anti-CTLA-4 antibodies, also referred to herein as activatable anti-CTLA-4 antibodies or CTLA-4 activatable antibodies, are used in methods of treating, preventing, delaying the progression of, ameliorating and/or alleviating a symptom of a disease or disorder, including but not limited to, a disease or disorder associated with aberrant CTLA-4 expression and/or activity. For example, the activatable anti-CTLA-4 antibodies are used in methods of treating, preventing, delaying the progression of, ameliorating and/or alleviating a symptom of a cancer or other neoplastic condition. Activatable antibodies are described in, for example, US Pat. Nos. 8,513,390, 8,518,404; 9,120,853; 9,127,053 and International Publ. No. WO 2016/149201.

[0097] In some embodiments, the activatable anti-CTLA-4 antibodies provided herein comprise (i) ipilimumab or antigen binding domain thereof (AB), such as an ipilimumab variable light chain (VL), (ii) a cleavable moiety (CM), and (iii) a masking moiety (MM). In some embodiments, the VL is coupled to the MM, such that coupling of the MM reduces the ability of the ipilimumab to bind to CTLA-4. In some embodiments, the MM is coupled to the VL via a cleavable moiety (CM) (also known as a substrate linker) that includes a substrate for a protease, for example, a protease that is over-expressed in the tumor microenvironment.

Antibody or Antigen Binding Fragment Thereof

[0098] In some embodiments, the antibody or antigen binding domain thereof (AB) comprises the complementarity determining regions (CDRs) of the anti-CTLA-4 antibody ipilimumab, identified as 10D1 in U.S. Patent Nos. 6,984,720 and 7,605,238, which are hereby incorporated by reference in their entireties. Ipilimumab (also formerly known as MDX-010 and BMS-734016) is marketed as YERVOY® and has been approved for the treatment of metastatic melanoma and is in clinical testing in other cancers. *See* Hoos *et al.* (2010) *Semin. Oncol.* 37:533; Hodi *et al.* (2010) *N. Engl. J. Med.* 363:711; Pardoll (2012) *Nat. Immunol.* 13(12): 1129.

[0099] Ipilimumab has a human IgG1 isotype, which binds best to most human Fc receptors (Bruhns *et al.* (2009) *Blood* 113: 3716) and is considered equivalent to murine

IgG2a with respect to the types of activating Fc receptors that it binds. Since IgG1 binds to the activating receptor CD16 (FcγRIIIa) expressed by human NK cells and monocytes, ipilimumab can mediate ADCC. The IgG1-isotype ipilimumab was originally isolated directly from a hybridoma but was subsequently cloned and expressed in Chinese hamster ovary (CHO) cells. Notwithstanding the consideration that an isotype that mediates ADCC and/or CDC might be undesirable in an antibody targeting a receptor on T cells that seeks to upregulate an immune response, the IgG1 isotype of the antibody was retained, in part, because it enhanced vaccine response in cynomolgus monkey and was considered functional. Ipilimumab has been shown to increase the numbers of activated T cells in the blood, as evidenced, for example, by a significant increase in the expression of HLA-DR on the surface of post-treatment CD4⁺ and CD8⁺ cells as well as increases in absolute lymphocyte count (Ku *et al.* (2010) *Cancer* 116:1767; Attia *et al.* (2005) *J. Clin. Oncol.* 23:6043; Maker *et al.* (2005) *J. Immunol.* 175:7746; Berman *et al.* (2009) *J. Clin. Oncol.* 27(suppl):15s.3020; Hamid *et al.* (2009) *J. Clin. Oncol.* 27(suppl): 15s.9008), indicating that depletion of T cells does not occur in the periphery in man. Ipilimumab demonstrated only modest levels of ADCC of activated T cells using IL-2-activated PBMCs as effector cells; however, use of T_{regs} as targets was not tested. Minor changes in peripheral T_{reg} frequency in the blood of patients treated with ipilimumab have been observed (Maker *et al.* (2005) *J. Immunol.* 175:7746), but little information of the effect of ipilimumab on intratumoral T_{regs} is available. However, a positive correlation between a high CD8⁺ to T_{reg} ratio and tumor necrosis in biopsies from metastatic melanoma lesions from patients treated with ipilimumab have been described. Hodi *et al.* (2008) *Proc. Nat'l Acad. Sci. (USA)* 105:3005. In addition, tumor tissue from ipilimumab-treated bladder cancer patients had lower percentages of CD4⁺ Foxp3⁺ T cells than tumors from untreated bladder cancer patients. Liakou *et al.* (2008) *Proc. Nat'l Acad. Sci. (USA)* 105:14987.

[0100] In some embodiments, the activatable anti-CTLA-4 antibody comprises a combination of a variable heavy chain CDR1 (VH CDR1, also referred to herein as CDRH1), CDR2 (VH CDR2, also referred to herein as CDRH2), and CDR3 (VH CDR3, also referred to herein as CDRH3), and a variable light chain CDR1 (VL CDR1, also referred to herein as CDRL1), CDR2 (VL CDR2, also referred to herein as CDRL2), and

CDR3 (VL CDR3, also referred to herein as CDRL3). These CDR sequences are provided at Table 2.

Table 2: CDR Sequences of heavy and light chains for Ipilimumab

CHAIN	CDR1	CDR2	CDR3
LIGHT	RASQSVGSSYLA (SEQ ID NO: 560)	GAFSRAT (SEQ ID NO: 561)	QQYGSSPWT (SEQ ID NO: 562)
HEAVY	SYTMH (SEQ ID NO: 557)	FISYDGNNKYYADSVKG (SEQ ID NO: 558)	TGWLGPFDY (SEQ ID NO: 559)

[0101] Ipilimumab-VL chain

EIVLTQSPGTLSPGERATLSCRASQSVGSSYLA WYQQKPGQAPRLLIYGAFSRA
TGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKVEIK (SEQ
ID NO: 344)

[0102] Ipilimumab-VH chain

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYTMHWVRQAPGKGLEWVTFISYD
GNNKYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAIYYCARTGWLGPFDY
WGQGTLLTVSS (SEQ ID NO: 345)

[0103] Various other sequences, as indicated, are provided below.

[0104] Human Kappa constant LC

RTVAAPSVFIFPPSDEQLKSGTASVVCLLNFPYPREAKVQWKVDNALQSGNSQES
VTEQDSKDSSTLSSTLTLSKADYEEKHKVYACEVTHQGLSSPVTKSFNRGEC
(SEQ ID NO: 346)

[0105] Mouse Kappa constant light chain

RADAAPT VSI FPPSSEQLTSGGASVVCFLNFPY PKDINVKWKIDG SERQNGVLNS
WTDQDSKDSSTYSMSSTLT LTKDEYERHNSYTCEATHKTSTSPIVKSFNRNEC
(SEQ ID NO: 347)

[0106] Ipilimumab—Human Kappa LC

EIVLTQSPGTLSPGERATLSCRASQSVGSSYLA WYQQKPGQAPRLLIYGAFSRA
TGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKVEIKRTVA

APSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQ
DSKDSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID
NO: 348)

[0107] Ipilimumab—Mouse Kappa LC

EIVLTQSPGTLSPGERATLSCRASQSVGSSYLAWYQQKPGQAPRLLIYGAFSRA
TGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKVEIKRADA
APTVSIFPPSSEQLTSGGASVVCFLNNFYPKDINVKWKIDGSRQNGVLNSWTDQ
DSKDSTYSMSSTLTTLTKDEYERHNSYTCEATHKTSTSPIVKSFNRNEC (SEQ ID
NO: 349)

[0108] Human IgG1 constant HC

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPA
VLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEPKSCDKTHTCPP
CPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVE
VHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIS
KAKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNY
KTTTPVLDSGDSFFLYSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSP
G (SEQ ID NO: 350)

[0109] Mouse IgG1 constant HC

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPA
VLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCP
PCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGV
EVHNAKTKPREEQYASTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI
SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENN
YKTTTPVLDSGDSFFLYSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLS
PG (SEQ ID NO: 351)

[0110] Mouse IgG2a constant HC

AKTTAPSVYPLAPVCGDTTGSSVTLGCLVKGYFPEPVTLTWNSGSLSSGVHTFPA
VLQSDLYTLSSSVTVTSSTWPSQSITCNVAHPASSTKVDKKIEPRGPTIKPCPPCKC
PAPNLLGGPSVFIFPPKIKDVLMISSLPIVTCVVDVSEDDPDVQISWVFNNEVH
TAQTQTHREDYNSTLRVVSALPIQHQQDWMSGKEFKCKVNNKDLPAPIERTISKPK

GSVRAPQVYVLPPPEEEMTKKQVTLTCTMVTDFMPEDIYVEWTNNGKTELNYKN
TEPVLDSGDSYFMYSKLRVEKKNWVERNSYSCSVVHEGLHNHHTTKSFSRTPGK
(SEQ ID NO: 352)

[0111] Ipilimumab-VH—Human IgG1 constant HC

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYTMHWVRQAPGKGLEWVTFISYD
GNNKYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAIYYCARTGWLGPFDY
WGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSG
ALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKRVEP
KSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEV
KFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSN
KALPAPIEKTISKAKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEW
ESNGQPENNYKTTTPVLDSGDSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNH
YTQKSLSLSPG (SEQ ID NO: 353)

[0112] Ipilimumab-VH—Mouse IgG1 constant HC

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYTMHWVRQAPGKGLEWVTFISYD
GNNKYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAIYYCARTGWLGPFDY
WGQGTLVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSG
ALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVE
PKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPE
VKFNWYVDGVEVHNAKTKPREEQYASTYRVVSVLTVLHQDWLNGKEYKCKVSN
KALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVE
WESNGQPENNYKTTTPVLDSGDSFFLYSKLTVDKSRWQQGNVFSCSVMHEALH
NHYTQKSLSLSPG (SEQ ID NO: 354)

[0113] Ipilimumab-VH—Mouse IgG2a constant HC

QVQLVESGGGVVQPGRSLRLSCAASGFTFSSYTMHWVRQAPGKGLEWVTFISYD
GNNKYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAIYYCARTGWLGPFDY
WGQGTLVTVSSAKTTAPSVYPLAPVCGDTTGSSVTLGCLVKGYFPEPVTLTWNS
GSLSSGVHTFPAVLQSDLYTLSSSVTVTSSTWPSQSITCNVAHPASSTKVDKKIEP
RGPTIKPCPPCKCPAPNLLGGPSVFIFPPKIKDVLMI SLSPIVTCVVDVSEDDPDV
QISW FVNNVEVHTAQTQTHREDYNSTLRVVSALPIQH QDWMSGKEFKCKVNNK

DLPAPIERTISKPKGSVRAPQVYVLPPEEEMTKKQVTLTCMVTDMPEDIYVEV
TNNGKTELNYKNTEPVLDSDGSYFMYSKLRVEKKNWVERNSYSCSVVHEGLHN
HHTTKSFSRTPGK (SEQ ID NO: 355)

- [0114] In some embodiments, the antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence comprises 1, 2, 3, 4 or more amino acid sequence differences compared with the CDR sequences shown in Table 2, including conservative amino acid differences.
- [0115] In some embodiments, the activatable anti-CTLA-4 antibody comprises a heavy chain variable domain that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the group consisting of SEQ ID NO: 345. In some embodiments, the activatable anti-CTLA-4 antibody comprises a light chain variable domain, not including any MM, CM, linker, spacer or other sequence added in creation of the activatable form of the antibody, that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the group consisting of SEQ ID NOs: 563 to 565.
- [0116] In some embodiments, the antibody or antigen-binding fragment thereof that binds CTLA-4 in the activatable antibodies can include modifications, particularly in the Fc region of the antibody or antigen-binding fragment thereof. For example, the interaction of antibodies with FcγRs can be enhanced by modifying the glycan moiety attached to each Fc fragment at the N297 residue. In particular, the absence of core fucose residues strongly enhances ADCC via improved binding of IgG to activating FcγRIIIA without altering antigen binding or CDC. Natsume *et al.* (2009) *Drug Des. Devel. Ther.* 3:7. There is convincing evidence that afucosylated tumor-specific antibodies translate into enhanced therapeutic activity in mouse models *in vivo*. Nimmerjahn & Ravetch (2005) *Science* 310:1510; Mossner *et al.* (2010) *Blood* 115:4393.
- [0117] Modification of antibody glycosylation can be accomplished by, for example, expressing the antibody in a host cell with altered glycosylation machinery. Cells with altered glycosylation machinery have been described in the art and can be used as host cells in which to express recombinant antibodies of this disclosure to thereby produce an antibody with altered glycosylation. For example, the cell lines Ms704, Ms705, and Ms709 lack the fucosyltransferase gene, FUT8 (α -(1,6) fucosyltransferase) (see U.S. Pat.

App. Publication No. 20040110704; Yamane-Ohnuki *et al.* (2004) *Biotechnol. Bioeng.* 87: 614), such that antibodies expressed in these cell lines lack fucose on their carbohydrates. As another example, EP 1176195 also describes a cell line with a functionally disrupted FUT8 gene as well as cell lines that have little or no activity for adding fucose to the N-acetylglucosamine that binds to the Fc region of the antibody, for example, the rat myeloma cell line YB2/0 (ATCC CRL 1662). PCT Publication WO 03/035835 describes a variant CHO cell line, Lec13, with reduced ability to attach fucose to Asn (297)-linked carbohydrates, also resulting in hypofucosylation of antibodies expressed in that host cell. See also Shields *et al.* (2002) *J. Biol. Chem.* 277:26733. Antibodies with a modified glycosylation profile can also be produced in chicken eggs, as described in PCT Publication No. WO 2006/089231. Alternatively, antibodies with a modified glycosylation profile can be produced in plant cells, such as Lemna. See e.g. U.S. Publication No. 2012/0276086. PCT Publication No. WO 99/54342 describes cell lines engineered to express glycoprotein-modifying glycosyl transferases (e.g., beta(1,4)-N-acetylglucosaminyltransferase III (GnTIII)) such that antibodies expressed in the engineered cell lines exhibit increased bisecting GlcNac structures which results in increased ADCC activity of the antibodies. See also Umaña *et al.* (1999) *Nat. Biotech.* 17:176. Alternatively, the fucose residues of the antibody may be cleaved off using a fucosidase enzyme. For example, the enzyme alpha-L-fucosidase removes fucosyl residues from antibodies. Tarentino *et al.* (1975) *Biochem.* 14:5516. Core fucosylation may also be reduced by culturing antibody-producing cells in the presence of small molecule fucose analogs, such as those described at EP2282773B1, or in the presence of castanospermine, as described at WO 08/052030.

Cleavable Moiety

[0118] In some embodiments, the CM is specific for a protease, which is useful in leveraging the dysregulated protease activity in tumor cells for targeted activatable antibody activation at the site of treatment and/or diagnosis. Numerous studies have demonstrated the correlation of aberrant protease levels, e.g., uPA, legumain, MT-SP1, matrix metalloproteases (MMPs), in solid tumors. (See e.g., Murthy R V, *et al.* "Legumain expression in relation to clinicopathologic and biological variables in colorectal cancer." *Clin Cancer Res.* 11 (2005): 2293-2299; Nielsen B S, *et al.*

"Urokinase plasminogen activator is localized in stromal cells in ductal breast cancer." *Lab Invest* 81 (2001): 1485-1501; Look O R, *et al.* "In situ localization of gelatinolytic activity in the extracellular matrix of metastases of colon cancer in rat liver using quenched fluorogenic DQ-gelatin." *J Histochem Cytochem.* 51 (2003): 821-829).

[0119] A general overview of this process is discussed in US Pat. Nos. 7,666,817, 8,513,390, and 9,120,853 and International Publication Nos. WO 2016/118629 and WO 2016/149201, which are hereby incorporated by reference in their entireties. The cleavable moiety selection process is used to identify cleavable moieties that have a number of desirable characteristics. For example, the selected cleavable moieties are systemically stable (i.e., stable in the systemic circulation of a subject), are generally not susceptible to cleavage by circulating proteases such as plasmin, thrombin, tissue plasminogen activator (tPA) or a kallikrein (KLK) such as KLK-5 and/or KLK-7, are non-toxic, are generally not susceptible to cleavage at potential sites of toxicity such as the skin by proteases such as ADAM 9, ADAM 10, ADAM 17 and/or kallikreins, such as KLK-5 and KLK-7, and are active at an intended site of treatment and/or diagnosis. In some embodiments, the identified cleavable moieties are selected for proteases that are overexpressed at an intended site of therapy and/or diagnosis but are not typically expressed at or in normal, healthy or otherwise non-diseased or non-damaged tissue, and then the selected substrates are subsequently counter-screened against proteases expressed in normal, e.g., non-diseased, tissue. Exemplary proteases and/or enzymes are provided in Table 1 as indicated earlier.

[0120] In some embodiments, the cleavable moiety is selected from the group consisting of 2001 and 3001, and derivatives thereof. In some embodiments, the cleavable moiety is selected from the group consisting of 2001 (SEQ ID NO: 297), 2006 (SEQ ID NO: 300), 2007 (SEQ ID NO: 301), 2008 (SEQ ID NO: 302), 2009 (SEQ ID NO: 303), 2012 (SEQ ID NO: 305), 2011 (SEQ ID NO: 304), 2003 (SEQ ID NO: 298), 3001 (SEQ ID NO: 306), 3006 (SEQ ID NO: 313), 3007 (SEQ ID NO: 308), 3008 (SEQ ID NO: 309), 3009 (SEQ ID NO: 310), 3012 (SEQ ID NO: 312), 3011 (SEQ ID NO: 311), and 2005 (SEQ ID NO: 299). Table 3 provides additional cleavable moieties that may be used with the activatable anti-CTLA-4 antibodies disclosed herein.

Table 3. Anti-CTLA-4 Activatable Cleavable Moieties

SEQUENCE IDENTIFIER	CM Sequence
313	LSGRSDNH
314	LSGRSANPRG
315	TGRGPSWV
316	PLTGRSGG
317	TARGPSFK
318	NTLSGRSENHSG
319	NTLSGRSGNHGS
320	TSTSGRSANPRG
321	TSGRSANP
322	VHMPLGFLGP
306	AVGLLAPPGGLSGRSDNH
307	AVGLLAPPGGLSGRSDDH
308	AVGLLAPPGGLSGRSDIH
309	AVGLLAPPGGLSGRSDQH
310	AVGLLAPPGGLSGRSDTH
338	AVGLLAPPGGLSGRSDYH
339	AVGLLAPPGGLSGRSANI
340	AVGLLAPPGGLSGRSDNI
312	AVGLLAPPGGLSGRSANP
311	AVGLLAPPGGLSGRSDNP
299	AVGLLAPPSGRSANPRG
323	AVGLLAPP
324	AQNLLGMV
325	QNQALRMA
326	LAAPLGLL
327	STFPFGMF
328	ISSGLLSS
329	PAGLWLDP
330	VAGRSMRP
331	VVPEGRRS
332	ILPRSPAF
333	MVLGRSLL
334	VAGRSMRP
335	QGRAITFI
336	SPRSIMLA
337	SMLRSMPL
297	ISSGLLSGRSDNH
300	ISSGLLSGRSDDH
301	ISSGLLSGRSDIH
302	ISSGLLSGRSDQH
303	ISSGLLSGRSDTH
341	ISSGLLSGRSDYH

342	ISSGLLSGRSANI
343	ISSGLLSGRSDNI
305	ISSGLLSGRSANP
304	ISSGLLSGRSDNP
298	ISSGLLSGRSANPRG

Masking Moiety

[0121] The activatable anti-CTLA-4 antibodies provided herein comprise a masking moiety (MM). In some embodiments, the MM is an amino acid sequence that is coupled, or otherwise attached, to the anti-CTLA-4 antibody and is positioned within the activatable anti-CTLA-4 antibody construct such that the MM reduces the ability of the anti-CTLA-4 antibody to specifically bind CTLA-4. In some embodiments, the MM binds specifically to the antigen binding domain. Suitable MMs are identified using any of a variety of known techniques. For example, peptide MMs are identified using the methods described in U.S. Patent Application Publication Nos. 2009/0062142 by Daugherty *et al.* and 2012/0244154 by Daugherty *et al.*, the contents of which are hereby incorporated by reference in their entirety.

[0122] In some embodiments, the MM is selected from the group consisting of YV01 to YV66 and comprises an amino acid sequence selected from Table 4 below.

Table 4: Anti-CTLA4 Masking Moieties (MM)

SEQUENCE IDENTIFIER	MM SEQUENCE	SEQUENCE IDENTIFIER	MM SEQUENCE
1	DFSC LHSMY NVCLDP	147	EHCDVWMFGFNLC PY
2	QPCAQMYGYSMCPHT	148	EPCDYWMFGVNLCPY
3	LHCRTQMYGYNLCPY	149	EQCTMWMYGFNLCPY
4	LHCRTQLYGYNLCPY	150	ESACSLRMYEVCLQP
5	CTYSFFNVC	151	ESCASMYGYSMCPRT
6	CAQMYGYSMC	152	ESCSYWMFGYNLC PY
7	CPNHPMC	153	FSNTCPHHPMCYDYR
8	GTACTYSFFNVCLDP	154	FWNTCPHHPMCHDYK
9	FGTACPNHPMCHDWQ	155	FYQNCYPPTWCSMFS
10	SACAYWMFGVNLCPY	156	GECSYWMFGYNLC PY
11	CRTQLYGYNLC	157	GGSCMY SFFNICLDP
12	CRTQIYGYNLC	158	GGSCVYVMY NVCLDP
13	LHCRTQIYGYNLCPY	159	GHCLMHMYGYNLCPK
14	CPNHPMCHDWQ	160	GHC RMSMYEMTLCPR
15	GTACPNHPMCHDWQ	161	GISC VHIMFNFCLDP
16	CAYWMFGVNLCPY	162	GLCVMYMFGVNLCPY
17	QECHLYMYGVNLCPY	163	GSCDYWMFGYNLC PY

18	CHLYMYGVNLCPY	164	GSYCMYVMYNNVCLDP
19	GQCQFYMFYGNLCPY	165	GTKCIYSFYNNVCLDP
20	LSTCMYSFFNVCLDP	166	GTSTCPYHPMCHDYR
21	CLHSMYNNVCLDP	167	GTTCTYSFFNVCLDP
22	CLHSMYNNVC	168	GVCHFFMYGVSMCPA
23	CLHSLYNNVCLDP	169	GVPCWYSMYNNVCLDP
24	CLHSAYNNVCLDP	170	GVSCMYSMFNICLDP
25	CMYSFFNVCLDP	171	HAKCVYSFFNVCLDP
26	CMYSFFNNVC	172	HDSCMYSMYNNFCLDP
27	QPCAQMYGYSMC	173	HGNTCPNHPMCHDYQ
28	CAQLYGYSMCPHT	174	HKGCLYSFYNNICLDP
29	CAQMYGYSMCAHT	175	HKGCLYSFYNNVCLDP
30	CAQMYGYSMCPAT	176	HLSCMYIMYNNVCLDP
31	CAQMYGYSMCPHT	177	HSSCIYSMFNVCLDP
32	CPNHPLCHDWQ	178	HTNMCPYHPMCHDYK
33	CPNHPMCADWQ	179	HTPCTYSFFNVCLDP
34	CPNHPMCHAWQ	180	IMNTCPYHPMCHDYQ
35	CPNHPMCHDAQ	181	IVPCTYMMFGVCLQP
36	CPNHPMCHDWA	182	KKCDYWFYGVNLCPY
37	GTACPNHPMC	183	KNTCVYSFFNVCLDP
38	LHCRTQLYGYNLC	184	KPCAQMYGYSMCPHP
39	CRTQLYGYNLCPY	185	KPSCMYSSFFNVCLDP
40	CRTQLYGYNLCAY	186	KRPCMYSFYNNVCLDP
41	CRTQLYGYNLCPA	187	KTSCMYSFYNNICLDP
42	FGTACPNHPLCHDWQ	188	KTTCTYSFFNVCLDP
43	CPNHPLCHDFQ	189	LDCQMYWWFGACGDM
44	CPNHPLCHDYQ	190	LHCAIYMYGYNLCPF
45	CPNHPLCPY	191	LHCPFQMYGYNLCPH
46	CPNHPLCPA	192	LHCSMYMYGFNLCPN
47	CMYSFFNVCYP	193	RECMAYMYGYNLCPY
48	CMYSFFNVCYA	194	RHCQMFMFGYDLCPY
49	CLYSFFNVCYP	195	LIHCRYVMYGMCLP
50	CLYSFFNVCYA	196	LLPCEVMGSPRCKHD
51	FGAACPNHPICHDWQ	197	LPCHAYMYGYSLCPY
52	FGAACPNHPLCHDWQ	198	LPCLAYMYGVNLCPN
53	FGAACPNHPMCHDAQ	199	LPCMAYMFGFNLCPH
54	CLHSAYNACLDP	200	LPCNFHMFGFNLCPY
55	CAHSAYNNVCLDP	201	LQCAMYMYGYNLCPY
56	CLHSAYNNVCADP	202	LSSCTYSFFNVCLDP
57	CLHSAYNNVCLAP	203	LTCPFQMYGYNLCPY
58	CLHSAYNNVCLDA	204	LTSQCSPWYWCQIYD
59	KNTCTYVMYNNVCLDP	205	LYCPYMMYGYNLCPY
60	YISDCPYHPMCHDYQ	206	LYHCTYSFYNNVCLDP
61	FRNTCPYHPMCHDYR	207	LYRCIYSFYNNVCLDP
62	RECHMWMFGVNLCPY	208	MGCSMRMWGMELCPE
63	AVCHMYMYGYNLCPF	209	MKCDYWLYGYNLCPY
64	RSCPQMYGYSMCPHT	210	MNHCTLHMYNICMDP
65	QPCAQMFGYSMCPHT	211	MNPECPPHMPCHNSN
66	TAKCTYSFFNVCLDP	212	MPACTYSFFNICLDP
67	DFSCLYSMYNNVCLDP	213	MPQCHVIMYNNICLDP
68	DVSCMYMMYNNFCLDP	214	MSTCTYSFFNVCLDP

69	CPNHPMC	215	MTCNYWIFYGVNLCPY
70	CMYSFFNVCPY	216	MYCHQSMFGFRMCPD
71	CMYSFFNVCPA	217	NACAQMYGYSMCPHT
72	CTYSFFNVCPY	218	NDCDISMFDQSLCPY
73	CTYSFFNVCPA	219	NFSCVYVMFNVCLDP
74	GFPCMYSMFNVCLDP	220	NFTCALTMYEVCCLDP
75	GLSCMYSMYGYCLDP	221	NLCHAFMFGFNLCPY
76	IPCDYWMFGVNLCPY	222	NLNNCPHHPMCHDYQ
77	QVCHAYMYGYNLCPY	223	NPPCMYSFFNICLDP
78	RMYCTYSFYNVCLDP	224	NSACTYSFFNVCLDP
79	ALSCMYIMYNVCLDP	225	NVCTVSMFGVMLCPS
80	DFSCMYVMFNVCLDP	226	PACATLMYSVPLCPA
81	DFSCVYSMFNVCLDP	227	PAPCMYSFYNVCLDP
82	DMNTCPNHPMCYDYR	228	PLCAEMYGYSMCPHN
83	DMNTCPRHPMCHDYH	229	PQCHLYMYGYNLCPY
84	DSRCMYVMYNVCLDP	230	PRPCMYSFYNVCLDP
85	EHLCTYSFYNVCLDP	231	QHCPFQMYGYNLCPY
86	ELSCVYSMFGFCLDP	232	QHCQMFMFGYNLCPY
87	FTNNCPYHPMCHDYL	233	QHSCMYSSFFNVCLDP
88	GFSCYIMYDVCLDP	234	QKCHSYLYGVNLCPY
89	GSSCMYSMYNVCLDP	235	QKCNMFMFGYNLCPY
90	HFSCMYIMYNVCLDP	236	QMNDCPNHPMCHDYH
91	LHCGMWMFGVNLCPK	237	QPCAQMYGYSMCPAT
92	LPCQMWMFGHNLCPH	238	QPCAQMYGYSMCPRT
93	LPCTMYMYGYNLCPY	239	RECHFFFYGVNLCPY
94	LTCHHWFMFGVNLCPY	240	LNCGMFMGYNLCPY
95	NFSCMYSMFNVCLDP	241	RLCTSYMFGYNLCPQ
96	NNHCMYSFFNICLDP	242	RLSCMYSMFNVCLDP
97	NRSCMYIMYNVCLDP	243	RNCPFVMFGVNLCPY
98	NSCTMFMFGVNLCPY	244	RNGCMYSFFNVCLDP
99	NTCELYMFGVNLCPY	245	RNGCVYSFFNVCLDP
100	QHCDMWMFGYNLCPY	246	RPCHLYMFGYNLCPD
101	QHCPMYMFGYNLCPF	247	RPCHSYMYGINLCPY
102	QVCHIQMYGFDLCPH	248	RSCDMIMFGFNLCPY
103	RACDYWMYGVNLCPY	249	RSCPMWFYGVNLCPY
104	RQCHMQMFGYDLCPF	250	RSTVCFYDFCGPWER
105	SGSCLYSFYNVCLDP	251	RTCHFYMFGVNLCPY
106	SNGCTYSFFNVCLDP	252	RTCSMVMFGVNLCPY
107	STCAQMYGYSMCPH	253	SGKCTYSFFNVCLDP
108	SYKCLYSFYNVCLDP	254	SIVCDLYWEATCLRP
109	VLYCTYVMYNVCLDP	255	SLSCTYSFFNICLDP
110	VNCGMWMFGYNLCPK	256	SMNTCPYHPMCFDYK
111	YGSCLYSFYNICLDP	257	SQCWMWMYGYNLCPK
112	YPCAQMYGYSMCPHT	258	SSSCMYSSFFNVCLDP
113	AACDLWMFGVNLCPY	259	STACTYSFYNVCLDP
114	AFCTLAPYNQACIAN	260	STCAQMYGYSMCPHT
115	AGSCLYSMYNVCLDP	261	STRCVYSFYNVCLDP
116	ALCENTMYGYHLCPW	262	TACGAWMFGVNLCPY
117	ALSCMYIMYGVCLDP	263	TGACMYSFYNVCLDP
118	APVCDVLMFGFCMQP	264	TLSCMYSMYNVCLDP
119	AQVCSIMMYGTCLMP	265	TSCTVTMYQISMCPY

120	ASTCMYSFYNVCLDP	266	VGGCRHSFYNVCLDP
121	AVCEFWMFGFNLCPY	267	VHCQMYMYGYNLCPY
122	DANTCPNHPMCYDYH	268	VHNCMYSFFNVCLDP
123	DFSCIYIMFDVCLDP	269	VMCKLHMYGIPVCPK
124	DFSCMYVMYGFCLDP	270	VNFCNYSMYGICLLP
125	DFTCMYSMYNVCLDP	271	VNFCYACYCMSCVFS
126	DFTCTYSMYNVCLDP	272	VNQCTYSFFNVCLDP
127	DHYCTYIMYSICLDP	273	VPCPFHMFGYNLCPY
128	DICTNFMFGVNLCPY	274	VRCQMWMYGFNLCPH
129	DINTCPYHPMCHDYH	275	VRPCTYSFFNVCLDP
130	DKNTCPLHPMCHDYR	276	VSGCTYSFFNICLDP
131	DMNMCPNHPMCHDWH	277	YCSSWDTMTIPACNN
132	DMNSCPNHPMCHDYH	278	YDCDLSMFGIEMCPQ
133	DMNSCPNHPMCYDYR	279	YGNTCPFHMPMCHDYK
134	DMNTCPNHPMCFDYR	280	YGYCMYSFFNVCLDP
135	DMNTCPNHPMCHDFQ	281	YHCTMHMFGYNLCPF
136	DMNTCPNHPMCHDYR	282	YMNTCPNHPMCFDYQ
137	DMNTCPNHPMCYDYH	283	YMNTCPYHPMCHDYL
138	DMNTCPNHPMCYDYK	284	YMNTCPYHPMCHDYR
139	DMSTCPNHPMCHDYM	285	YNNCTYSFFNVCLDP
140	DRNMCPYHPMCYDYR	286	YPGCQYSFFNVCLDP
141	DSCAFMMFGVNLCPY	287	YRSCTHIMYNVCLDP
142	DSCRSVFDMMVWNCWN	288	YSFCDMLMYDVCLVP
143	DTPNCPHHPMCHNHM	289	YSIDCGLSWWCGGMT
144	DVSCLYVMYSVCLDP	290	YSTTCPYHPMCHDYH
145	DWCASMMFGYNLCPY	291	YVNTCPHHPMCHDYH
146	EFSCMYSMFNVCLDP	292	YVNTCPYHPMCHDYN

[0123] In some embodiments, the K_d of the activatable anti-CTLA-4 antibody, comprising a MM disclosed herein, towards the target is at least 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, 1,000 times greater than, or between 5-10, 10-100, 10-200, 10-500, 10-1,000 times greater than the K_d of the AB not modified with a MM or of the parental AB towards the target.

[0124] In some embodiments, the MM is not a natural binding partner of the activatable antibody. In some embodiments, the MM contains no or substantially no homology to any natural binding partner of the activatable antibody. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% identical to any natural binding partner of the activatable antibody. In some embodiments, the MM is no more than 50%, 25%, 20%, or 10% identical to any natural binding partner of the activatable antibody. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% identical to human CTLA-4. In some embodiments, the MM is no more than 50%, 25%, 20%, or 10% identical to human CTLA-4.

Exemplary Activatable anti-CTLA-4 Antibodies

- [0125] Particular antibodies described herein are activatable anti-CTLA-4 antibodies comprising any combination of the masking moieties, cleavable moieties, light chain variable domains (VL) (or the corresponding CDRs), and heavy chain variable domains (VH) (or the corresponding CDRs) provided in Tables 2-6. In some embodiments, the activatable anti-CTLA-4 antibody comprises a light chain comprising YV01 (SEQ ID NO: 1) as the masking moiety, LSGRSDNH (SEQ ID NO: 313) as the cleavable moiety, and the light chain variable domain (VL) of ipilimumab (SEQ ID NO: 344). In some embodiments, the activatable anti-CTLA-4 antibody comprises a light chain comprising YV01 (SEQ ID NO: 1) as the masking moiety, ISSGLLSGRSDNH (2001) (SEQ ID NO: 297) as the cleavable moiety, and the CDRs of the light chain variable domain (VL) of ipilimumab (SEQ ID NOs: 560, 561, and 562, respectively). In some embodiments, the activatable anti-CTLA-4 antibody comprises the heavy chain variable domain (VH) of ipilimumab (SEQ ID NO: 345) or just the corresponding CDRs (SEQ ID NOs: 557, 558, and 559).
- [0126] In some embodiments, the activatable anti-CTLA-4 comprises YV39 (SEQ ID NO: 39) as the masking moiety, and ISSGLLSGRSDNP ("2011") (SEQ ID NO: 304) as the cleavable moiety, and the heavy and light chain variable domains of ipilimumab ((SEQ ID NOs: 345 and 344, respectively), wherein the MM and CM are linked to the VL in the arrangement MM-CM-VL.
- [0127] In some embodiments, the activatable anti-CTLA-4 antibody includes a signal peptide. The signal peptide can be linked to the activatable anti-CTLA-4 antibody by a spacer. In some embodiments, the spacer is conjugated to the activatable antibody in the absence of a signal peptide. In some embodiments, the spacer is joined directly to the MM of the activatable antibody. In some embodiments, the spacer has amino acid sequence QGQSGS (SEQ ID NO: 546). In some embodiments, an activatable antibody comprises a spacer of sequence QGQSGS (SEQ ID NO: 546) joined directly to a MM sequence CRTQLYGYNLCPY (YV39) (SEQ ID NO: 39) in the structural arrangement from N-terminus to C-terminus of "spacer-MM-CM-VL" or "spacer-MM-CM-AB."
- [0128] In some embodiments, the activatable anti-CTLA-4 antibody comprises a linker peptide (LP) between the MM and the CM. In some embodiments, the activatable anti-CTLA-4 antibody comprises a linker peptide between the CM and the antibody or antigen

binding domain thereof (AB). In some embodiments, the activatable anti-CTLA-4 antibody comprises a first linker peptide (LP1) and a second linker peptide (LP2), and wherein the activatable anti-CTLA-4 antibody has the structural arrangement from N-terminus to C-terminus as follows: MM-LP1-CM-LP2-AB. In some embodiments, the light chain of the activatable anti-CTLA-4 antibody has the structural arrangement from N-terminus to C-terminus as follows: MM-LP1-CM-LP2-VL. In some embodiments, the two linker peptides need not be identical to each other. Examples of linker peptides that may be used with the activatable anti-CTLA-4 antibodies as disclosed herein are provided in U.S. Patent Publication No. 2016/0193332 and International Publication No. WO 2016/149201, *ibid*.

[0129] The disclosure also comprises a modified anti-CTLA-4 antibody that comprises a MM that is joined to the light chain of the antibody via a non-protease cleavable linker. In some embodiments, the non-protease cleavable linker comprises the amino acid sequence set forth in SEQ ID NO: 570. In some embodiments, such a modified anti-CTLA-4 antibody has a light chain comprising YV39 and a non-protease cleavable linker. In some embodiments, the light chain of the modified anti-CTLA-4 antibody comprises the amino acid sequence:

QGQSGSCRTQLYGYNLCPYGGGSSGGSGGGSGGGSGGGSGGGSGGGSEIVLT
QSPGTLSPGERATLSCRASQSVGSSYLAWYQQKPGQAPRLLIYGAFSR
ATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKV
EIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNA
LQSGNSQESVTEQDSKSTYLSSTLTLSKADYEKHKVYACEVTHQGLS
SPVTKSFNRGEC (SEQ ID NO: 530) or

CRTQLYGYNLCPYGGGSSGGSGGGSGGGSGGGSGGGSGGGSEIVLTQSPGTL
SLSPGERATLSCRASQSVGSSYLAWYQQKPGQAPRLLIYGAFSRATGIPD
RFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKVEIKRTV
AAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNS
QESVTEQDSKSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSF
NRGEC (SEQ ID NO: 531).

[0130] Linkers suitable for use in compositions described herein are generally ones that provide flexibility of the activatable anti-CTLA-4 antibody to facilitate the inhibition of the binding of the activatable antibody to the target. Such linkers are generally referred to as flexible linkers (also referred to as linker peptides herein). Suitable linkers can be readily selected and can be of any of a suitable of different lengths, such as from 1 amino acid (e.g. , Gly) to 20 amino acids, from 2 amino acids to 15 amino acids, from 3 amino acids to 12 amino acids, including 4 amino acids to 10 amino acids, 5 amino acids to 9 amino acids, 6 amino acids to 8 amino acids, or 7 amino acids to 8 amino acids, and may

be 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acids in length.

[0131] Exemplary flexible linkers include glycine polymers (G)_n, glycine-serine polymers (including, for example, (GS)_n, (GSGGS)_n (GSGGS is SEQ ID NO: 534) and (GGGS)_n (GGGS is SEQ ID NO: 535), where n is an integer of at least one), glycine-alanine polymers, alanine-serine polymers, and other flexible linkers known in the art. Glycine and glycine-serine polymers are relatively unstructured, and therefore may be able to serve as a neutral tether between components. Glycine accesses significantly more phi-psi space than even alanine, and is much less restricted than residues with longer side chains (*see* Scheraga, *Rev. Computational Chem.* 11173-142 (1992)). Exemplary flexible linkers include, but are not limited to Gly-Gly-Ser-Gly (SEQ ID NO: 536), Gly-Gly-Ser-Gly-Gly (SEQ ID NO: 537), Gly-Ser-Gly-Ser-Gly (SEQ ID NO: 538), Gly-Ser-Gly-Gly-Gly (SEQ ID NO: 539), Gly-Gly-Gly-Ser-Gly (SEQ ID NO: 540), Gly-Ser-Ser-Ser-Gly (SEQ ID NO: 541), and the like. The ordinarily skilled artisan will recognize that design of an activatable antibodies can include linkers that are all or partially flexible, such that the linker can include a flexible linker as well as one or more portions that confer less flexible structure to provide for a desired activatable antibodies structure.

[0132] In some embodiments, the activatable anti-CTLA-4 antibodies comprise the VL and VH (or the corresponding CDRs) of ipilimumab and a combination of MMs and CMs provided in Table 5 below, such that any MM in column 2 can be combined with any CM in column 4.

Table 5. Activatable anti-CTLA-4 Antibody Combinations

SEQ ID NO.	Masking Moiety (MM)	SEQ ID NO.	Cleavable Moiety (CM)
1	(YV01) DFSLHSMYNVCLDP	313	LSGRSDNH
2	(YV02) QPCAQMYGYSMCPHT	314	LSGRSANPRG
3	(YV03) LHCRTQMYGYNLCPY	315	TGRGPSWV
4	(YV04) LHCRTQLYGYNLCPY	316	PLTGRSGG
5	(YV05) CTYSFFNVC	317	TARGPSFK
6	(YV06) CAQMYGYSMC	318	NLSGRSENHSG
7	(YV07) CPNHPMC	319	NLSGRSGNHGS
8	(YV08) GTACTYSFFNVCLDP	320	TSTSGRSANPRG
9	(YV09) FGTACPNHPMCHDWQ	321	TSGRSANP
10	(YV10) SACAYWMFGVNLCPY	322	VHMPGLGLGP
11	(YV11) CRTQLYGYNLC	323	AVGLLAPP
12	(YV12) CRTQIYGYNLC		
13	(YV13) LHCRTQIYGYNLCPY		

14	(YV14) CPNHPMCHDWQ	324	AQNLLGMV
15	(YV15) GTACPNHPMCHDWQ	325	QNQALRMA
16	(YV16) CAYWMFGVNLCPY	326	LAAPLGLL
17	(YV17) QECHLYMYGVNLCPY	327	STFPFGMF
18	(YV18) CHLYMYGVNLCPY	328	ISSGLLSS
19	(YV19) GQCQFYMFGYNLCPY	329	PAGLWLDP
20	(YV20) LSTCMYSFFNVCLDP	330	VAGRSMRP
21	(YV21) CLHSMYNVCLDP	331	VVPEGRRS
22	(YV22) CLHSMYNV	332	ILPRSPAF
23	(YV23) CLHSLYNVCLDP	333	MVLGRSLL
24	(YV24) CLHSAYNVCLDP	334	VAGRSMRP
25	(YV25) CMYSFFNVCLDP	335	QGRAITFI
26	(YV26) CMYSFFNV	336	SPRSIMLA
27	(YV27) QPCAQMYGYSMC	337	SMLRSMPL
28	(YV28) CAQLYGYSMCPHT	297	ISSGLLSGRSDNH
29	(YV29) CAQMYGYSMCAHT	300	ISSGLLSGRSDDH
30	(YV30) CAQMYGYSMCPAT	301	ISSGLLSGRSDIH
31	(YV31) CAQMYGYSMCPHT	302	ISSGLLSGRSDQH
32	(YV32) CPNHPLCHDWQ	303	ISSGLLSGRSDTH
33	(YV33) CPNHPMCADWQ	341	ISSGLLSGRSDYH
35	(YV34) CPNHPMCHAWQ	342	ISSGLLSGRSANI
35	(YV35) CPNHPMCHDAQ	343	ISSGLLSGRSDNI
36	(YV36) CPNHPMCHDWA	305	ISSGLLSGRSANP
37	(YV37) GTACPNHPMC	304	ISSGLLSGRSDNP
38	(YV38) LHCRTQLYGYNLC	298	ISSGLLSGRSANPRG
39	(YV39) CRTQLYGYNLCPY	306	AVGLLAPPGGLSGRSDNH
40	(YV40) CRTQLYGYNLCAY	307	AVGLLAPPGGLSGRSDDH
41	(YV41) CRTQLYGYNLCAPA	308	AVGLLAPPGGLSGRSDIH
42	(YV42) FGTACPNHPLCHDWQ	309	AVGLLAPPGGLSGRSDQH
43	(YV43) CPNHPLCHDFQ	310	AVGLLAPPGGLSGRSDTH
44	(YV44) CPNHPLCHDYQ	338	AVGLLAPPGGLSGRSDYH
45	(YV45) CPNHPLCPY	339	AVGLLAPPGGLSGRSANI
46	(YV46) CPNHPLCPA	340	AVGLLAPPGGLSGRSDNI
47	(YV47) CMYSFFNVCYP	312	AVGLLAPPGGLSGRSANP
48	(YV48) CMYSFFNVCYA	311	AVGLLAPPGGLSGRSDNP
49	(YV49) CLYSFFNVCYP	299	AVGLLAPPSGRSANPRG
50	(YV50) CLYSFFNVCYA		
51	(YV51) FGAACPNHPICHDWQ		
52	(YV52) FGAACPNHPLCHDWQ		
53	(YV53) FGAACPNHPMCHDAQ		
54	(YV54) CLHSAYNACLDP		
55	(YV55) CAHSAYNVCLDP		
56	(YV56) CLHSAYNVCADP		
57	(YV57) CLHSAYNVCLAP		
58	(YV58) CLHSAYNVCLDA		
59	(YV60) KNTCTYVMYNVCLDP		
60	(YV61) YISDCPYHPMCHDYQ		
61	(YV62) FRNTCPYHPMCHDYR		
62	(YV63) RECHMWMFGVNLCPY		
63	(YV64) AVCHMYMYGYNLCPF		
64	(YV65) RSCPQMYGYSMCPHT		
65	(YV66) QPCAQMFGYSMCPHT		

[0133] In some embodiments, the activatable anti-CTLA-4 antibodies comprise the specific combination of MMs and CMs provided in Table 6 below.

Table 6. Exemplary Activatable Anti-CTLA-4 Antibody Combination

Comb. No.	Masking Moiety (MM)	Cleavable Moiety (CM)
1	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSDNH (SEQ ID NO: 297)
2	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSDNP (SEQ ID NO: 304)
3	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSANP (SEQ ID NO: 305)
4	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSDQH (SEQ ID NO: 302)
5	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSDDH (SEQ ID NO: 300)
6	CRTQLYGYNLCPY (SEQ ID NO: 39)	ISSGLLSGRSDTH (SEQ ID NO: 303)
7	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	ISSGLLSGRSDNH (SEQ ID NO: 297)
8	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
9	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	ISSGLLSGRSDDH (SEQ ID NO: 300)
10	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	ISSGLLSGRSDIH (SEQ ID NO: 301)
11	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	ISSGLLSGRSDQH (SEQ ID NO: 302)
12	LHCRTQMYGYNLCPY (SEQ ID NO: 3)	ISSGLLSGRSDTH (SEQ ID NO: 303)
13	CAQMYGYSMC (SEQ ID NO: 06)	ISSGLLSGRSDNH (SEQ ID NO: 297)
14	CAQMYGYSMC (SEQ ID NO: 06)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
15	FGTACPNHPMCHDWQ (SEQ ID NO: 09)	ISSGLLSGRSDNH (SEQ ID NO: 297)
16	FGTACPNHPMCHDWQ (SEQ ID NO: 09)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
17	CLHSLYNVCLDP (SEQ ID NO: 23)	ISSGLLSGRSDNH (SEQ ID NO: 297)
18	CLHSLYNVCLDP (SEQ ID NO: 23)	ISSGLLSGRSDDH (SEQ ID NO: 300)
19	CLHSLYNVCLDP (SEQ ID NO: 23)	ISSGLLSGRSDIH (SEQ ID NO: 301)
20	CLHSLYNVCLDP (SEQ ID NO: 23)	ISSGLLSGRSDQH (SEQ ID NO: 302)

21	CLHSLYNVCLDP (SEQ ID NO: 23)	ISSGLLSGRSDTH (SEQ ID NO: 303)
22	CLHSLYNVCLDP (SEQ ID NO: 23)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
23	CLHSAYNVCLDP (SEQ ID NO: 24)	ISSGLLSGRSDNH (SEQ ID NO: 297)
24	CLHSAYNVCLDP (SEQ ID NO: 24)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
25	QPCAQMYGYSMC (SEQ ID NO: 27)	ISSGLLSGRSDNH (SEQ ID NO: 297)
26	QPCAQMYGYSMC (SEQ ID NO: 27)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
27	CAQMYGYSMCAHT (SEQ ID NO: 29)	ISSGLLSGRSDNH (SEQ ID NO: 297)
28	CAQMYGYSMCAHT (SEQ ID NO: 29)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
29	CPNHPLCHDWQ (SEQ ID NO: 32)	ISSGLLSGRSDNH (SEQ ID NO: 297)
30	CPNHPLCHDWQ (SEQ ID NO: 32)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
31	CPNHPMCADWQ (SEQ ID NO: 33)	ISSGLLSGRSDNH (SEQ ID NO: 297)
32	CPNHPMCADWQ (SEQ ID NO: 33)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
33	CPNHPMCHDAQ (SEQ ID NO: 35)	ISSGLLSGRSDNH (SEQ ID NO: 297)
34	CPNHPMCHDAQ (SEQ ID NO: 35)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
35	CRTQLYGYNLCPY (SEQ ID NO: 39)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
36	CRTQLYGYNLCPA (SEQ ID NO: 41)	ISSGLLSGRSDNH (SEQ ID NO: 297)
37	CRTQLYGYNLCPA (SEQ ID NO: 41)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
38	FGAACPNHPICHDWQ (SEQ ID NO: 51)	ISSGLLSGRSDNH (SEQ ID NO: 297)
39	FGAACPNHPICHDWQ (SEQ ID NO: 51)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
40	FGAACPNHPLCHDWQ (SEQ ID NO: 52)	ISSGLLSGRSDNH (SEQ ID NO: 297)
41	FGAACPNHPLCHDWQ (SEQ ID NO: 52)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
42	FGAACPNHPMCHDAQ (SEQ ID NO: 53)	ISSGLLSGRSDNH (SEQ ID NO: 297)
43	FGAACPNHPMCHDAQ (SEQ ID NO: 53)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
44	CLHSAYNACLDP (SEQ ID NO: 54)	ISSGLLSGRSDNH (SEQ ID NO: 297)

45	CLHSAYNACLDP (SEQ ID NO: 54)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
46	CAHSAYNVCLDP (SEQ ID NO: 55)	ISSGLLSGRSDNH (SEQ ID NO: 297)
47	CAHSAYNVCLDP (SEQ ID NO: 55)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
48	CLHSAYNVCADP (SEQ ID NO: 56)	ISSGLLSGRSDNH (SEQ ID NO: 297)
49	CLHSAYNVCADP (SEQ ID NO: 56)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
50	CLHSAYNVCLAP (SEQ ID NO: 57)	ISSGLLSGRSDNH (SEQ ID NO: 297)
51	CLHSAYNVCLAP (SEQ ID NO: 57)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
52	CLHSAYNVCLDA (SEQ ID NO: 58)	ISSGLLSGRSDNH (SEQ ID NO: 297)
53	CLHSAYNVCLDA (SEQ ID NO: 58)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306)
54	YISDCPYHPMCHDYQ (SEQ ID NO: 60)	ISSGLLSGRSDNH (SEQ ID NO: 297)
55	FRNTCPYHPMCHDYR (SEQ ID NO: 61)	ISSGLLSGRSDNH (SEQ ID NO: 297)
56	AVCHMYMYGYNLCPF (SEQ ID NO: 63)	ISSGLLSGRSDNH (SEQ ID NO: 297)
57	RSCPQMYGYSMCPHT (SEQ ID NO: 64)	ISSGLLSGRSANP (SEQ ID NO: 305)
58	QPCAQMFGYSMCPHT (SEQ ID NO: 65)	ISSGLLSGRSANP (SEQ ID NO: 305)

[0134] In some embodiments, the activatable anti-CTLA-4 antibodies described herein also include an agent conjugated to the activatable antibody. In some embodiments, the conjugated agent is a therapeutic agent, such as an anti-neoplastic agent. In some embodiments, the agent is conjugated to a carbohydrate moiety of the activatable antibody, preferably where the carbohydrate moiety is located outside the antigen-binding region of the antibody or antigen-binding fragment in the activatable antibody. In some embodiments, the agent is conjugated to a sulfhydryl group of the antibody or antigen-binding fragment in the activatable antibody. In some embodiments, the agent is conjugated to an amino group of the antibody or antigen-binding fragment of the activatable antibody. In some embodiments, the agent is conjugated to a carboxylic acid group of the antibody or antigen-binding fragment of the activatable antibody.

- [0135] In some embodiments, the agent is a cytotoxic agent such as a toxin (e.g., an enzymatically active toxin of bacterial, fungal, plant, or animal origin, or fragments thereof), or a radioactive isotope (i.e., a radioconjugate).
- [0136] In some embodiments, the conjugated activatable antibody can be modified for site-specific conjugation through modified amino acid sequences inserted or otherwise included in the activatable antibody sequence. These modified amino acid sequences are designed to allow for controlled placement and/or dosage of the conjugated agent within a conjugated activatable anti-CTLA-4 antibody. For example, the activatable antibody can be engineered to include cysteine substitutions at positions on light and heavy chains that provide reactive thiol groups and do not negatively impact protein folding and assembly, nor alter antigen binding. In some embodiments, the activatable antibody can be engineered to include or otherwise introduce one or more non-natural amino acid residues within the activatable antibody to provide suitable sites for conjugation. In some embodiments, the activatable antibody can be engineered to include or otherwise introduce enzymatically activatable peptide sequences within the activatable antibody sequence.
- [0137] In some embodiments, the agent is a detectable moiety such as, for example, a label or other marker. For example, the agent is or includes a radiolabeled amino acid, one or more biotinyl moieties that can be detected by marked avidin (e.g., streptavidin containing a fluorescent marker or enzymatic activity that can be detected by optical or calorimetric methods), one or more radioisotopes or radionuclides, one or more fluorescent labels, one or more enzymatic labels, and/or one or more chemiluminescent agents. In some embodiments, detectable moieties are attached by linker molecules.
- [0138] Conjugates of the antibody and cytotoxic agent are made using a variety of bifunctional protein-coupling agents such as N-succinimidyl-3-(2-pyridyldithiol) propionate (SPDP), iminothiolane (IT), bifunctional derivatives of imidoesters (such as dimethyl adipimidate HCL), active esters (such as disuccinimidyl suberate), aldehydes (such as glutaraldehyde), bis-azido compounds (such as bis (p-azidobenzoyl) hexanediamine), bis-diazonium derivatives (such as bis-(p-diazoniumbenzoyl)-ethylenediamine), diisocyanates (such as toluene 2,6-diisocyanate), and bis-active fluorine compounds (such as 1,5-difluoro-2,4-dinitrobenzene). For example, a ricin immunotoxin can be prepared as described in Vitetta *et al.*, *Science* 238: 1098 (1987).

Carbon-14-labeled 1-isothiocyanatobenzyl-3-methyldiethylene triaminepentaacetic acid (MX-DTPA) is an exemplary chelating agent for conjugation of radionucleotide to the antibody. (See WO94/11026).

[0139] Those of ordinary skill in the art will recognize that a large variety of possible moieties can be coupled to the resultant antibodies of the invention. (*See, e.g.*, "Conjugate Vaccines", Contributions to Microbiology and Immunology, J. M. Cruse and R. E. Lewis, Jr (eds), Carger Press, New York, (1989), the entire contents of which are incorporated herein by reference).

II. Uses of Anti-CTLA-4 Activatable Antibodies

[0140] Therapeutic formulations of the invention, which include an activatable anti-CTLA-4 antibody, are used to prevent, treat or otherwise ameliorate a disease or disorder, including but not limited to, a disease or disorder associated with aberrant CTLA-4 expression and/or activity. For example, therapeutic formulations of the invention, which include an activatable anti-CTLA-4 antibody, are used as cancer immunotherapy, e.g., potentiating an endogenous immune response in a subject afflicted with a cancer so as to thereby treat the subject, which method comprises administering to the subject therapeutically effective amount of any of the activatable anti-CTLA-4 antibodies described herein.

[0141] Examples of cancers that may be treated using the immunotherapeutic methods of the disclosure include bone cancer, pancreatic cancer, skin cancer, cancer of the head or neck, breast cancer, lung cancer, cutaneous or intraocular malignant melanoma, unresectable or metastatic melanoma, renal cancer, uterine cancer, ovarian cancer, colorectal cancer, colon cancer, rectal cancer, cancer of the anal region, stomach cancer, testicular cancer, uterine cancer, carcinoma of the fallopian tubes, carcinoma of the endometrium, carcinoma of the cervix, carcinoma of the vagina, carcinoma of the vulva, cancer of the esophagus, cancer of the small intestine, cancer of the endocrine system, cancer of the thyroid gland, cancer of the parathyroid gland, cancer of the adrenal gland, sarcoma of soft tissue, cancer of the urethra, cancer of the penis, a hematological malignancy, solid tumors of childhood, lymphocytic lymphoma, cancer of the bladder, cancer of the kidney or ureter, carcinoma of the renal pelvis, neoplasm of the central nervous system (CNS), primary CNS lymphoma, tumor angiogenesis, spinal axis tumor,

brain stem glioma, pituitary adenoma, Kaposi's sarcoma, epidermoid cancer, squamous cell cancer, environmentally induced cancers including those induced by asbestos, metastatic cancers, and any combinations of said cancers. In some embodiments, the cancer is selected from MEL, RCC, squamous NSCLC, non-squamous NSCLC, CRC, CRPC, squamous cell carcinoma of the head and neck, and carcinomas of the esophagus, ovary, gastrointestinal tract and breast. The present methods are also applicable to treatment of metastatic cancers.

[0142] Other cancers include hematologic malignancies including, for example, multiple myeloma, B-cell lymphoma, Hodgkin lymphoma/primary mediastinal B-cell lymphoma, non-Hodgkin's lymphomas, acute myeloid lymphoma, chronic myelogenous leukemia, chronic lymphoid leukemia, follicular lymphoma, diffuse large B-cell lymphoma, Burkitt's lymphoma, immunoblastic large cell lymphoma, precursor B-lymphoblastic lymphoma, mantle cell lymphoma, acute lymphoblastic leukemia, mycosis fungoides, anaplastic large cell lymphoma, T-cell lymphoma, and precursor T-lymphoblastic lymphoma, and any combinations of said cancers.

[0143] Increased proteolysis is known to be a hallmark of cancer. (*See e.g., Affara N I, et al. "Delineating protease functions during cancer development." Methods Mol Biol. 539 (2009): 1-32.* Progression, invasion and metastasis of tumors result from several interdependent processes in which proteases are implicated. This process is described generally in U.S. Publication No. 2016/0193332 A1, which is incorporated in its entirety.

[0144] In some embodiments of these methods for treating a cancer subject, the activatable antibodies of the present invention, e.g. activatable ipilimumab, is administered to the subject as monotherapy. In some embodiments, stimulation or blockade of immunomodulatory targets may be effectively combined with standard cancer treatments, including chemotherapeutic regimes, radiation, surgery, hormone deprivation and angiogenesis inhibitors. The activatable anti-CTLA-4 antibody can be linked to an anti-neoplastic agent (as an immunoconjugate) or can be administered separately from the agent. In the latter case (separate administration), the antibody can be administered before, after or concurrently with the agent or can be co-administered with other known therapeutic agents. Chemotherapeutic drugs include, among others, doxorubicin (ADRIAMYCIN®), cisplatin, carboplatin, bleomycin sulfate, carmustine, chlorambucil (LEUKERAN®), cyclophosphamide (CYTOXAN®; NEOSAR®),

lenalidomide (REVLIMID®), bortezomib (VELCADE®), dexamethasone, mitoxantrone, etoposide, cytarabine, bendamustine (TREANDA®), rituximab (RITUXAN®), ifosfamide, vincristine (ONCOVIN®), fludarabine (FLUDARA®), thalidomide (THALOMID®), alemtuzumab (CAMPATH®), ofatumumab (ARZERRA®), everolimus (AFINITOR®, ZORTRESS®), and carfilzomib (KYPROLISTM). Co-administration of anti-cancer agents that operate via different mechanisms can help overcome the development of resistance to drugs or changes in the antigenicity of tumor cells.

[0145] Activatable anti-CTLA-4 antibodies of the present invention, such as the activatable ipilimumab, may also be used in combination with other immunomodulatory agents, such as antibodies against other immunomodulatory receptors or their ligands. Several other co-stimulatory and inhibitory receptors and ligands that regulate T cell responses have been identified. Examples of stimulatory receptors include Inducible T cell Co-Stimulator (ICOS), CD137 (4-1BB), CD134 (OX40), CD27, Glucocorticoid-Induced TNFR-Related protein (GITR), and Herpes Virus Entry Mediator (HVEM), whereas examples of inhibitory receptors include Programmed Death-1 (PD-1), Programmed Death Ligand-1 (PD-L1), B and T Lymphocyte Attenuator (BTLA), T cell Immunoglobulin and Mucin domain-3 (TIM-3), Lymphocyte Activation Gene-3 (LAG-3), adenosine A2a receptor (A2aR), Killer cell Lectin-like Receptor G1 (KLRG-1), Natural Killer Cell Receptor 2B4 (CD244), CD160, T cell Immunoreceptor with Ig and ITIM domains (TIGIT), and the receptor for V-domain Ig Suppressor of T cell Activation (VISTA). Mellman *et al.* (2011) *Nature* 480:480; Pardoll (2012) *Nat. Rev. Cancer* 12:252; Baitsch *et al.* (2012) *PloS One* 7:e30852.

[0146] Anti-PD-1 antibodies OPDIVO® (nivolumab) and KEYTRUDA® (pembrolizumab), as well as anti-PD-L1 antibody TECENTRIQ® (atezolizumab), have been approved for use in treating cancer, and may be combined with the activatable anti-CTLA-4 antibodies of the present invention, e.g. activatable ipilimumab. These receptors and their ligands provide targets for therapeutics designed to stimulate, or prevent the suppression, of an immune response so as to thereby attack tumor cells. Weber (2010) *Semin. Oncol.* 37:430; Flies *et al.* (2011) *Yale J. Biol. Med.* 84:409; Mellman *et al.* (2011) *Nature* 480:480; Pardoll (2012) *Nat. Rev. Cancer* 12:252. Stimulatory receptors or receptor ligands are targeted by agonist agents, whereas inhibitory receptors or receptor ligands are targeted by blocking agents. Among the most promising approaches to

enhancing immunotherapeutic anti-tumor activity is the blockade of so-called "immune checkpoints," which refer to the plethora of inhibitory signaling pathways that regulate the immune system and are crucial for maintaining self-tolerance and modulating the duration and amplitude of physiological immune responses in peripheral tissues in order to minimize collateral tissue damage. *See e.g. Weber (2010) Semin. Oncol. 37:430; Pardoll (2012) Nat. Rev. Cancer 12:252.* Because many of the immune checkpoints are initiated by ligand-receptor interactions, they can be readily blocked by antibodies or modulated by recombinant forms of ligands or receptors.

Anti-PD-1 Antibodies Useful for the Invention

[0147] Any anti-PD-1 antibody that is known in the art can be used in the presently described methods. In particular, various human monoclonal antibodies that bind specifically to PD-1 with high affinity have been disclosed in U.S. Patent No. 8,008,449. Each of the anti-PD-1 humanized antibodies disclosed in U.S. Patent No. 8,008,449 has been demonstrated to exhibit one or more of the following characteristics: (a) binds to human PD-1 with a K_D of 1×10^{-7} M or less, as determined by surface plasmon resonance using a Biacore biosensor system; (b) does not substantially bind to human CD28, CTLA-4 or ICOS; (c) increases T-cell proliferation in a Mixed Lymphocyte Reaction (MLR) assay; (d) increases interferon- γ production in an MLR assay; (e) increases IL-2 secretion in an MLR assay; (f) binds to human PD-1 and cynomolgus monkey PD-1; (g) inhibits the binding of PD-L1 and/or PD-L2 to PD-1; (h) stimulates antigen-specific memory responses; (i) stimulates antibody responses; and (j) inhibits tumor cell growth *in vivo*. Anti-PD-1 antibodies usable in the present invention include monoclonal antibodies that bind specifically to human PD-1 and exhibit at least one, in some embodiments, at least five, of the preceding characteristics.

[0148] Other anti-PD-1 monoclonal antibodies have been described in, for example, U.S. Patent Nos. 6,808,710, 7,488,802, 8,168,757 and 8,354,509, US Publication No. 2016/0272708, and PCT Publication Nos. WO 2012/145493, WO 2008/156712, WO 2015/112900, WO 2012/145493, WO 2015/112800, WO 2014/206107, WO 2015/35606, WO 2015/085847, WO 2014/179664, WO 2017/020291, WO 2017/020858, WO 2016/197367, WO 2017/024515, WO 2017/025051, WO 2017/123557, WO 2016/106159, WO 2014/194302, WO 2017/040790, WO 2017/133540, WO

2017/132827, WO 2017/024465, WO 2017/025016, WO 2017/106061, each of which is incorporated by reference in its entirety.

[0149] In some embodiments, the anti-PD-1 antibody is selected from the group consisting of nivolumab (also known as "OPDIVO®"; formerly designated 5C4, BMS-936558, MDX-1106, or ONO-4538), pembrolizumab (Merck, also known as "KEYTRUDA®", lambrolizumab, and MK-3475. *See* WO2008156712A1), PDR001 (Novartis; *see* WO 2015/112900), MEDI-0680 (AstraZeneca; AMP-514; *see* WO 2012/145493), REGN-2810 (Regeneron; *see* WO 2015/112800), JS001 (TAIZHOU JUNSHI PHARMA; *see* Si-Yang Liu et al., *J. Hematol. Oncol.* 10:136 (2017)), BGB-A317 (Beigene; *see* WO 2015/35606 and US 2015/0079109), INCSHR1210 (SHR-1210; Jiangsu Hengrui Medicine; *see* WO 2015/085847; Si-Yang Liu et al., *J. Hematol. Oncol.* 10:136 (2017)), TSR-042 (ANB011; Tesaro Biopharmaceutical; *see* WO2014/179664), GLS-010 (WBP3055; Wuxi/Harbin Gloria Pharmaceuticals; *see* Si-Yang Liu et al., *J. Hematol. Oncol.* 10:136 (2017)), AM-0001 (Armo), STI-1110 (Sorrento Therapeutics; *see* WO 2014/194302), AGEN2034 (Agenus; *see* WO 2017/040790), and MGD013 (Macrogenics).

[0150] In one embodiment, the anti-PD-1 antibody is nivolumab. Nivolumab is a fully human IgG4 (S228P) PD-1 immune checkpoint inhibitor antibody that selectively prevents interaction with PD-1 ligands (PD-L1 and PD-L2), thereby blocking the down-regulation of antitumor T-cell functions (U.S. Patent No. 8,008,449; Wang et al., 2014 *Cancer Immunol Res.* 2(9):846-56).

[0151] In another embodiment, the anti-PD-1 antibody is pembrolizumab. Pembrolizumab is a humanized monoclonal IgG4 antibody directed against human cell surface receptor PD-1 (programmed death-1 or programmed cell death-1). Pembrolizumab is described, for example, in U.S. Patent Nos. 8,354,509 and 8,900,587; *see also* www.cancer.gov/drugdictionary?cdrid=695789 (last accessed: December 14, 2014). Pembrolizumab has been approved by the FDA for the treatment of relapsed or refractory melanoma.

[0152] Anti-PD-1 antibodies usable in the disclosed methods also include isolated antibodies that bind specifically to human PD-1 and cross-compete for binding to human PD-1 with any anti-PD-1 antibody disclosed herein, e.g., nivolumab (*see, e.g.,* U.S. Patent No. 8,008,449 and 8,779,105; WO 2013/173223). In some embodiments, the anti-PD-1

antibody binds the same epitope as any of the anti-PD-1 antibodies described herein, e.g., nivolumab. The ability of antibodies to cross-compete for binding to an antigen indicates that these monoclonal antibodies bind to the same epitope region of the antigen and sterically hinder the binding of other cross-competing antibodies to that particular epitope region. These cross-competing antibodies are expected to have functional properties very similar those of the reference antibody, e.g., nivolumab, by virtue of their binding to the same epitope region of PD-1. Cross-competing antibodies can be readily identified based on their ability to cross-compete with nivolumab in standard PD-1 binding assays such as Biacore analysis, ELISA assays or flow cytometry (*see, e.g.*, WO 2013/173223).

[0153] In certain embodiments, the antibodies that cross-compete for binding to human PD-1 with, or bind to the same epitope region of human PD-1 antibody, nivolumab, are monoclonal antibodies. For administration to human subjects, these cross-competing antibodies are chimeric antibodies, engineered antibodies, or humanized or human antibodies. Such chimeric, engineered, humanized or human monoclonal antibodies can be prepared and isolated by methods well known in the art.

[0154] Anti-PD-1 antibodies usable in the methods of the disclosed invention also include antigen-binding portions of the above antibodies. It has been amply demonstrated that the antigen-binding function of an antibody can be performed by fragments of a full-length antibody.

[0155] Anti-PD-1 antibodies suitable for use in the disclosed methods or compositions are antibodies that bind to PD-1 with high specificity and affinity, block the binding of PD-L1 and or PD-L2, and inhibit the immunosuppressive effect of the PD-1 signaling pathway. In any of the compositions or methods disclosed herein, an anti-PD-1 "antibody" includes an antigen-binding portion or fragment that binds to the PD-1 receptor and exhibits the functional properties similar to those of whole antibodies in inhibiting ligand binding and up-regulating the immune system. In certain embodiments, the anti-PD-1 antibody or antigen-binding portion thereof cross-competes with nivolumab for binding to human PD-1.

Anti-PD-L1 Antibodies Useful for the Invention

[0156] Any anti-PD-L1 antibody can be used in the methods of the present disclosure. Examples of anti-PD-L1 antibodies useful in the methods of the present disclosure include the antibodies disclosed in US Patent No. 9,580,507. Each of the anti-PD-L1

human monoclonal antibodies disclosed in U.S. Patent No. 9,580,507 have been demonstrated to exhibit one or more of the following characteristics: (a) binds to human PD-L1 with a K_D of 1×10^{-7} M or less, as determined by surface plasmon resonance using a Biacore biosensor system; (b) increases T-cell proliferation in a Mixed Lymphocyte Reaction (MLR) assay; (c) increases interferon- γ production in an MLR assay; (d) increases IL-2 secretion in an MLR assay; (e) stimulates antibody responses; and (f) reverses the effect of T regulatory cells on T cell effector cells and/or dendritic cells. Anti-PD-L1 antibodies usable in the present invention include monoclonal antibodies that bind specifically to human PD-L1 and exhibit at least one, in some embodiments, at least five, of the preceding characteristics.

- [0157] In certain embodiments, the anti-PD-L1 antibody is selected from the group consisting of BMS-936559 (formerly 12A4 or MDX-1105; *see, e.g.*, U.S. Patent No. 7,943,743 and WO 2013/173223), MPDL3280A (also known as RG7446, atezolizumab, and TECENTRIQ®; US 8,217,149; *see, also*, Herbst et al. (2013) J Clin Oncol 31(suppl):3000), durvalumab (IMFINZI™; MEDI-4736; AstraZeneca; *see* WO 2011/066389), avelumab (Pfizer; MSB-0010718C; BAVENCIO®; *see* WO 2013/079174), STI-1014 (Sorrento; *see* WO2013/181634), CX-072 (CytomX; *see* WO2016/149201), KN035 (3D Med/Alphamab; *see* Zhang et al., *Cell Discov.* 7:3 (March 2017), LY3300054 (Eli Lilly Co.; *see, e.g.*, WO 2017/034916), and CK-301 (Checkpoint Therapeutics; *see* Gorelik et al., AACR:Abstract 4606 (Apr 2016)).
- [0158] In certain embodiments, the PD-L1 antibody is atezolizumab (TECENTRIQ®). Atezolizumab is a fully humanized IgG1 monoclonal anti-PD-L1 antibody.
- [0159] In certain embodiments, the PD-L1 antibody is durvalumab (IMFINZI™). Durvalumab is a human IgG1 kappa monoclonal anti-PD-L1 antibody.
- [0160] In certain embodiments, the PD-L1 antibody is avelumab (BAVENCIO®). Avelumab is a human IgG1 lambda monoclonal anti-PD-L1 antibody.
- [0161] In other embodiments, the anti-PD-L1 monoclonal antibody is selected from the group consisting of 28-8, 28-1, 28-12, 29-8, 5H1, and any combination thereof.
- [0162] Anti-PD-L1 antibodies usable in the disclosed methods also include isolated antibodies that bind specifically to human PD-L1 and cross-compete for binding to human PD-L1 with any anti-PD-L1 antibody disclosed herein, *e.g.*, atezolizumab and/or avelumab. In some embodiments, the anti-PD-L1 antibody binds the same epitope as any

of the anti-PD-L1 antibodies described herein, *e.g.*, atezolizumab and/or avelumab. The ability of antibodies to cross-compete for binding to an antigen indicates that these antibodies bind to the same epitope region of the antigen and sterically hinder the binding of other cross-competing antibodies to that particular epitope region. These cross-competing antibodies are expected to have functional properties very similar those of the reference antibody, *e.g.*, atezolizumab and/or avelumab, by virtue of their binding to the same epitope region of PD-L1. Cross-competing antibodies can be readily identified based on their ability to cross-compete with atezolizumab and/or avelumab in standard PD-L1 binding assays such as Biacore analysis, ELISA assays or flow cytometry (*see, e.g.*, WO 2013/173223).

[0163] In certain embodiments, the antibodies that cross-compete for binding to human PD-L1 with, or bind to the same epitope region of human PD-L1 antibody as, atezolizumab and/or avelumab, are monoclonal antibodies. For administration to human subjects, these cross-competing antibodies are chimeric antibodies, engineered antibodies, or humanized or human antibodies. Such chimeric, engineered, humanized or human monoclonal antibodies can be prepared and isolated by methods well known in the art.

[0164] Anti-PD-L1 antibodies usable in the methods of the disclosed invention also include antigen-binding portions of the above antibodies. It has been amply demonstrated that the antigen-binding function of an antibody can be performed by fragments of a full-length antibody.

[0165] Anti-PD-L1 antibodies suitable for use in the disclosed methods or compositions are antibodies that bind to PD-L1 with high specificity and affinity, block the binding of PD-1, and inhibit the immunosuppressive effect of the PD-1 signaling pathway. In any of the compositions or methods disclosed herein, an anti-PD-L1 "antibody" includes an antigen-binding portion or fragment that binds to PD-L1 and exhibits the functional properties similar to those of whole antibodies in inhibiting receptor binding and up-regulating the immune system. In certain embodiments, the anti-PD-L1 antibody or antigen-binding portion thereof cross-competes with atezolizumab and/or avelumab for binding to human PD-L1.

[0166] Efficaciousness of prevention, amelioration or treatment is determined in association with any known method for diagnosing or treating the disease or disorder, including but not limited to, a disease or disorder associated with aberrant CTLA-4

expression and/or activity. Prolonging the survival of a subject or otherwise delaying the progression of the disease or disorder, including but not limited to, a disease or disorder associated with aberrant CTLA-4 expression and/or activity in a subject, indicates that the activatable antibody confers a clinical benefit.

[0167] It will be appreciated that therapeutic entities in accordance with the invention will be administered with suitable carriers, excipients, and other agents that are incorporated into formulations to provide improved transfer, delivery, tolerance, and the like. A multitude of appropriate formulations can be found in the formulary known to all pharmaceutical chemists: Remington's Pharmaceutical Sciences (15th ed, Mack Publishing Company, Easton, Pa. (1975)), particularly Chapter 87 by Blaug, Seymour, therein. These formulations include, for example, powders, pastes, ointments, jellies, waxes, oils, lipids, lipid (cationic or anionic) containing vesicles (such as Lipofectin™), DNA conjugates, anhydrous absorption pastes, oil-in-water and water-in-oil emulsions, emulsions carbowax (polyethylene glycols of various molecular weights), semi-solid gels, and semi-solid mixtures containing carbowax. Any of the foregoing mixtures may be appropriate in treatments and therapies in accordance with the present invention, provided that the active ingredient in the formulation is not inactivated by the formulation and the formulation is physiologically compatible and tolerable with the route of administration. *See also* Baldrick P. "Pharmaceutical excipient development: the need for preclinical guidance." *Regul. Toxicol Pharmacol.* 32(2):210-8 (2000), Wang W. "Lyophilization and development of solid protein pharmaceuticals." *Int. J. Pharm.* 203(1-2):1-60 (2000), Charman W N "Lipids, lipophilic drugs, and oral drug delivery-some emerging concepts." *J Pharm Sci.* 89(8):967-78 (2000), Powell *et al.* "Compendium of excipients for parenteral formulations" *PDA J Pharm Sci Technol.* 52:238-311 (1998) and the citations therein for additional information related to formulations, excipients and carriers well known to pharmaceutical chemists.

[0168] Activatable anti-CTLA-4 antibodies can be administered in the form of pharmaceutical compositions. Principles and considerations involved in preparing such compositions, as well as guidance in the choice of components are provided, for example, in Remington: The Science And Practice Of Pharmacy 19th ed. (Alfonso R. Gennaro, *et al.*, editors) Mack Pub. Co., Easton, Pa.: 1995; Drug Absorption Enhancement: Concepts, Possibilities, Limitations, And Trends, Harwood Academic Publishers, Langhorne, Pa.,

1994; and Peptide And Protein Drug Delivery (Advances In Parenteral Sciences, Vol. 4), 1991, M. Dekker, New York.

- [0169] The formulation can also contain more than one active compound as necessary for the particular indication being treated, preferably those with complementary activities that do not adversely affect each other. Alternatively, or in addition, the composition can comprise an agent that enhances its function, such as, for example, a cytotoxic agent, cytokine, chemotherapeutic agent, or growth-inhibitory agent. Such molecules are suitably present in combination in amounts that are effective for the purpose intended.
- [0170] The active ingredients can also be entrapped in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxymethylcellulose or gelatin-microcapsules and poly-(methylmethacrylate) microcapsules, respectively, in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles, and nanocapsules) or in macroemulsions.
- [0171] The formulations to be used for *in vivo* administration must be sterile. This is readily accomplished by filtration through sterile filtration membranes.
- [0172] Sustained-release preparations can be prepared. Suitable examples of sustained-release preparations include semipermeable matrices of solid hydrophobic polymers containing the antibody, which matrices are in the form of shaped articles, e.g., films, or microcapsules. Examples of sustained-release matrices include polyesters, hydrogels (for example, poly(2-hydroxyethyl-methacrylate), or poly(vinylalcohol)), polylactides (U.S. Pat. No. 3,773,919), copolymers of L-glutamic acid and γ ethyl-L-glutamate, non-degradable ethylene-vinyl acetate, degradable lactic acid-glycolic acid copolymers such as the LUPRON DEPOT™ (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and poly-D-(-)-3-hydroxybutyric acid. While polymers such as ethylene-vinyl acetate and lactic acid-glycolic acid enable release of molecules for over 100 days, certain hydrogels release proteins for shorter time periods.
- [0173] In some embodiments, the activatable antibody contains a detectable label. An intact antibody, or a fragment thereof (e.g., Fab, scFv, or F(ab)2) can be used. The term "labeled", with regard to the probe or antibody, is intended to encompass direct labeling of the probe or antibody by coupling (i.e., physically linking) a detectable substance to the probe or antibody, as well as indirect labeling of the probe or antibody by reactivity

with another reagent that is directly labeled. Examples of indirect labeling include detection of a primary antibody using a fluorescently-labeled secondary antibody and end-labeling of a DNA probe with biotin such that it can be detected with fluorescently-labeled streptavidin. The term "biological sample" is intended to include tissues, cells and biological fluids isolated from a subject, as well as tissues, cells and fluids present within a subject. Included within the usage of the term "biological sample", therefore, is blood and a fraction or component of blood including blood serum, blood plasma, or lymph. For example, the antibody can be labeled with a radioactive marker whose presence and location in a subject can be detected by standard imaging techniques.

III. *Pharmaceutical Compositions*

[0174] The activatable anti-CTLA-4 antibodies of the invention (also referred to herein as "active compounds"), and derivatives, fragments, analogs and homologs thereof, can be incorporated into pharmaceutical compositions suitable for administration. Such compositions typically comprise the activatable antibody and a pharmaceutically acceptable carrier. As used herein, the term "pharmaceutically acceptable carrier" is intended to include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like, compatible with pharmaceutical administration. Suitable carriers are described in the most recent edition of Remington's Pharmaceutical Sciences, a standard reference text in the field, which is incorporated herein by reference. Preferred examples of such carriers or diluents include, but are not limited to, water, saline, ringer's solutions, dextrose solution, and 5% human serum albumin. Liposomes and non-aqueous vehicles such as fixed oils may also be used. The use of such media and agents for pharmaceutically active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active compound, use thereof in the compositions is contemplated. Supplementary active compounds can also be incorporated into the compositions.

[0175] A pharmaceutical composition of the invention is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, e.g., intravenous, intradermal, subcutaneous, oral (e.g., inhalation), transdermal (i.e., topical), transmucosal, and rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the

following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid (EDTA); buffers such as acetates, citrates or phosphates, and agents for the adjustment of tonicity such as sodium chloride or dextrose. The pH can be adjusted with acids or bases, such as hydrochloric acid or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

[0176] Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL™ (BASF, Parsippany, N.J.) or phosphate buffered saline (PBS). In all cases, the composition must be sterile and should be fluid to the extent that easy syringeability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars, polyalcohols such as manitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent which delays absorption, for example, aluminum monostearate and gelatin.

[0177] Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a

basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, methods of preparation are vacuum drying and freeze-drying that yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

[0178] For administration by inhalation, the compounds are delivered in the form of an aerosol spray from pressured container or dispenser which contains a suitable propellant, e.g., a gas such as carbon dioxide, or a nebulizer.

[0179] Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in the art.

[0180] Activatable antibodies of the present invention may also be administered subcutaneously in conjunction with agents to facilitate injection of large volumes at a single site (interstitial drug dispersion agents) such as soluble neutral-active hyaluronidase glycoproteins (sHASEGP), for example, human soluble PH-20 hyaluronidase glycoproteins, such as rHuPH20 (HYLENEX®, Baxter International, Inc.). Certain exemplary sHASEGPs and methods of use, including rHuPH20, are described in US Patent Publication Nos. 2005/0260186 and 2006/0104968. In one aspect, a sHASEGP is combined with one or more additional glycosaminoglycanases such as chondroitinases.

[0181] The compounds can also be prepared in the form of suppositories (e.g., with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

[0182] In some embodiments, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid.

Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Pat. No. 4,522,811.

[0183] It is especially advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the invention are dictated by and directly dependent on the unique characteristics of the active compound and the particular therapeutic effect to be achieved, and the limitations inherent in the art of compounding such an active compound for the treatment of individuals.

[0184] The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

[0185] Embodiments of the present disclosure can be further defined by reference to the following non-limiting examples, which describe in detail preparation of certain antibodies of the present disclosure and methods for using antibodies of the present disclosure. It will be apparent to those skilled in the art that many modifications, both to materials and methods, may be practiced without departing from the scope of the present disclosure.

[0186] The invention will be further described in the following examples, which do not limit the scope of the invention described in the claims.

EXAMPLES

Example 1: Identification of Masking Moieties for the Activatable anti-CTLA-4 Antibody

[0187] In order to identify masking moieties (MM) that reduce the binding of anti-CTLA-4 antibodies to their target protein, anti-CTLA-4 antibody (i.e., ipilimumab) was used to

screen peptide libraries using methods similar to that described in PCT International Publication Nos. WO 2009/025846, WO 2010/081173, and WO 2016/149201, the contents of which are hereby incorporated by reference in their entireties. The screening consisted of two rounds of magnetic-activated cell sorting (MACS) purification followed by three rounds of fluorescence-activated cell sorting (FACS).

[0188] The initial MACS purification was done with protein-A Dynabeads® (Invitrogen) and anti-CTLA-4 antibody at a concentration of 100 nM. Approximately 10^{11} cells were screened for binding, and 6×10^6 cells were collected. The second MACS purification was done with streptavidin DYNABEADS® (Thermo Fisher Scientific) and biotinylated anti-CTLA-4 antibody at a concentration of 100 nM. The eluate from the initial MACS purification was expanded, approximately 10^{11} cells were screened for binding, and approximately 10^7 cells were collected. The output of the previously described MACS purification was subjected to serial rounds of FACS sorting with decreasing concentrations of anti-CTLA-4 labeled with Alexa Fluor® 488 (Thermo Fisher Scientific). Labeled anti-CTLA4 antibody was used at concentrations of 10 nM, 1 nM, and 200 pM for the first, second, and third sorts, respectively. Individual peptide clones, from the third sort were identified by sequence analysis and subsequently verified for their ability to bind the anti-CTLA4 antibody. Two peptide consensus sequences were selected for affinity maturation: XXCXXXMYGYNLCPY (SEQ ID NO: 554) and XXXCXHSMYNVCLDP (SEQ ID NO: 555).

[0189] Affinity maturation libraries were built on these consensus sequences as described in Table 7. Rows 1 and 3 represent the consensus sequence and rows 2 and 4 represent the nucleotide sequences encoding the peptide libraries that were inserted into the display system using a method similar to that described in PCT International Publication Number WO 2010/081173, *ibid*.

Table 7: Maturation Libraries

1	X	X	C	X	X	X	M	Y	G	Y	N	L	C	P	Y
2	NNK	NNK	TGC	NNK	NNK	NNK	NTT	TWT	GGG	KWT	AAT	CTG	TGC	CCG	TAT
3	X	X	X	C	X	H	S	M	Y	N	V	C	L	D	P
4	NNK	NNK	NNK	TGC	NNK	NWT	AGT	NTT	TWT	AAT	NTT	TGC	CTT	GAT	CCT

[0190] The maturation libraries were screened in a manner similar to that described for the naïve libraries described above. The screening consisted of one round of MACS and

subsequent rounds of FACS sorting. The MACS was done with protein-A DYNABEADS® (Thermo Fisher Scientific) and the anti-CTLA-4 antibody at a concentration of 100 nM. For MACS, 10^{11} cells were screened for binding, and approximately 10^8 cells were selected. The eluate from the MACS was expanded, and approximately 10^{11} cells were subjected to serial rounds of FACS sorting with decreasing concentrations of Alexa Fluor® 488-labeled anti-CTLA4 antibody. Labeled anti-CTLA4 antibody was used at concentrations of 100 nM, 20 nM, 5 nM, 1 nM, and 1 nM for the first, second, third, fourth and fifth sorts, respectively. Individual peptide clones from the fourth and fifth sorts were identified by sequence analysis and subsequently verified for their ability to bind the anti-CTLA4 antibody. The sequences of the anti-CTLA-4 masking moieties identified through the methods described above are provided in Tables 4 and 5. Four consensus sequences can be derived from the mask sequences listed in Tables 4 and 5:

Consensus 1. C(L/M/V/T)Y(S/V/I)(F/L/M/A)(Y/F)N(V/I)CLDP (SEQ ID NO: 566)

Consensus 2. CAQMYGYSMC(P/A)(H/R/A)T (SEQ ID NO: 567)

Consensus 3. CX(M/I/Y/L/N/F)(Y/W/F/Q/T)(M/Y)YG(Y/V/F)(N/D)LCP(Y/F) (SEQ ID NO: 568)

Consensus 4. (N/T)(S/T/M/A)CP(N/Y)HP(M/L)C(H/F/Y)D(Y/F/W) (SEQ ID NO: 569)

Example 2: Construction and Characterization of Activatable Anti-muCTLA-4 Antibodies

[0191] In order to show a proof-of-concept that the activatable anti-CTLA-4 antibodies can be used to treat tumors, six activatable anti-mouse CTLA-4 antibodies (based on clone 9D9) were constructed using techniques similar to those disclosed in Examples 1 and 3 herein. These antibodies comprise either MY11 or MY03 as the masking moiety, and cleavable moiety “0003” having amino acid sequence TSTSGRSANPRG (SEQ ID NO: 320), “1004” having amino acid sequence AVGLLAPP (SEQ ID NO: 323), or “2001” having amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 297). The antibodies were all mouse IgG2a isotype. As controls, anti-mouse CTLA-4 monoclonal antibody (9D9) (“9D9 mg2a”) and a human anti-diphtheria toxin antibody with a mIgG2a isotype (“mg2a”) were used.

- [0192] On day 0, BALB/c mice were subcutaneously injected with 1×10^6 CT26 tumor cells. Administration of the different antibodies began on day 7 post tumor implantation. Prior to administration, tumor size was measured and the mice were randomized into different treatment groups, so as to have comparable mean tumor volumes (e.g., 39-44 mm³). Tumors were measured with calipers two-dimensionally, and tumor volume was calculated as $L \times (W^2/2)$, L = length (the longer of the 2 measurements), W = width. The mice were then treated intraperitoneally (i.p.) with the designated antibody (e.g., 25 µg/dose). Tumor volume was measured twice weekly. At day 12 post tumor implantation, some of the mice from each group were sacrificed, and tumor and spleen were harvested for immunomonitoring to investigate the effects of the antibodies on the T cell populations. Some or all of the remaining mice from the different groups were used for subsequent pharmacokinetic (PK) and/or pharmacodynamics (PD) analysis.
- [0193] As shown in FIG. 1A, mice that received the unrelated mouse IgG2a antibody (i.e., the human anti-diphtheria toxin antibody) failed to control the tumor. In contrast, as shown in FIG. 1C, mice that received the activatable anti-mouse CTLA-4 antibody (comprising MY11 as the masking moiety and 2001 as the cleavable moiety) controlled tumor size almost as well as those mice that received the anti-mouse CTLA-4 mAb (9D9) (FIG. 1B). These data demonstrate that tumor-specific protease can cleave the cleavable moiety, resulting in the removal of the masking moiety and the binding of the released antibody to its target protein.
- [0194] To determine whether or not activatable anti-mouse CTLA-4 antibodies are active in the periphery, proliferation and activity of Foxp3⁺ regulatory T cells were determined in the spleen, and regulatory T cell abundance was determined in tumor samples for comparison, as described in Example 5, *infra*. In agreement with the data from FIGs. 1B and 1C, all the activatable anti-CTLA-4 antibodies behaved similarly to the anti-mouse CTLA-4 mAb (9D9) in the tumor (FIG. 2A). In contrast, the activatable antibodies resembled the unrelated mouse IgG2a antibody in the spleen (FIGs. 2B and 2C). Such data suggest that the masking moiety-containing prodomain of the activatable anti-mouse CTLA-4 antibodies remains intact and attached to the antibody in the spleen, blocking the activity of the antibody, whereas the prodomain is cleaved off by tumor specific proteases to generate fully active anti-CTLA-4 antibody in the tumor.

Example 3:
Construction of Activatable Anti-human CTLA-4 Antibodies

[0195] Activatable anti-CTLA4 antibodies comprising an anti-CTLA4 masking moiety, a cleavable moiety, and an anti-CTLA4 antibody (e.g., ipilimumab) of the disclosure were produced according to methods similar to those described in PCT Publication Nos. WO 2009/025846 *ibid.* and WO 2010/081173 *ibid.*, and WO 2016/118629, *ibid.* Activatable anti-CTLA4 antibodies were expressed in EXPI293™ cells (Thermo Fisher Scientific) and purified by protein A chromatography (MabSelect SuRe, GE Healthcare) as per manufacturers' protocols. Quality control of the resultant activatable antibodies indicated that most comprise at least 95% monomer.

[0196] To assess the feasibility of using the activatable anti-CTLA-4 antibodies disclosed herein in a human setting, the antibodies were produced as human IgG1 (hIgG1) heavy chain (Hc) and human kappa (hK) light chain (Lc) format. The activatable antibodies all comprise the antibody or antigen binding domain thereof of ipilimumab. The cleavable moiety was selected from the group consisting of a cleavable moiety referred to herein as "2001" and comprising the sequence ISSGLLSGRSDNH (SEQ ID NO: 297) and derivatives thereof and a cleavable moiety referred to herein as "3001" and comprising the sequence AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306) and derivatives thereof. In some embodiments, the cleavable moiety was selected from the group consisting of ISSGLLSGRSDNH (SEQ ID NO: 297), also referred to herein as "2001"; ISSGLLSGRSDDH (SEQ ID NO: 300), also referred to herein as "2006"; ISSGLLSGRSDIH (SEQ ID NO: 301), also referred to herein as "2007"; ISSGLLSGRSDQH (SEQ ID NO: 302), also referred to herein as "2008"; ISSGLLSGRSDTH (SEQ ID NO: 303), also referred to herein as "2009"; ISSGLLSGRSANP (SEQ ID NO: 305), also referred to herein as "2012"; ISSGLLSGRSDNP (SEQ ID NO: 304), also referred to herein as "2011"; ISSGLLSGRSANPRG (SEQ ID NO: 298), also referred to herein as "2003"; AVGLLAPPGGLSGRSDNH (SEQ ID NO: 306), also referred to herein as "3001"; AVGLLAPPGGLSGRSDDH (SEQ ID NO: 307), also referred to herein as "3006"; AVGLLAPPGGLSGRSDIH (SEQ ID NO: 308), also referred to herein as "3007"; AVGLLAPPGGLSGRSDQH (SEQ ID NO: 309), also referred to herein as "3008"; AVGLLAPPGGLSGRSDTH (SEQ ID NO: 310), also referred to herein as "3009";

AVGLLAPPGGLSGRSANP (SEQ ID NO: 312), also referred to herein as "3012"; AVGLLAPPGGLSGRSDNP (SEQ ID NO: 311), also referred to herein as "3011"; and AVGLLAPPSGRSANPRG (SEQ ID NO: 299), also referred to herein as "2005". The masking moiety was selected from the group of masking moieties provided in Tables 4 and 5. In some embodiments, the masking moiety was CRTQLYGYNLCPY (SEQ ID NO: 39), referred to herein as YV39. Some of the activatable anti-CTLA-4 antibodies also included spacer sequences and/or linker peptides.

Example 4:

In vitro Characterization of Activatable Anti-Human CTLA-4 Antibodies

[0197] In order to assess the ability of the activatable antibodies to bind to CTLA-4 in the absence of protease activity, an enzyme-linked immunosorbent assay (ELISA) was used to measure binding affinity. Briefly, Nunc MaxiSorp[®] plates were coated overnight at 40°C with 100 μ L/well of a 1 μ g/mL solution of human CTLA-4 protein (Sino Biological) in PBS, pH 7.4. Plates were then washed three times with PBST (PBS, pH 7.4, 0.05% Tween-20), and the wells were blocked with 200 μ L/well, 10 mg/mL bovine serum albumin (BSA) in PBST for 2 hours at room temperature. Afterwards, the plates were washed three more times with PBST. The activatable antibodies were then serially diluted, as shown below in Table 8.

Table 8. Serial Dilution of Activatable Anti-CTLA-4 Antibodies for Binding Analysis

	[Antibody] = nM Columns 1-3	[activatable antibody 1] = nM Columns 4-6	[activatable antibody 2] = nM Columns 7-9	[activatable antibody 3] = nM Columns 10-12
A	10	1000	1000	1000
B	3.33	333	333	333
C	1.11	111	111	111
D	0.37	37	37	37
E	0.123	12.3	12.3	12.3
F	0.041	4.1	4.1	4.1
G	0.0137	1.34	1.34	1.34
H	.0046	0.45	0.45	Blank

[0198] In the current Example, the highest concentration used for the parental antibody and the activatable antibodies were 10 nM and 100 nM, respectively. However, the

concentrations can be increased or decreased to give full saturation binding curves for activatable antibodies with stronger or weaker masking.

[0199] The diluted antibodies were added to the plates and incubated for 1 hour at room temperature. Afterwards, the plates were washed three times with PBST. Then, 100 μ L of goat-anti-human IgG (Fab specific, Sigma cat # A0293; diluted at 1:4,000 in 10 mg/mL BSA in PBST) was added to each well, and the plate was incubated for an additional 1 hour at room temperature. Next, the plates were developed with tetramethylbenzidine (TMB) and 1N HCl. Absorbance at 450 nm was then measured and reported as optical density (OD 450 nm).

[0200] As shown in FIGs. 3A to 3E, anti-CTLA-4 activatable antibodies typically had reduced binding to CTLA-4 as compared to ipilimumab ("YV1"). *See also* FIGs. 4A to 4D, FIGs. 5A to 5F, and FIGs. 6A to 6B. Such data demonstrate that the masking moieties effectively conceal the antigen binding domain on the anti-CTLA-4 activatable antibodies.

[0201] To further assess the binding ability, the activatable human anti-CTLA-4 antibodies were serially diluted (e.g., 60 μ g/mL to 0.0003 μ g/mL) and added to 58 α - β -CTLA-4/CD3 ζ cells, which stably express human CTLA-4. After 30 minutes of incubation at 4°C, an allophycocyanin (APC)-labeled anti-human secondary antibody was added and binding of the activatable anti-human CTLA-4 antibodies to human CTLA-4 was assessed using a Canto flow cytometer. The geometric mean fluorescence intensity (GMFI) was determined using FlowJo[®] analysis software. Ipilimumab was used as a control. As shown in FIGs. 7A and 7B, the activatable human anti-CTLA-4 antibodies did not bind to human CTLA-4 as effectively as ipilimumab. These data further demonstrate that in the absence of specific proteases, the masking moiety of the activatable antibodies inhibits binding of such activatable antibodies to human CTLA-4.

[0202] To confirm that the reduced binding observed with the activatable anti-CTLA-4 antibodies was due to the masking moiety, studies were performed on mono-clipped, MMP fully-clipped and uPA fully-clipped forms of the activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety. The mono-clipped form of the antibody was produced by expressing a construct producing one intact light chain (including the mask moiety) and a second light chain truncated at the same position as if it had been cleaved by MMP14. The MMP or uPA fully clipped forms were expressed

from constructs with both light chains truncated as if they had been cleaved by MMP or μ PA, respectively. As shown in FIGs. 7C and 7D, the mono-clipped activatable antibody had intermediate binding ($EC_{50} = 2.8$ nM) as compared to the non-clipped activatable antibody ($EC_{50} = 22$ nM) and ipilimumab ($EC_{50} = 0.54$ nM). In contrast, the MMP or μ PA fully-clipped activatable antibodies behaved similarly to ipilimumab (MMP clipped: $EC_{50} = 0.65$ nM; μ PA clipped: $EC_{50} = 0.76$). Such data confirm that the reduced binding observed with the activatable anti-CTLA-4 antibody is due to the masking moiety.

[0203] Next, to determine whether the observed reduced binding to CTLA-4 correlated with reduced activity, the activity of an activatable human anti-CTLA-4 antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011") was characterized in an *in vitro* functional assay using staphylococcal enterotoxin B (SEB). SEB is a superantigen that strongly activates T cells and stimulates cytokine secretion. Whole fresh peripheral blood mononuclear cells (PBMC) were isolated from healthy human donors using a standard Ficoll-Paque separation method. Serial dilution of the antibodies (e.g., 40 μ g/mL to 0.01 μ g/mL) were performed and plated in triplicate in a 96-well flat-bottom tissue culture plate. The antibodies used included (i) Ipi YV39 2011, (ii) ipilimumab, and (iii) an unrelated isotype control. Next, the isolated PBMC were resuspended in T-cell assay media (RPMI media + 10% heat-inactivated fetal bovine serum (HI-FBS) + 1% HEPES buffer + 1% MEM non-essential amino acid + 1% Na-pyruvate) and added to the plate at 1×10^5 cells/well. The cells were stimulated with a suboptimal concentration (e.g., 85 ng/mL – determined by titrating SEB and observing the stimulation on T-cell proliferation) of SEB. The cells were incubated at 37°C for 3 days. Then, the IL-2 concentration in the supernatants was measured by homogeneous time-resolved fluorescence (HTRF). The HTRF data were analyzed using Softmax Pro and graphed using GraphPad Prism.

[0204] As shown in FIG. 8, ipilimumab enhanced the SEB-mediated IL-2 production by the PBMC in a dose-dependent manner. In contrast, the Ipi YV39 2011 activatable antibody had activity similar to that of the isotype control, suggesting that the masking moiety (YV39) is effective in blocking the functional activity of ipilimumab. These data are in agreement with the binding data described above and demonstrate that in the absence of specific proteases, the activatable anti-human CTLA-4 antibodies exhibit reduced activity.

Example 5:

In vivo Characterization of Activatable Anti-Human CTLA-4 Antibodies

[0205] In order to characterize the antibodies disclosed herein *in vivo*, four activatable human anti-human CTLA-4 antibodies (based on ipilimumab) were prepared using mouse IgG2a. The antibodies comprise YV04, YV23, YV24, or YV39 as the masking moiety, and 2001 as the cleavable moiety ("Ipi YV04 2001", "Ipi YV23 2001", "Ipi YV24 2001", and "Ipi YV39 2001", respectively). As controls, ipilimumab ("Ipi mg2a") and an unrelated human anti-diphtheria toxin ("control mg2a") were used. The activity of these activatable anti-CTLA-4 antibodies was assessed using the MC38 tumor model as described below.

[0206] Briefly, on day 0, human CTLA-4 knock-in C57BL/6 mice were subcutaneously injected with 2×10^6 MC38 colon adenocarcinoma cells into their left lower abdominal quadrant. Tumors were measured with calipers two-dimensionally, and tumor volume was calculated as $L \times (W^2/2)$, L = length (the longer of the 2 measurements), W = width. Next, the mice were randomized into different groups, so as to have similar mean tumor volumes (e.g., 37 mm³). Administration of the antibodies began on day 7 post tumor implantation with the mice receiving a single dose (e.g., 200 µg/mouse) of the relevant antibody via intraperitoneal (i.p.) injection. At day 12 post tumor implantation, several of the mice from each group were sacrificed, and tumor and spleen were harvested for immunomonitoring to investigate the effect of the antibodies on the T cell populations. Some or all of the remaining mice from the different groups were used for subsequent pharmacokinetic (PK) and/or pharmacodynamics (PD) analysis.

Immunomonitoring of T Cell Populations

[0207] The harvested tumor and spleen were processed on a gentleMACS Octo Dissociator™ (Miltenyi, San Diego, CA). Single cell suspensions were stained with the following T cell markers: CD4, CD8, CD19, ICOS, CD45, FoxP3, CTLA-4, CD3, Ki-67, PD-1, Granzyme B, and LIVE/DEAD®.

PK/PD Analysis

[0208] The mice were checked daily for postural, grooming, and respiratory changes, as well as lethargy. Tumors and group body weights were recorded twice a week until death,

euthanasia, or end of the study period. The response to the treatments was measured as a function of tumor growth inhibition (TGI), which was calculated as follows: $\% \text{ TGI} = \{1 - [(T_t - T_o)/(C_t - C_o)]\} \times 100$, T_t = tumor volume of the treatment group on a given day, T_o = initial tumor volume, C_t = tumor volume of the control group on a given day, C_o = initial tumor volume of the control group. Animals were euthanized if the tumor reached a volume greater than approximately 2500 mm³ or appeared ulcerated.

Statistical Analysis

- [0209] Microsoft Excel was used to calculate the mean, standard deviation (SD), and median values of tumor volumes and body weights. The mean and median values were calculated when 100% and at least 60% of the study animals remained in each treatment group, respectively. GraphPad Prism[®] v.4 software was used to plot data.
- [0210] As expected, mice that received the unrelated control antibody failed to control tumor growth (FIG. 9A) whereas all the mice that received ipilimumab effectively controlled tumor growth (FIG. 9B). Mice that received the different activatable human anti-CTLA-4 antibodies controlled tumor growth comparably with ipilimumab (FIGs. 9C to 9F). Of the activatable antibodies, Ipi YV39 2001 most closely resembled the efficacy of ipilimumab in controlling tumor growth (FIG. 9F).
- [0211] In regard to the frequency of regulatory T cells in the tumor and spleen of the treated mice, as observed earlier with the activatable anti-mouse CTLA-4 antibodies (*see* Example 2), activatable anti-human CTLA-4 antibodies (mouse IgG2a isotype) behaved similarly to ipilimumab in tumors (FIGs. 12A and 12B), but in the spleen, the activatable antibodies were more comparable to the unrelated control antibody (FIGs. 12C to 12F).
- [0212] The data shown here collectively demonstrate that the activatable human anti-CTLA-4 antibodies disclosed herein can effectively control tumors like the traditional ipilimumab while exhibiting less risk of undesirable side effects.

Example 6:

In vivo Characterization of Activatable Anti-Human CTLA-4 Antibodies Comprising Modified Cleavable Moieties

- [0213] To address a possible deamidation site in certain cleavable moiety sequences (*see* Example 10), activatable human anti-CTLA4 antibodies were prepared using a human

IgG1 and various CM sequences. The activatable antibodies comprise YV39 as the masking moiety and one of several variants of the 2001 cleavable moiety: WT (2001), ANP (2012), DNP (2011), or Q (2008) ("Ipi YV39 2001", "Ipi YV39 2012", "Ipi YV39 2011", and "Ipi YV39 2008", respectively). Ipilimumab and the unrelated human anti-diphtheria toxin were again used as controls.

[0214] To measure the activity of the activatable anti-CTLA-4 antibodies, the MC38 tumor model was used as described above in Example 5. For the dose titration study (FIGs. 11A to 11F), the mice were treated with ipilimumab or the activatable antibody comprising YV39 as the masking moiety and 2011 as the cleavable moiety ("Ipi YV39 2011") at doses of 200 µg/dose, 60 µg/dose, and 20 µg/dose.

[0215] As shown in FIGs. 10A and 10B, mice treated with the control antibody failed to control the tumor, whereas 6 out of 10 mice treated with ipilimumab were tumor-free at the end of the experiment. Mice treated with the different activatable antibodies were able to control tumor as observed with the traditional ipilimumab (FIGs. 10C to 10F). *See also* FIGs. 11B – 11G.

[0216] In regard to the frequency of regulatory T cells in the tumor and spleen of the treated mice, as observed earlier, tumor-specific protease was required to cleave the 2001 cleavable moiety variants. In the tumors, these activatable antibodies behaved like ipilimumab in reducing the frequency of Foxp3+ regulatory T cells (FIGs. 13A, 13B, 14A, and 14B). *See also* FIG. 15. In the spleen, the antibodies more closely mirrored the unrelated control antibody (FIGs. 13C to 13E, 14D to 14G, and 16A to 16B), demonstrating that the masking moiety remains coupled to the activatable antibody in the absence of the specific tumor-associated proteases.

Example 7:

In vivo Characterization of A Non-Fucosylated Version of Activatable Anti-Human CTLA-4 Antibodies

[0217] As described above, the absence of core fucose residues can strongly enhance ADCC via improved binding of IgG to activating FcγRIIIA without altering antigen binding or CDC. Natsume *et al.* (2009) *Drug Des. Devel. Ther.* 3:7. Non-fucosylated forms of ipilimumab ("Ipi NF") and ipi YV39 2011 ("Ipi YV39 2011 NF") were prepared. Binding of Ipi and Ipi NF were determined for various mouse, human and

cynomolgus monkey Fc receptors. Results are provided at FIG. 19. As expected, Ipi NF showed dramatically enhanced affinity (i.e., lower K_d) for activating receptors human CD16a (FcγRIIIa), cyno CD16 (FcγRIII) and mouse FcγRIV.

[0218] Ipi YV39 2011 NF and Ipi-NF were tested at various doses in the MC38 tumor model described in Example 5. Ipilimumab and an unrelated hIgG1 were used as controls. Results are provided at FIGs. 17A - D. Ipi NF was somewhat more effective at limiting or preventing tumor growth than ipilimumab (compare FIGs. 17B and 17C), and Ipi YV39 2011 NF was equivalent to Ipi NF (compare FIGs. 17C and 17D). In addition, FoxP3+ regulatory T cells were also similarly depleted in the tumors of mice treated with Ipi NF and Ipi YV39 2011 antibody (*see* FIG. 18). In both experiments, the Ipi YV39 2011 NF is shown to be fully activated in the tumor.

[0219] These results confirm that the methods of the present invention are equally applicable to non-fucosylated forms of ipilimumab, including non-fucosylated activatable CTLA-4 antibodies such as YV39 2011 NF.

Example 8: *In vivo* Characterization of Activatable Anti-Human CTLA-4 Antibodies in Cynomolgus Monkeys

[0220] To assess the anti-CTLA-4 antibodies in a primate, cynomolgous monkeys were administered activatable antibody comprising YV39 as the masking moiety and 2001 as the cleavable moiety. Vehicle and ipilimumab were used as controls. Each monkey received 10 mg of antibody or anti-CTLA-4 activatable antibody, and blood was collected on days 0, 4, 8, 15, 22, 36, and 43 post-antibody administration. As shown in FIG. 20, in monkeys that received ipilimumab, there was a spike in CD4+ T cell proliferation as measured by Ki67-staining at around days 8-15 post antibody administration. In contrast, activatable anti-CTLA-4 antibody behaved similarly to the vehicle control and did not induce CD4+ T cell proliferation in the monkeys. These data demonstrate that even in primates, the activatable anti-CTLA-4 antibody shows little if any activation, indicating the absence of specific proteases.

[0221] Collectively, the data presented at FIGs. 1 - 20 demonstrate that the activatable anti-CTLA-4 antibodies described herein offer an improvement over ipilimumab. The

activatable antibodies control tumor growth just as effectively as ipilimumab while reducing the risk of serious adverse events often observed with ipilimumab treatment.

Example 9:
K_{app} and ME Values for Activatable CTLA-4 Antibodies

[0222] Table 9 provides the K_{app} and masking efficiency (ME) values for activatable antibodies, disclosed herein, comprising a variety of masking moieties and cleavable moieties in a human IgG1 format. The values provided in this Table were calculated from the data depicted in the Figures. K_{app} represents the binding affinity of the activatable antibody under the conditions of the measurement, in this example binding by ELISA; it is to be appreciated, however, that binding affinity can also be measured by binding to CTLA-4 expressed on primary or transfected cells or by other physical methods such as, but not limited to, surface plasmon resonance or equilibrium dialysis. Masking efficiency (ME) is calculated by dividing the K_{app} of the activatable antibody by the K_D of ipilimumab, measured under the same conditions.

Table 9: K_{app} and ME Values

	CM 2001		CM 3001		CM 2008		CM 2011		CM 2012		NSUB	
	K _{app} nM	ME	K _{app} nM	ME	K _{app} nM	ME	K _{app} nM	ME	K _{app} nM	ME	K _{app} nM	ME
YV04-YV1	17.8	57										
YV06-YV1	0.6	2										
YV09-YV1	33.6	112	44.4	126								
YV23-YV1	11.4	38	13.8	39								
YV24-YV1			9.0	29								
YV27-YV1	0.7	2.3	0.8	2.3								
YV29-YV1	0.7	2.3	0.8	2.3								
YV32-YV1	0.9	3.0	1.2	3.4								
YV33-YV1	1.3	4.3	1.9	5								
YV35-YV1	3.7	12.3	5.3	15								
YV39-YV1	16.9	56	14.3	41	31.4	135	13.2	57	14.9	64	31.8	137
YV41-YV1	14.4	48	22.6	65								
YV51-YV1	4.4	15	4.9	14								
YV52-YV1	0.8	2.7	0.9	2.6								
YV53-YV1	4.1	14	5.3	15								
YV54-YV1	0.6	2	1.0	2.8								
YV55-YV1	4.8	16	6.0	18								
YV56-YV1	0.4	1.3	0.4	1								
YV57-YV1	0.4	1.3	1.6	4.6								
YV58-YV1	0.3	1	0.4	1								

[0223] Table 10 provides the K_{app} and ME values for the activatable antibodies disclosed herein, comprising a variety of masking moieties and cleavable moieties in a YV1 mouse Ig2a format. The values provided were calculated from the data depicted in the Figures.

Table 10: K_{app} and ME values

	CM 2001		CM 2006		CM 2007		CM 2008		CM 2009	
	K_{app} nM	ME	K_{app} nM	ME	K_{app} nM	ME	K_{app} nM	ME	K_{app} nM	ME
YV04-YV1	5.7	16.2	26.4	75	19.3	55	19.1	54	16.4	47
YV23-YV1			12.5	36	7.8	22	2.7	8	9.4	27
YV39-YV1	18.0	51	23.9	68			17.6	50	18.0	51

[0224] Table 11 provides K_{app} and ME values for the activatable antibodies comprising masking moieties having higher ME values and the 2012 cleavable moiety in aYV1 mouse IgG2a format. The values provided were calculated from the data depicted in the Figures.

Table 11: K_{app} and ME values

	CM 2001		CM 2011		CM 2012		NSUB	
	K_{app} nM	ME	K_{app} nM	ME	K_{app} nM	ME	K_{app} nM	ME
YV39-YV1	18.0	51	18.0	51	12.9	144	29.8	85
YV61-YV1					17.9	200		
YV62-YV1					15.5	173		
YV63-YV1					104	1170		
YV64-YV1					56.5	631		
YV65-YV1					12.3	156		
YV66-YV1					18.9	242		
YV01-YV1					38.6	493		
YV02-YV1					14.8	189		

Example 10:

Deamidation, Isomerization, and Stabilization Assessment for Activatable CTLA-4 Antibodies

[0225] As suggested in Example 6, to address a possible deamidation site in certain cleavable moiety (CM) sequences in certain activatable human anti-CTLA-4 antibodies, such activatable antibodies were prepared using various CM sequences (i.e., 2001, 2011, 2012, and 2008). In the cleavable moieties 2011, 2012, and 2008, the DNH sequence found in the 2001 cleavable moiety was replaced with DNP, ANP, and DQH, respectively.

[0226] These activatable CTLA-4 antibodies were produced by transient transfection of the relevant constructs in HEK 293 cells, and subjected to peptide mapping liquid chromatography - mass spectroscopy (LC-MS) to detect potential breakdown products. The 2001 (DNH) cleavable moiety, which was initially selected for use in the activatable

anti-CTLA-4 antibodies of the present invention, showed deamidation of the asparagine (N) residue (6.4%) after 7 days in PBS at 4°C. Forced stability studies showed an increase from 18.5% to 32.8% deamidation when stored at 25°C for 4 weeks, and to 36.5% and 66.6% when stored at 40°C for one week and four weeks, respectively.

[0227] Cleavable moieties 2008, 2011 and 2012 were selected to try to overcome the deamidation problem with 2001 in these activatable CTLA-4 antibodies. All of these had 0.1% or less deamidation when stored 40°C for one week in PBS, compared with 6.4% deamidation of 2001. However, further stability analysis (also by LC-MS) showed that while these activatable CTLA-4 antibodies comprising the 2008 (DQH) cleavable moiety exhibited minimal deamidation, it showed significant aspartate isomerization at the aspartate residue under various conditions (*see* Table 12). In contrast, 2011 (DNP) exhibited minimal aspartate isomerization. Aspartate isomerization was not relevant for 2012 (ANP), in which the aspartate residue is replaced with alanine.

Table 12: Isomerization values

Temperature	Time	Cleavable Moiety – Isomerization Values		
		2011 (DNP)	2012 (ANP)	2008 (DQH)
-80°C	0 days (T ₀)	0.1%	N/A	1.8%
4°C	0 days (T ₀)	0.1%	N/A	2.4%
25°C	3 months	0.2%	N/A	8.2%
40°C	3 months	0.2%	N/A	34.5%

[0228] However, *in vitro* stability studies in mouse, rat, and cynomolgus monkey serum showed substantial clipping between asparagine and proline residues for 2012 (ANP) (*see* Table 13) in these activatable CTLA-4 antibodies. 2011 (DNP) remained as the cleavable moiety with acceptably low levels of deamidation, aspartate isomerization, and light chain clipping.

Table 13: Degree of clipping observed between the asparagine and proline residues

Serum	Cleavable Moiety – Clipping Between Asparagine and Proline Residues	
	2011 (DNP)	2012 (ANP)
Mouse	-	++
Cyno	+/-	+++

[0229] All publications, patents, patent applications, internet sites, and accession numbers/database sequences (including both polynucleotide and polypeptide sequences)

cited herein are hereby incorporated by reference in their entirety for all purposes to the same extent as if each individual publication, patent, patent application, internet site, or accession number/database sequence were specifically and individually indicated to be so incorporated by reference.

What is claimed:

1. An activatable anti-human CTLA-4 antibody comprising:
 - (i) a heavy chain comprising a heavy chain variable domain (VH) comprising CDRH1: SYTMH (SEQ ID NO: 557); CDRH2: FISYDGNNKYYADSVKG (SEQ ID NO: 558); and CDRH3: TGWLGPFDY (SEQ ID NO: 559); and
 - (ii) a light chain comprising:
 - (a) a light chain variable domain (VL) comprising CDRL1: RASQSVGSSYLA (SEQ ID NO: 560); CDRL2: GAFSRAT (SEQ ID NO: 561); and CDRL3: QQYGSSPWT (SEQ ID NO: 562);
 - (b) a cleavable moiety (CM); and
 - (c) a masking moiety (MM),wherein the light chain has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-VL.
2. The activatable anti-human CTLA-4 antibody of Claim 1, wherein the MM is selected from the group consisting of YV01, YV02, YV03, YV04, YV09, YV23, YV24, YV35, YV39, YV51, YV61, YV62, YV63, YV64, YV65, and YV66.
3. The activatable anti-human CTLA-4 antibody of Claim 2, wherein the CM is a substrate for a protease selected from the group consisting of MMP1, MMP2, MMP3, MMP8, MMP9, MMP11, MMP13, MMP14, MMP17, legumain, matriptase, and uPA.
4. The activatable anti-human CTLA-4 antibody of Claim 3, wherein the CM is selected from a group consisting of 2001, 2003, 2005, 2006, 2007, 2008, 2009, 2011, 2012, 3001, 3006, 3007, 3008, 3009, 3011, and 3012.
5. The activatable anti-human CTLA-4 antibody of Claim 4, wherein the MM is YV04, YV23, YV24, YV39, YV61, YV62, YV63, or YV64.
6. The activatable anti-human CTLA-4 antibody of Claim 5, wherein the MM is YV39.
7. The activatable anti-human CTLA-4 antibody of Claim 6, wherein the CM is 2001, 2011, or 2012.

8. The activatable anti-human CTLA-4 antibody of claim 7, wherein the CM is 2011.
9. The activatable anti-human CTLA-4 antibody of any one of claims 1 to 8 comprising:
 - (i) a heavy chain comprising the amino acid sequence of SEQ ID NO: 345; and
 - (ii) a light chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 563, 564, and 565.
10. The activatable anti-human CTLA-4 antibody of claim 9 comprising:
 - (i) a heavy chain comprising the amino acid sequence of SEQ ID NO: 345; and
 - (ii) a light chain comprising the amino acid sequence of SEQ ID NO: 564.
11. The activatable anti-human CTLA-4 antibody of claim 10, wherein:
 - (i) the heavy chain further comprises the human IgG1 constant domain sequence of SEQ ID NO: 350; and
 - (ii) the light chain further comprises the human light chain kappa constant domain sequence of SEQ ID NO: 346.
12. The activatable anti-human CTLA-4 antibody of any one of claims 1 to 11 further comprising a second linker peptide (LP2) as disclosed herein, and wherein the activatable anti-human CTLA-4 antibody has the structural arrangement, from N-terminus to C-terminus, MM-LP1-CM-LP2-VL or MM-LP2-CM-LP1-VL.
13. The activatable anti-human CTLA-4 antibody of claim 12, wherein the LP1 and the LP2 are not identical to each other.
14. The activatable anti-human CTLA-4 antibody of any one of claims 1 to 13 further comprising a spacer, and having the structural arrangement, from N-terminus to C-terminus, spacer-MM-CM-VL.
15. The activatable anti-human CTLA-4 antibody of any one of claims 1 to 14 further comprising a toxic agent.

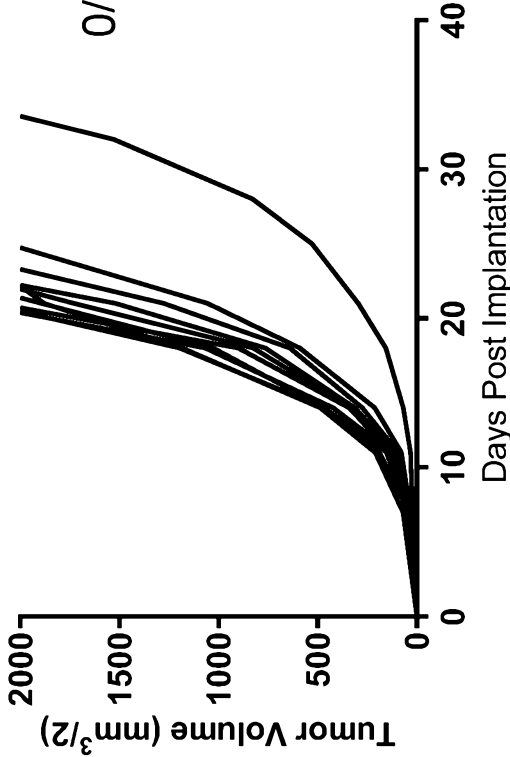
16. The activatable anti-human CTLA-4 antibody of claim 15, wherein the toxic agent is conjugated to the activatable antibody via a cleavable linker.
17. The activatable anti-human CTLA-4 antibody of any one of claims 1 to 14 further comprising a detectable moiety.
18. The activatable anti-human CTLA-4 antibody of claim 178, wherein the detectable moiety is a diagnostic agent.
19. A pharmaceutical composition comprising the activatable anti-human CTLA-4 antibody of any one of claims 1 to 18 and a carrier.
20. The pharmaceutical composition of claim 19 comprising an additional therapeutic agent.
21. An isolated nucleic acid molecule encoding the heavy chain and/or the light chain of the activatable anti-human CTLA-4 antibody of any one of claims 1 to 14.
22. A vector comprising the isolated nucleic acid molecule of claim 21.
23. A method of producing an activatable anti-human CTLA-4 antibody comprising:
 - (i) culturing a cell comprising the vector of claim 22 under conditions that lead to expression of the activatable antibody; and
 - (ii) recovering the activatable antibody.
24. A method of reducing CTLA-4 activity in a subject in need thereof comprising administering an effective amount of the pharmaceutical composition of claim 19 or 20 to the subject.
25. A method of treating, alleviating a symptom of, or delaying the progression of a cancer in a subject comprising administering a therapeutically effective amount of the pharmaceutical composition of claim 19 or 20 to the subject.
26. The method of claim 25, wherein the cancer is a bladder cancer, a bone cancer, a breast cancer, a carcinoid, a cervical cancer, a colon cancer, an endometrial cancer, a glioma, a

head and neck cancer, a liver cancer, a lung cancer, a lymphoma, a melanoma, an ovarian cancer, a pancreatic cancer, a prostate cancer, a renal cancer, a sarcoma, a skin cancer, a stomach cancer, a testis cancer, a thyroid cancer, a urogenital cancer, or a urothelial cancer.

27. The method of claim 26, wherein the cancer is melanoma.

FIG. 1A

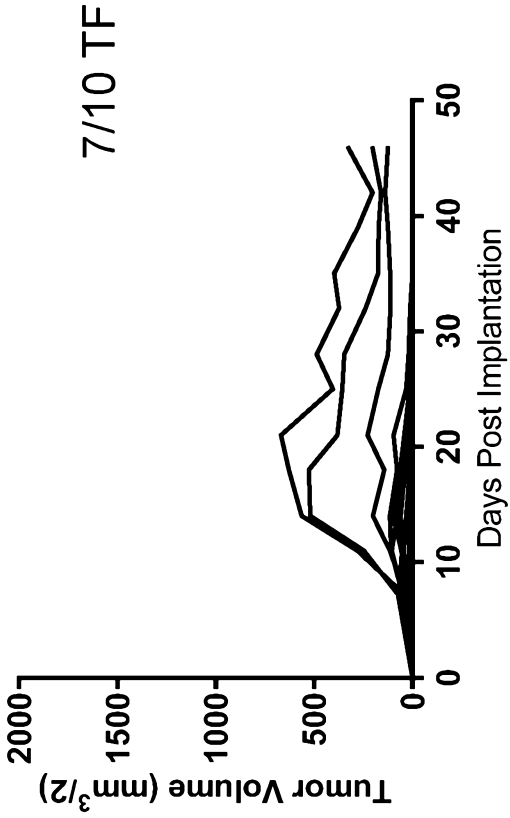
Control mlgG2a



0/10 TF

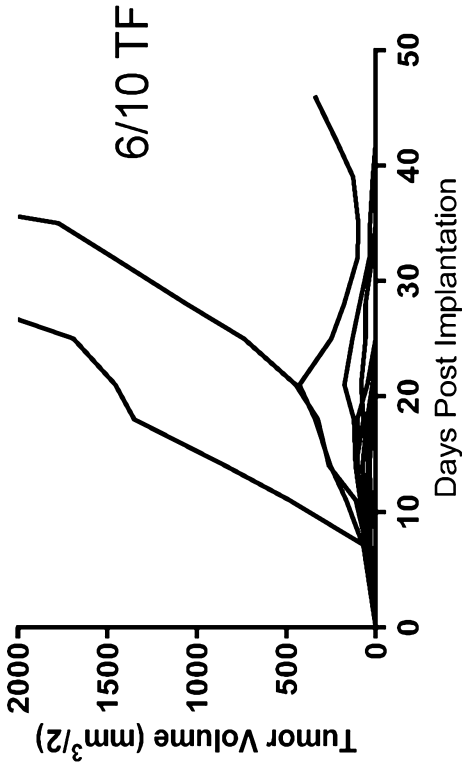
FIG. 1B

9D9 mlgG2a



7/10 TF

FIG. 1C 9D9 mlgG2MY11 2001



6/10 TF

FIG. 2A Tumor Foxp3+ of CD4+

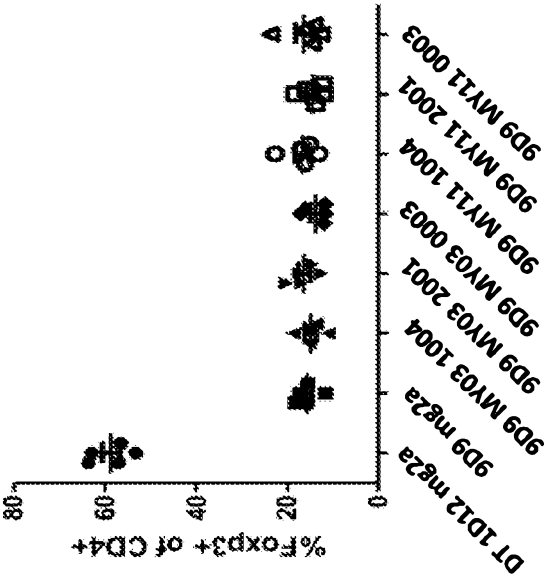


FIG. 2B Spleen Ki-67+ of Foxp3+

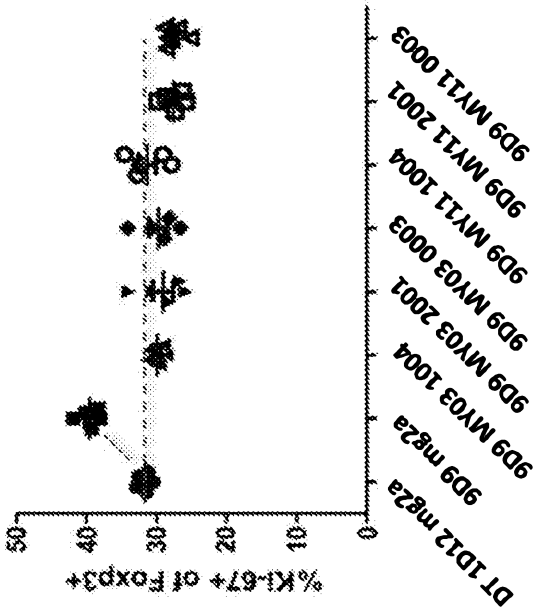
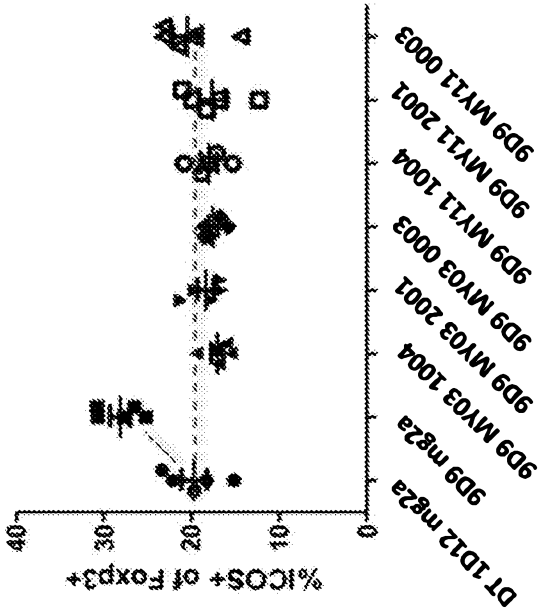


FIG. 2C Spleen ICOS+ of Foxp3+



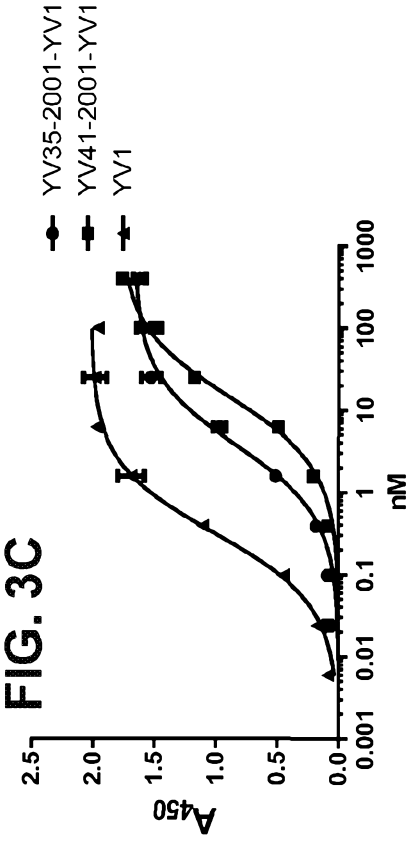
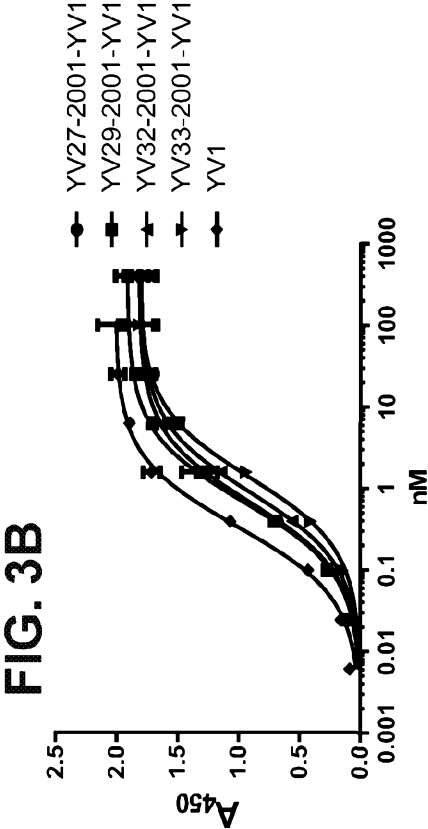
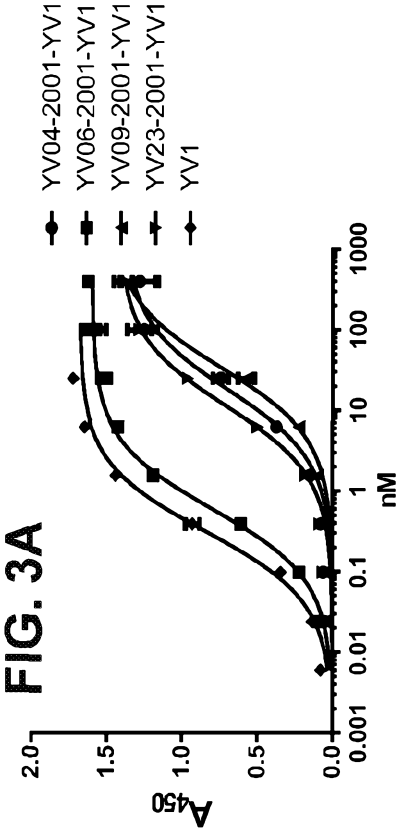
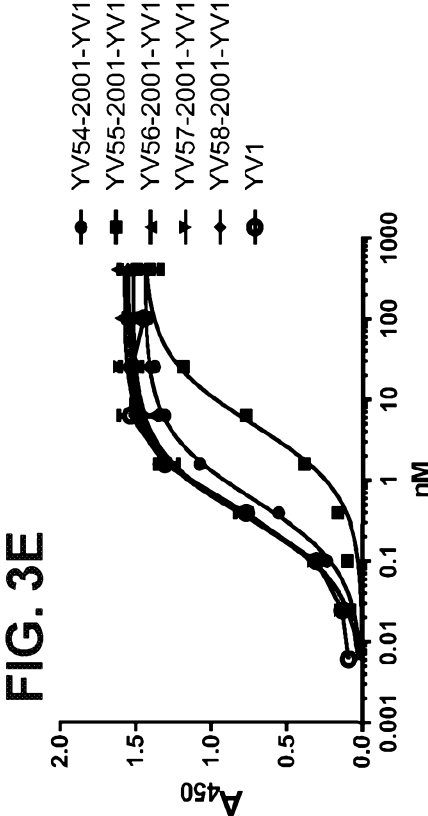
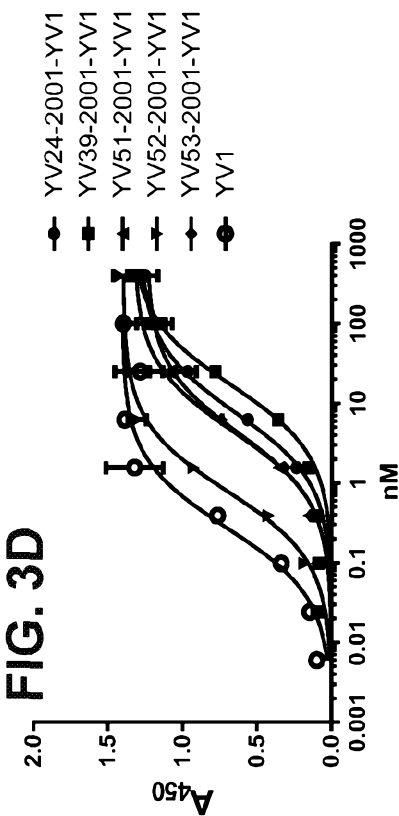


FIG. 4A

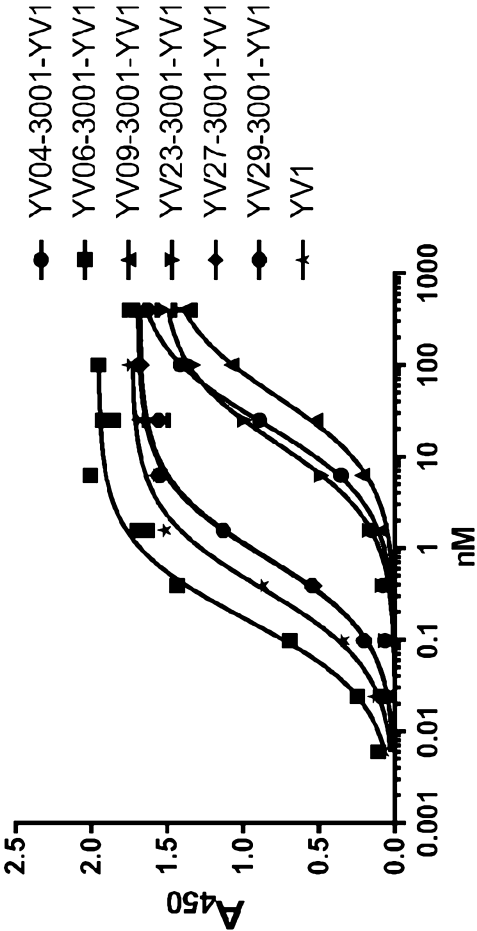


FIG. 4C

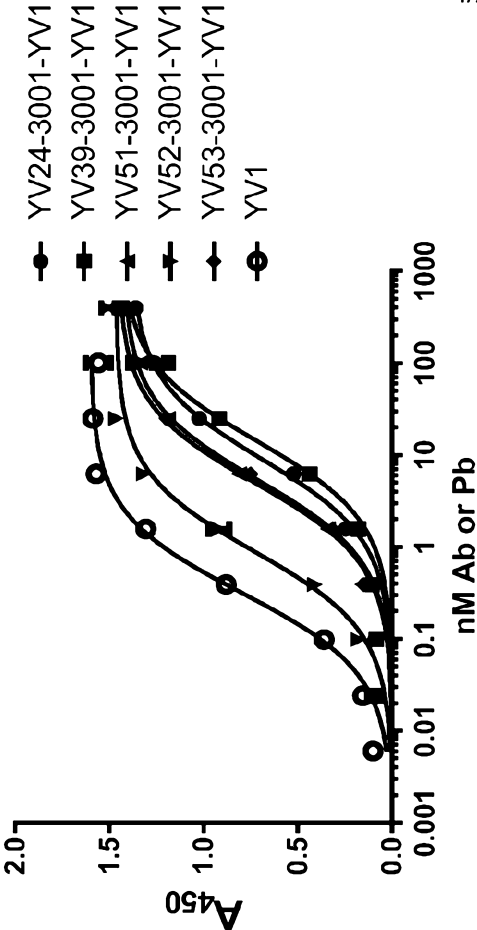


FIG. 4B

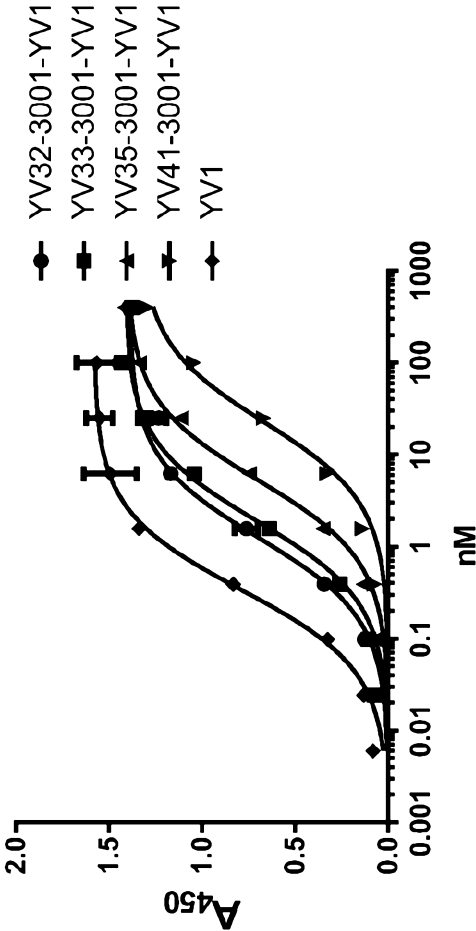


FIG. 4D

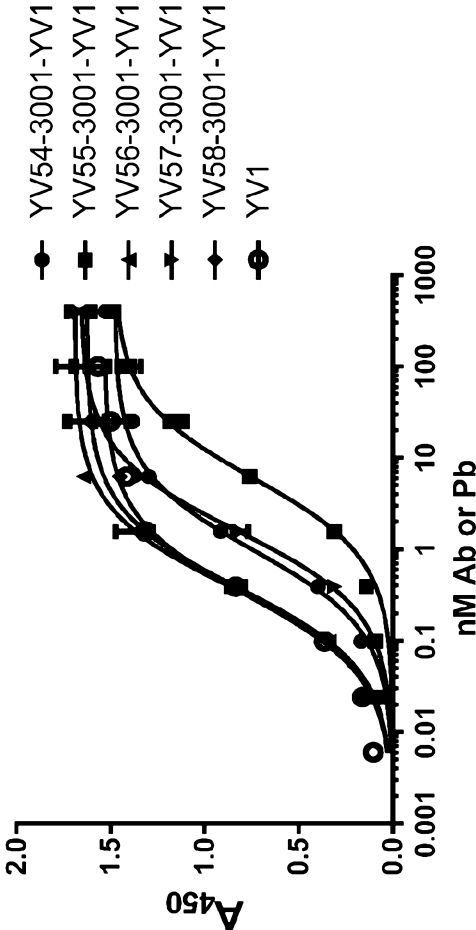


FIG. 5A

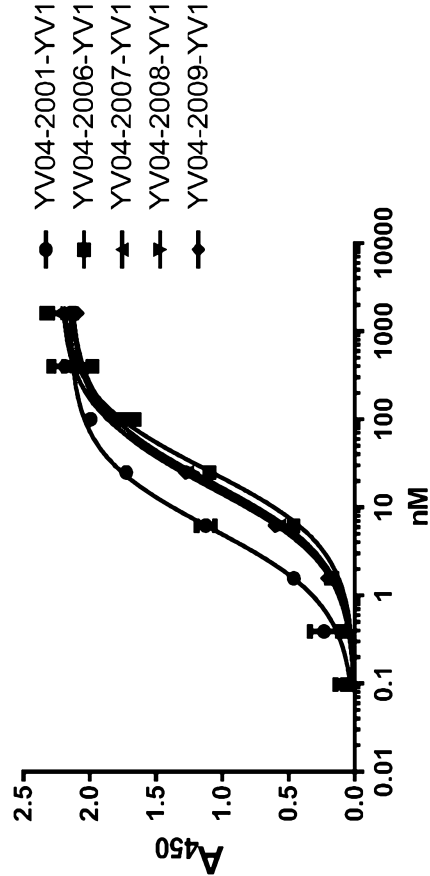


FIG. 5B

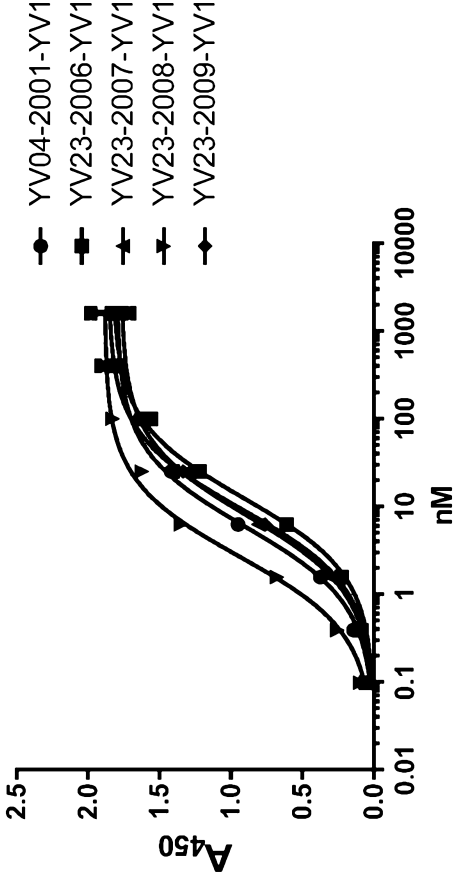


FIG. 5C

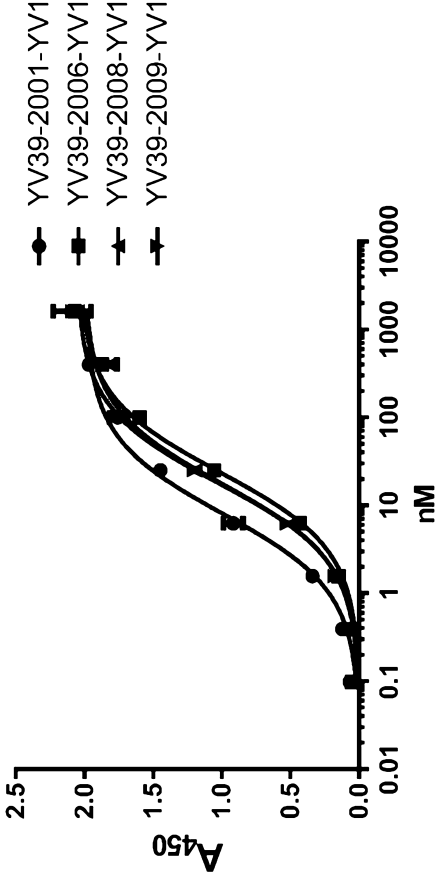


FIG. 5D

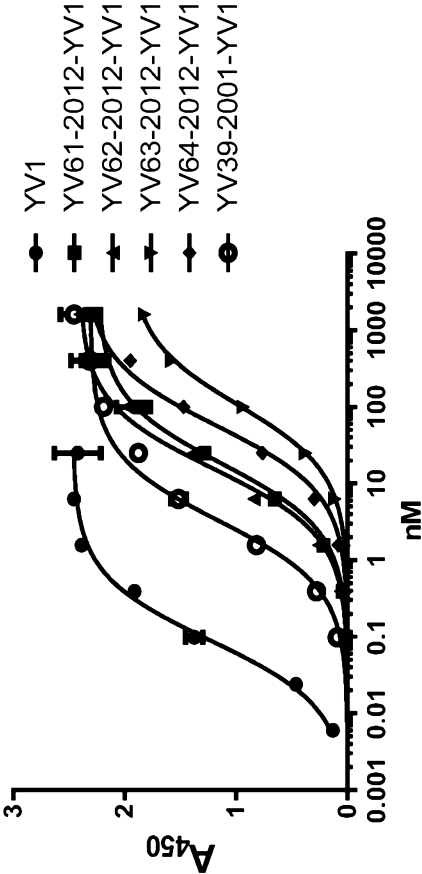


FIG. 5E

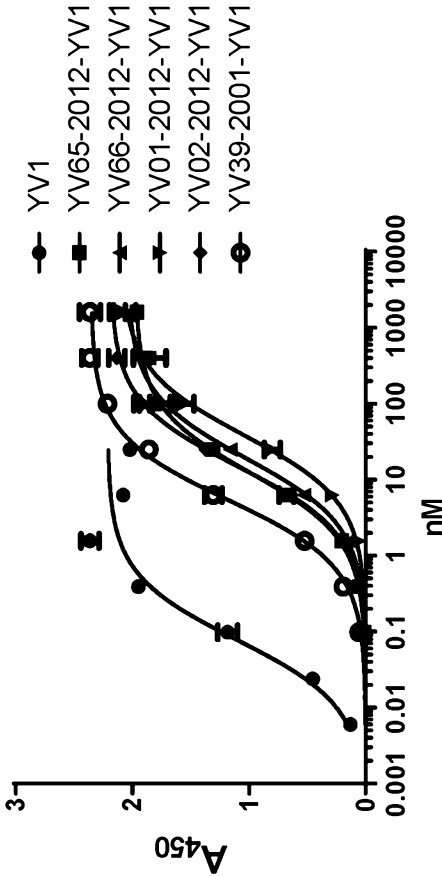


FIG. 5F

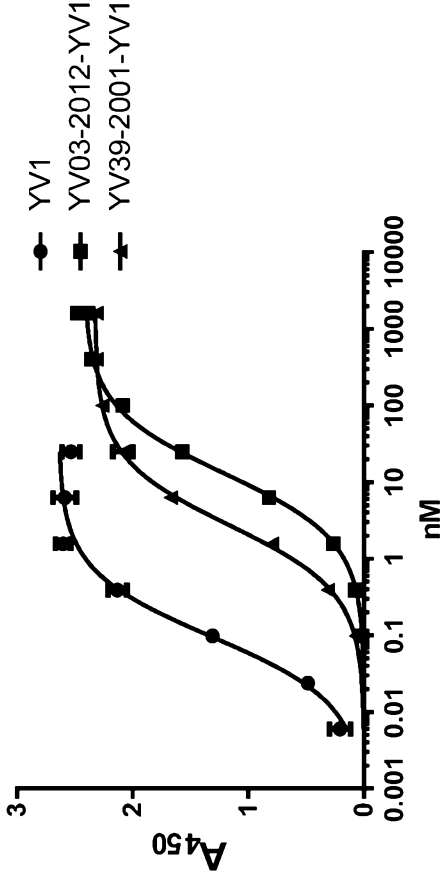


FIG. 6A

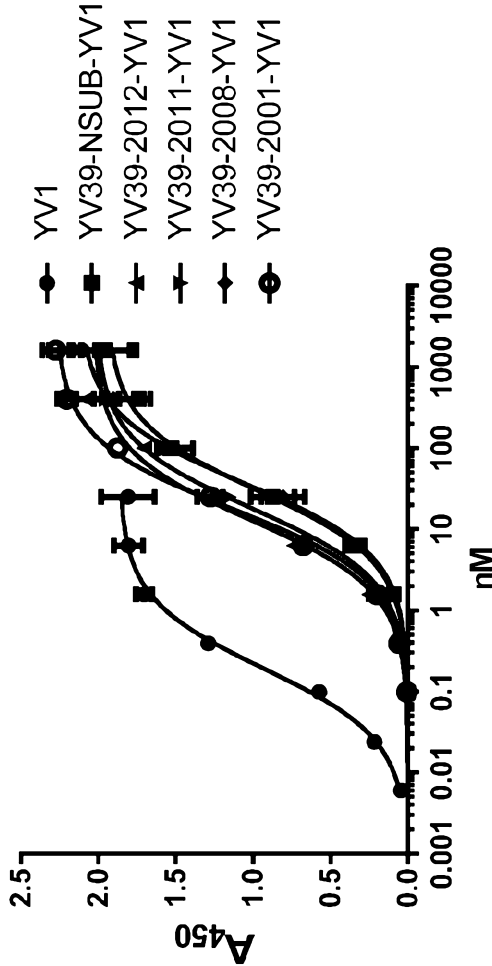


FIG. 6B

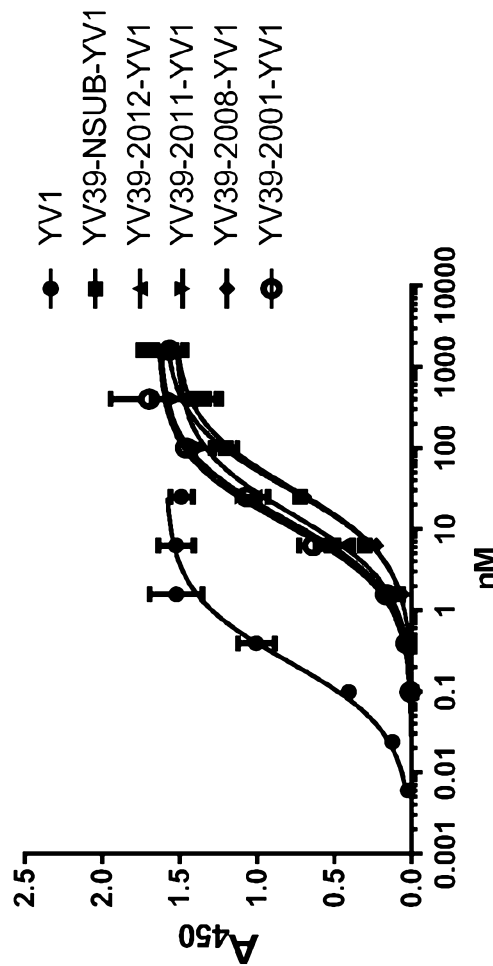


FIG. 7B

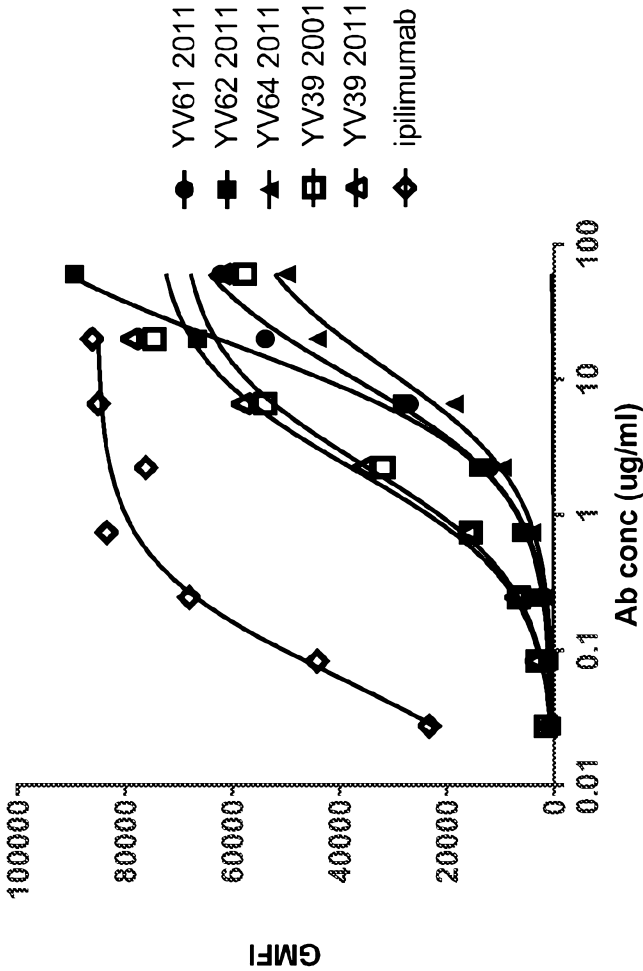


FIG. 7A

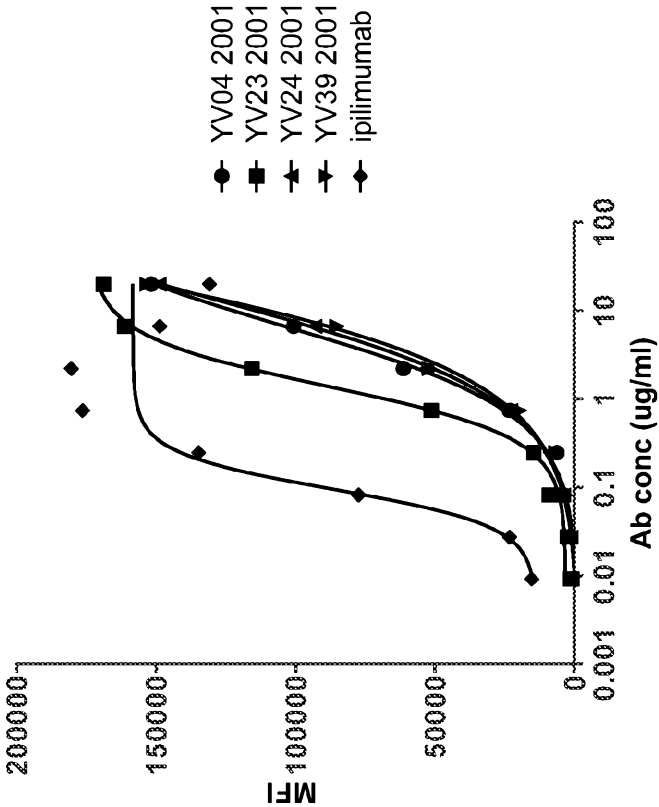


FIG. 7C

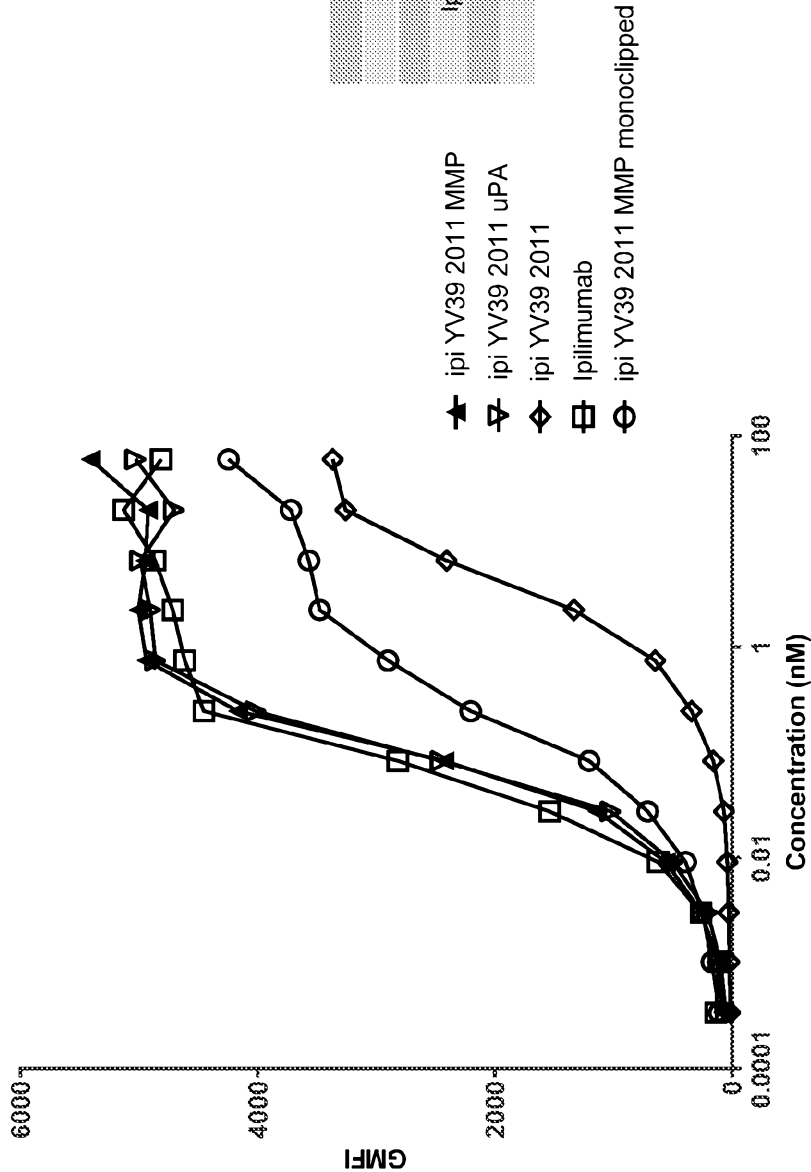
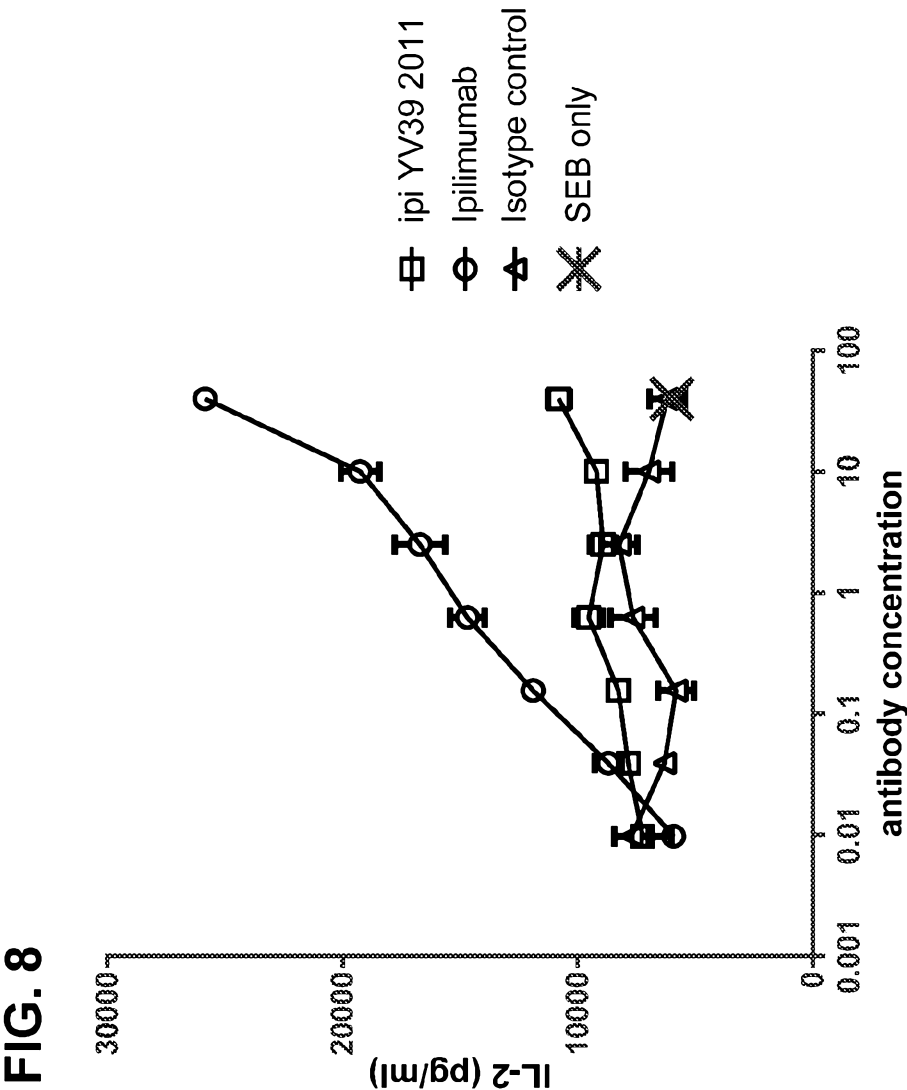


FIG. 7D

Sample	EC50 (nM)
ipi YV39 2011	22
Ipilimumab	0.54
ipi YV39 2011 MMP monoclonal	2.8
ipi YV39 2011 MMP clipped	0.65
ipi YV39 2011 μ PA clipped	0.76

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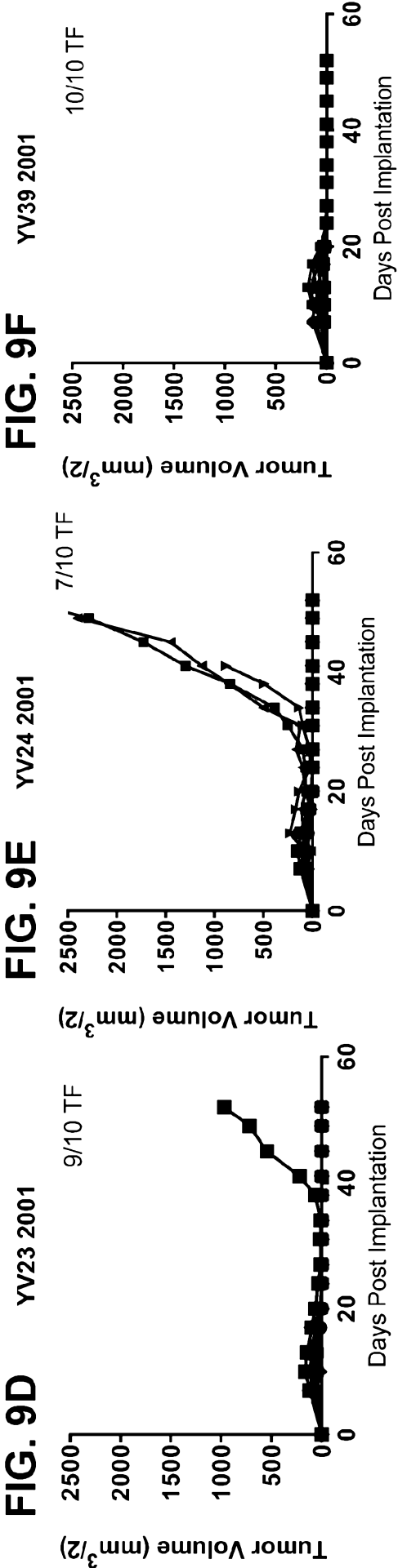
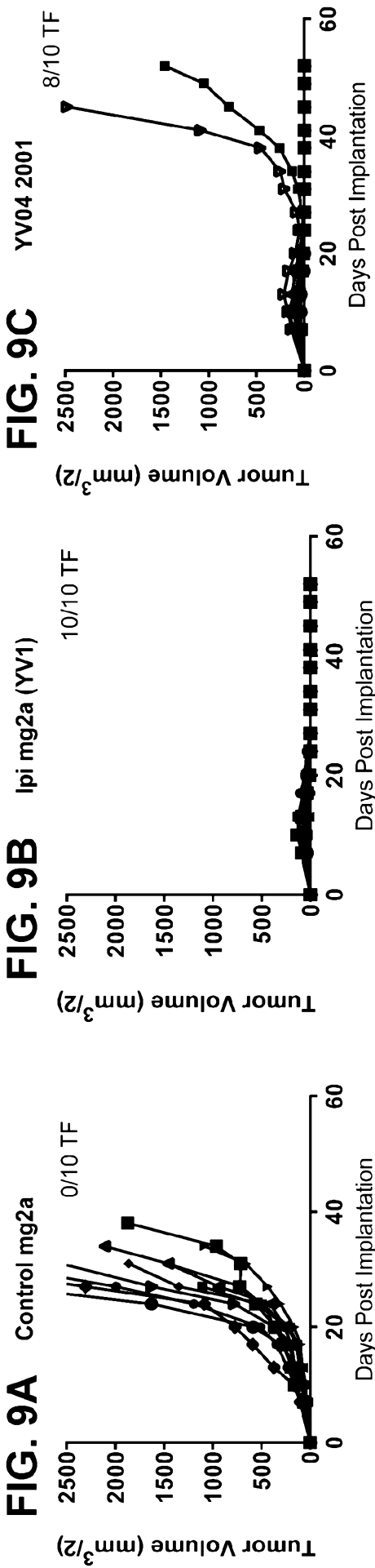


FIG. 10A

hg1

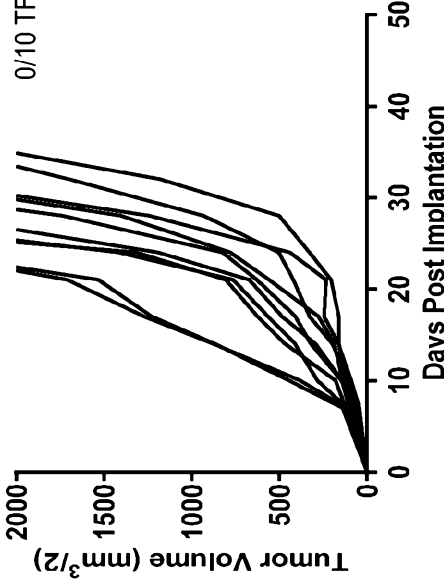


FIG. 10B

lpi hg1

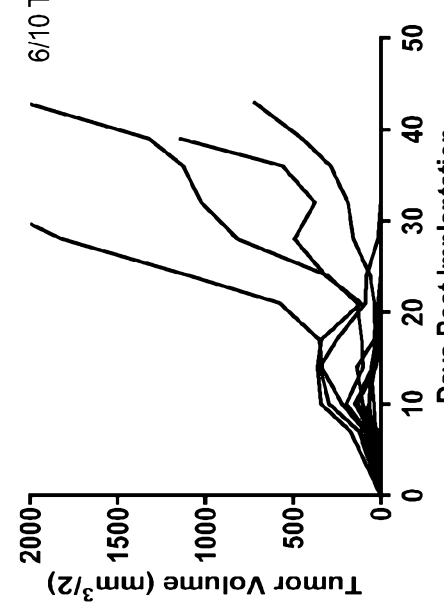


FIG. 10C

YV39 2001

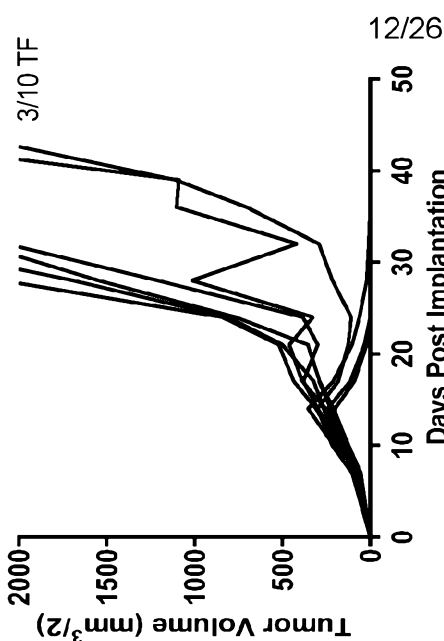


FIG. 10D

YV39 2012

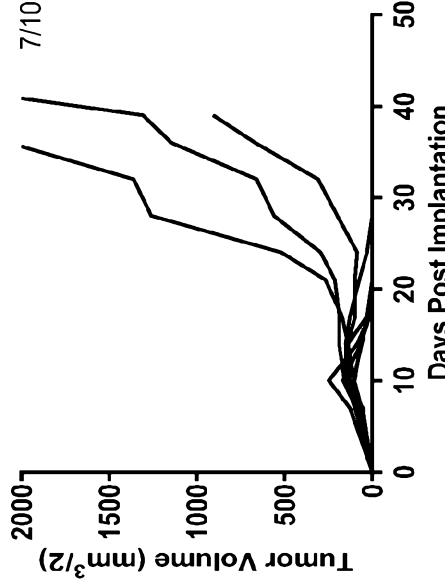


FIG. 10E

YV39 2011

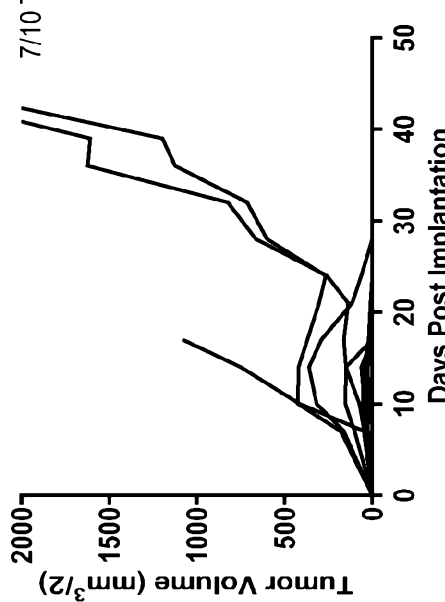
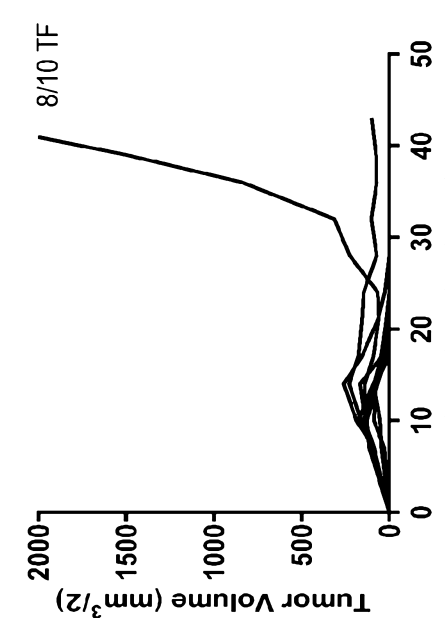
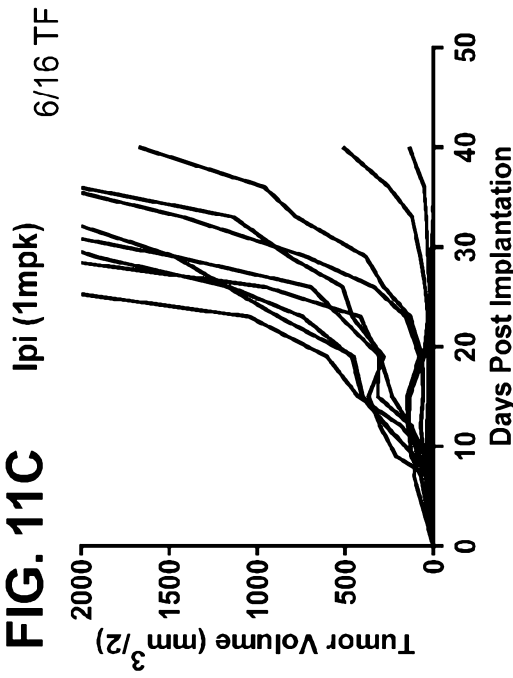
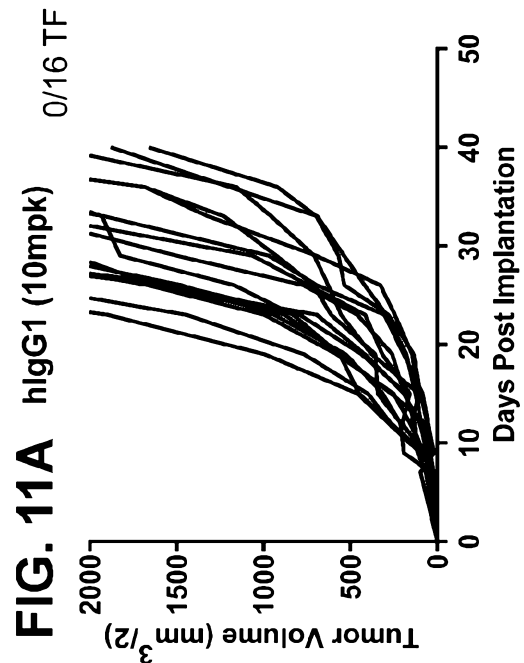
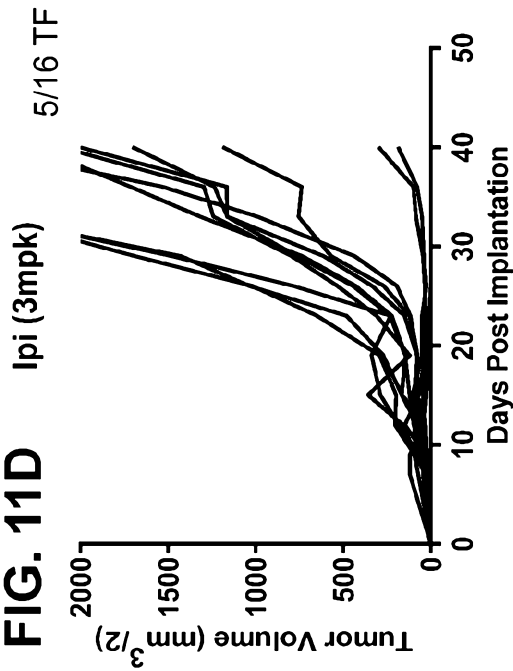
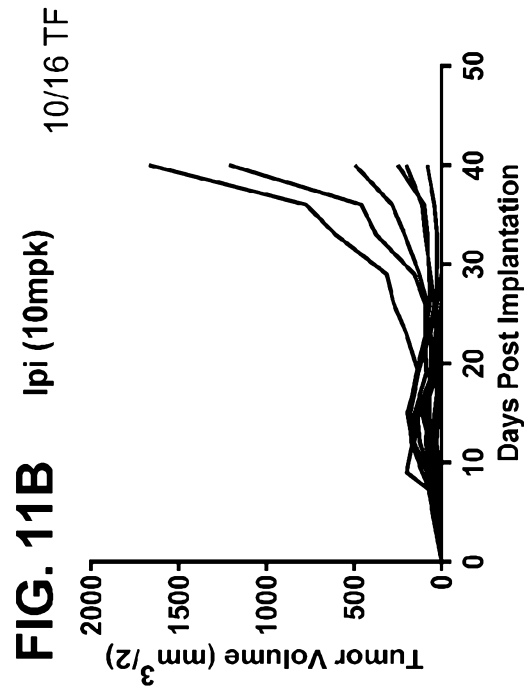


FIG. 10F

YV39 2008





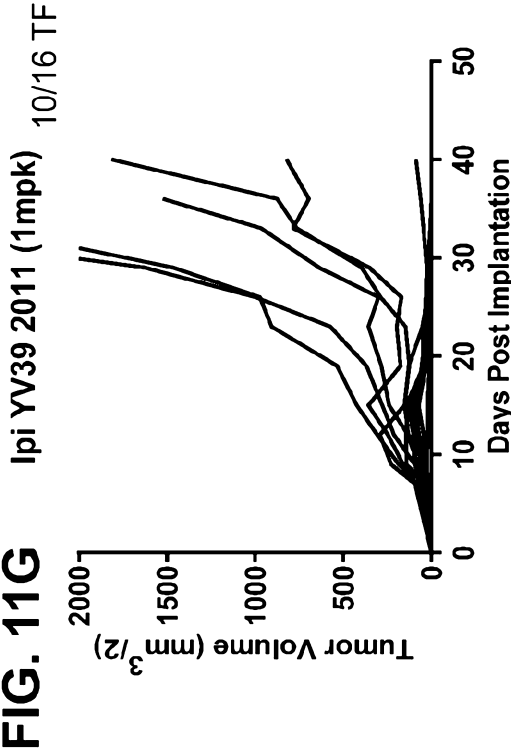
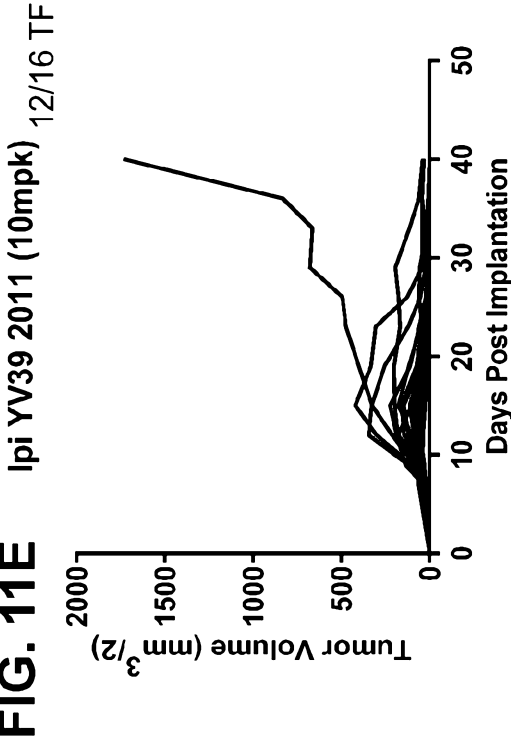
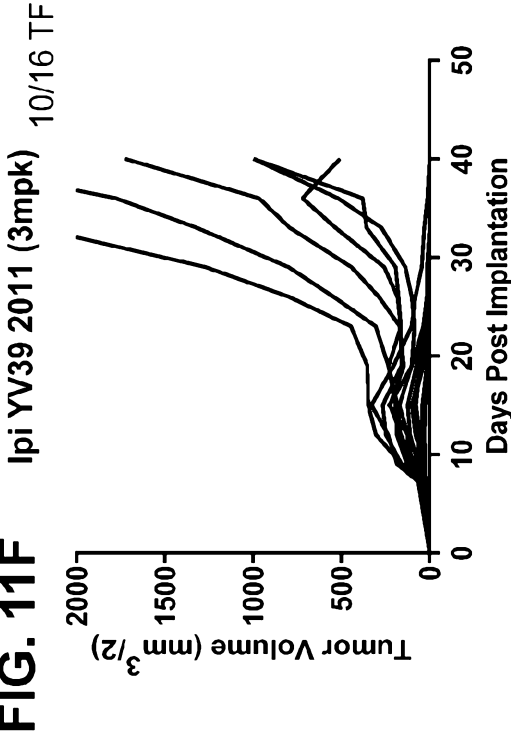


FIG. 12A FoxP3+ of CD4+ (Tumor)

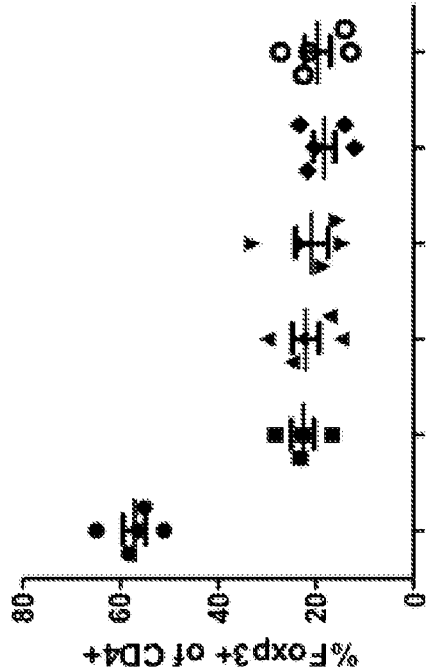


FIG. 12B FoxP3+ of CD45+ (Tumor)



FIG. 12C FoxP3+ of CD4+ (Spleen)

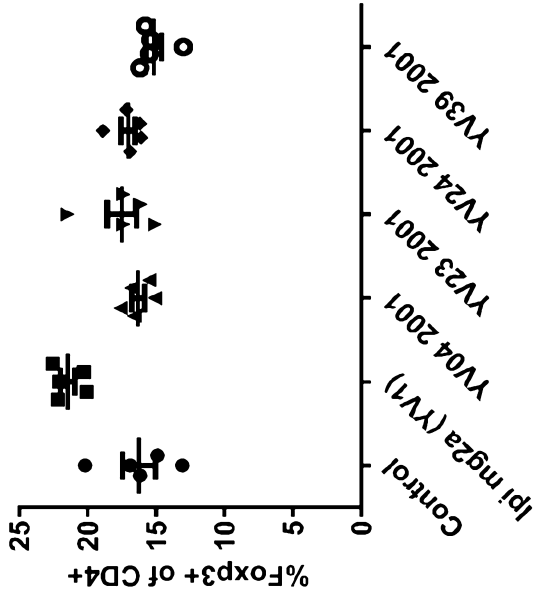


FIG. 12D FoxP3+ of CD45+ (Spleen)

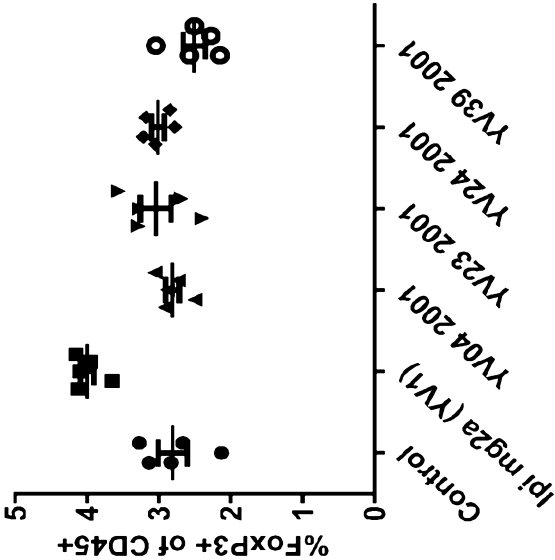


FIG. 12E ICOS+ of Tregs (Spleen)

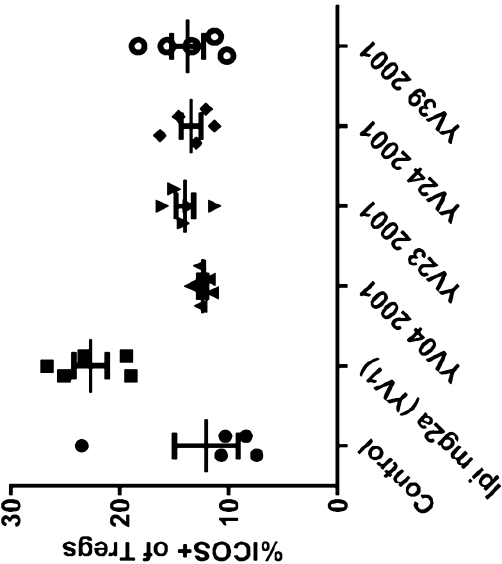


FIG. 12F Ki-67+ of Tregs (Spleen)

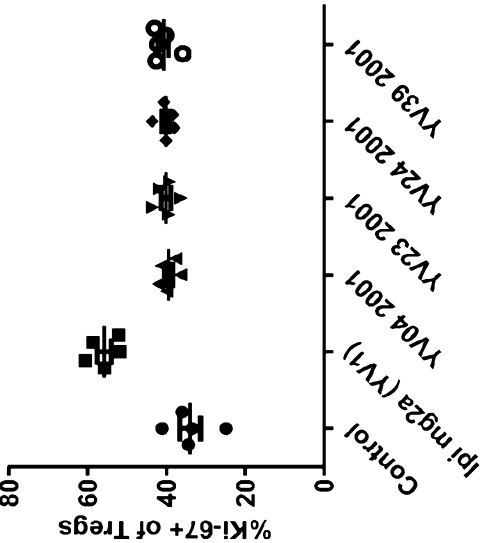


FIG. 13A

FoxP3+ of CD4+ (Tumor)

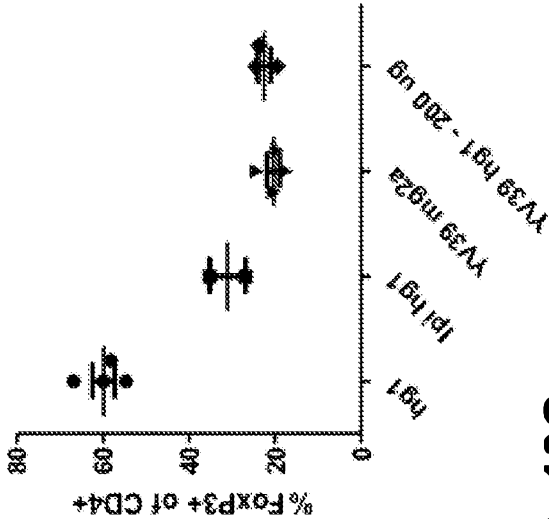


FIG. 13B

FoxP3+ of CD45+ (Tumor)

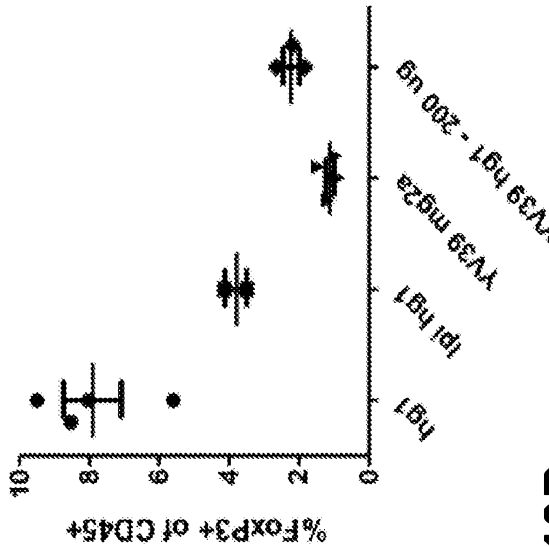


FIG. 13C

FoxP3+ of CD4+ (Spleen)

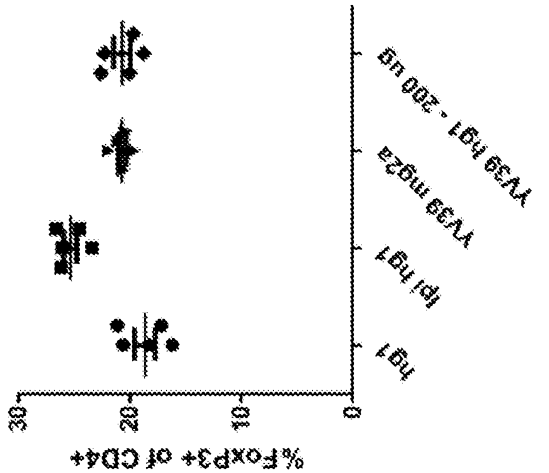


FIG. 13D

ICOS+ of Tregs (Spleen)

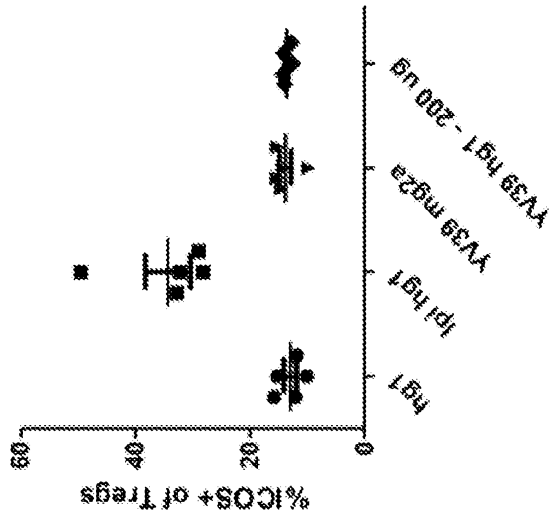


FIG. 13E

Ki-67+ of Tregs (Spleen)

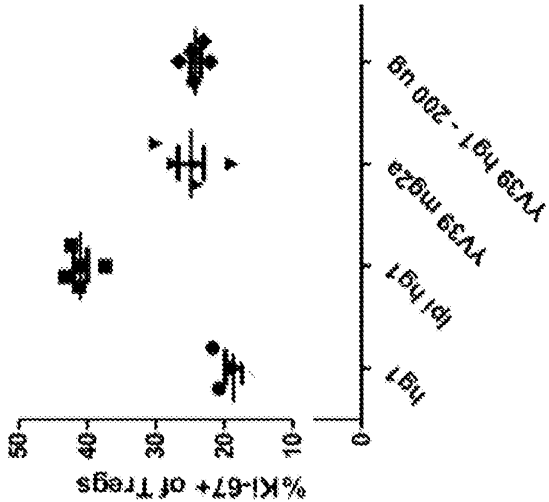


FIG. 14A FoxP3+ of CD4+ (Tumor)

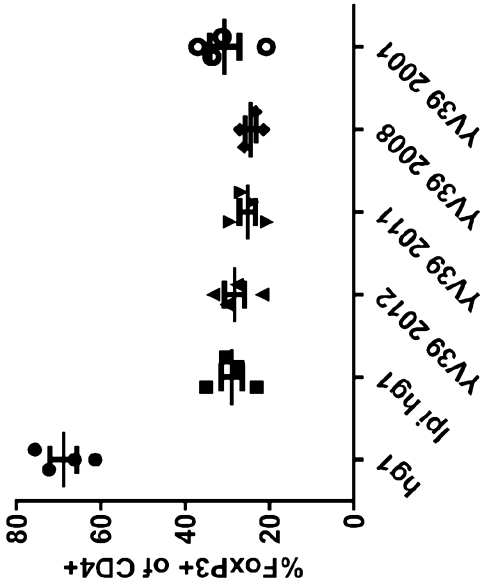


FIG. 14B FoxP3+ of CD45+ (Tumor)

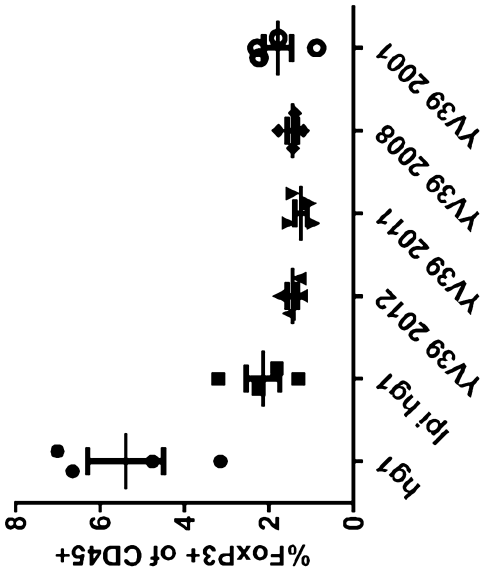


FIG. 14C CD4 effector+ of CD45+ (Tumor)

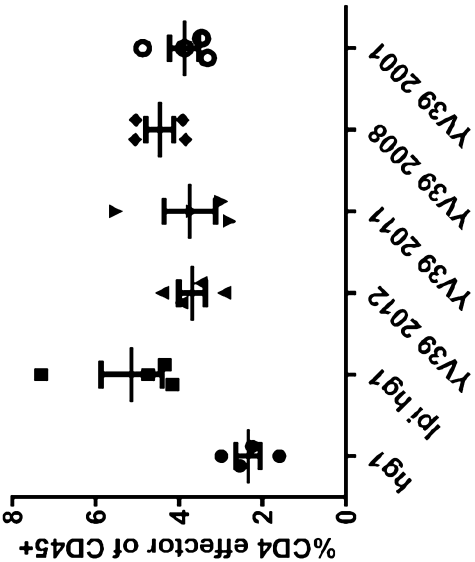


FIG. 14D FoxP3+ of CD4+ (Spleen)

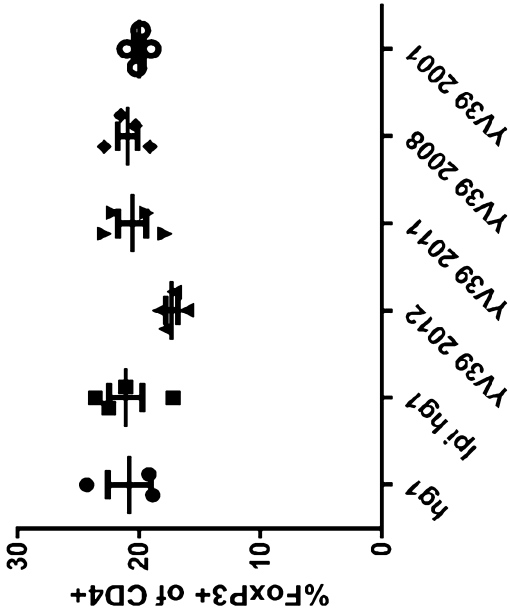


FIG. 14E FoxP3+ of CD45+ (Spleen)

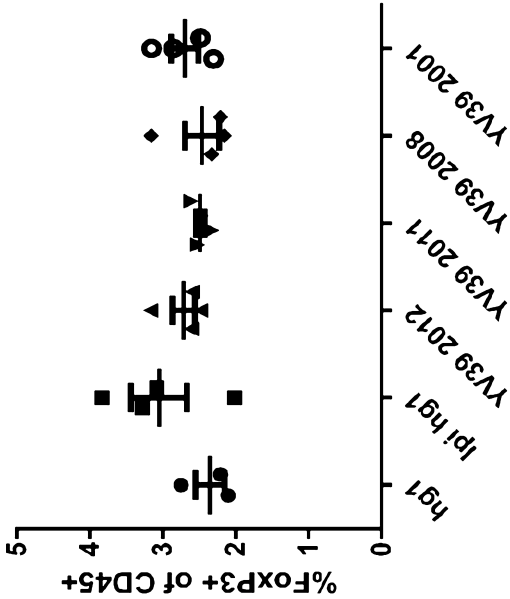


FIG. 14F ICOS+ of Tregs (Spleen)

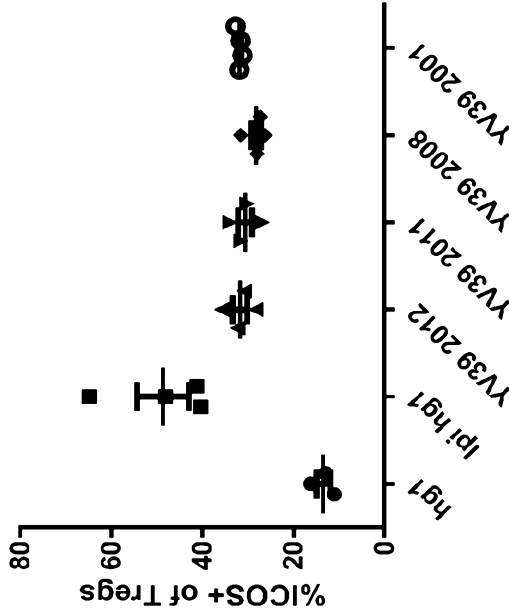


FIG. 14G Ki-67+ of Tregs (Spleen)

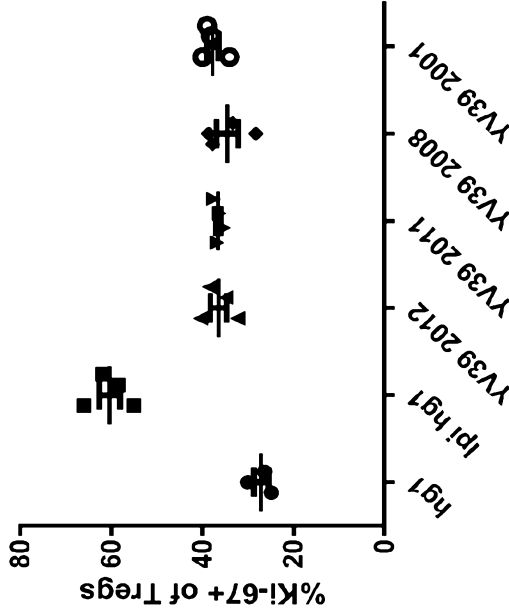


FIG. 15
Tumor Tregs

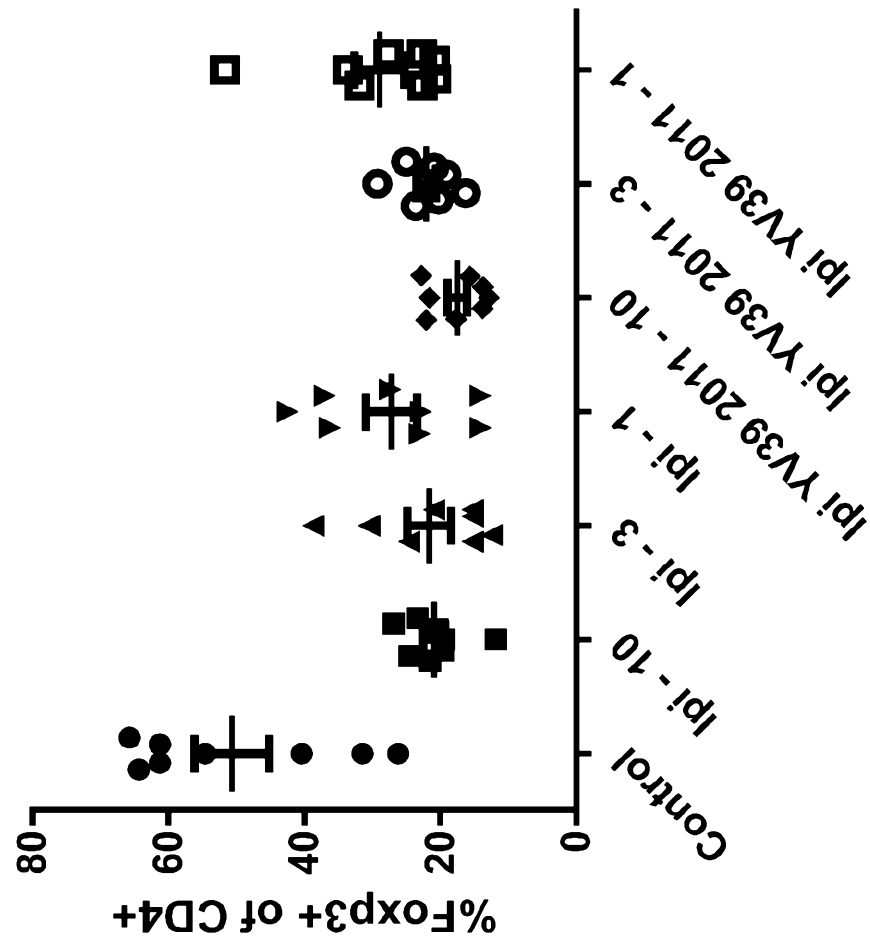


FIG. 16A

Spleen-ICOS on Tregs

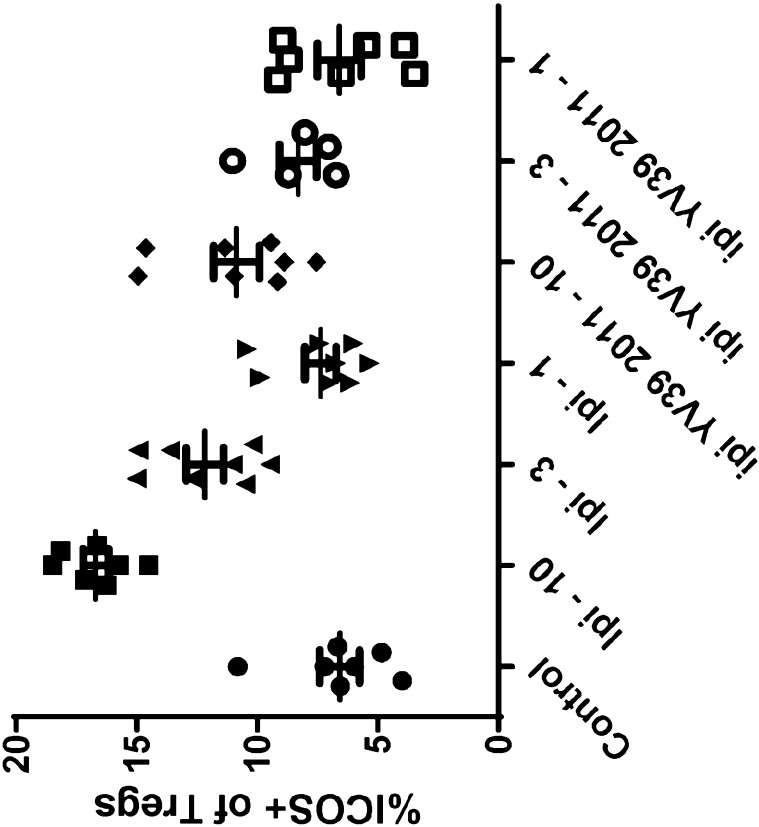


FIG. 16B

Spleen Ki-67 on Tregs

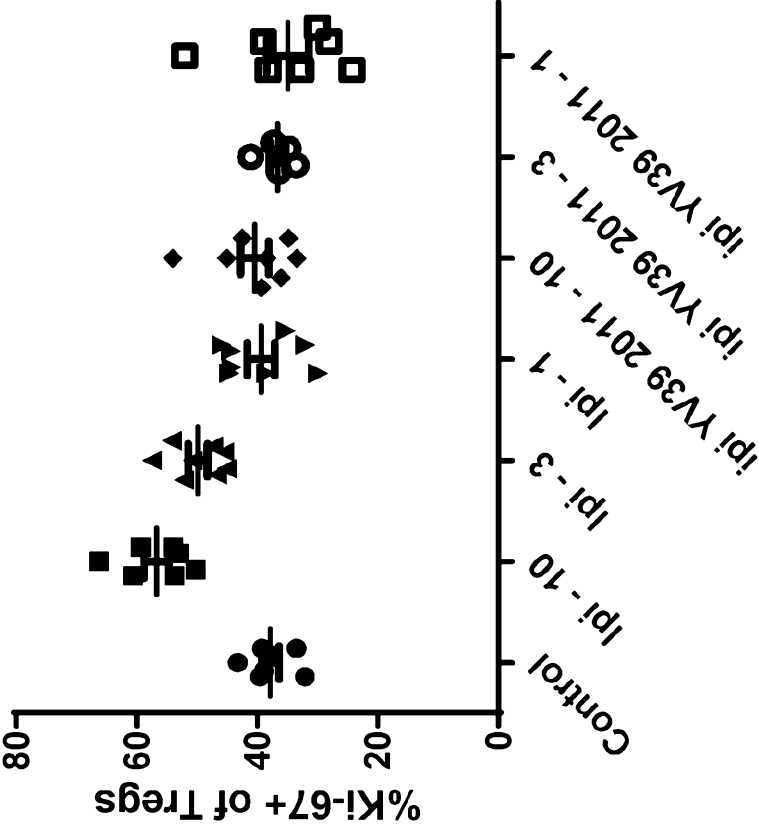


FIG. 17A

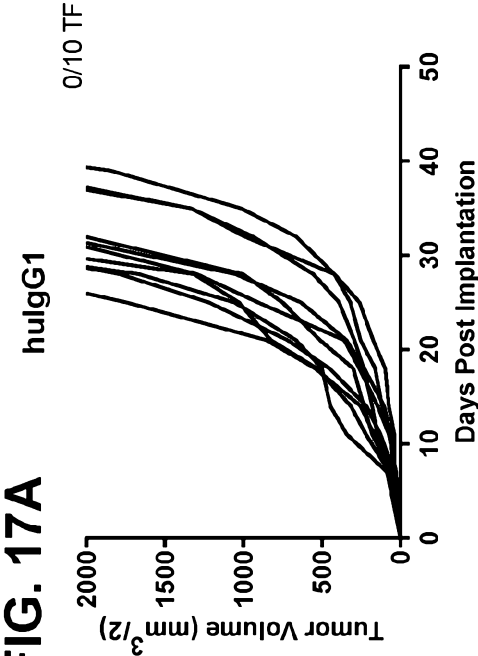


FIG. 17B

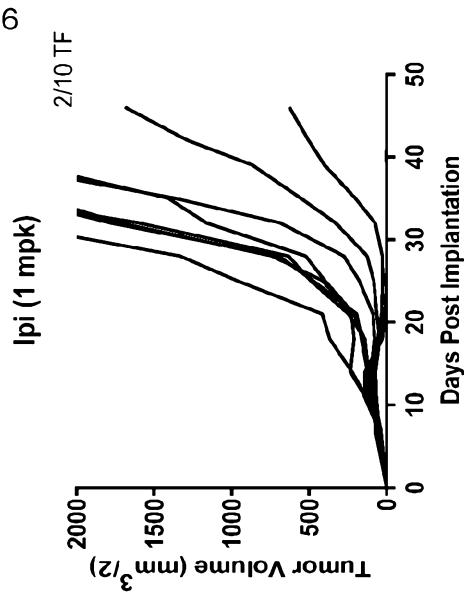
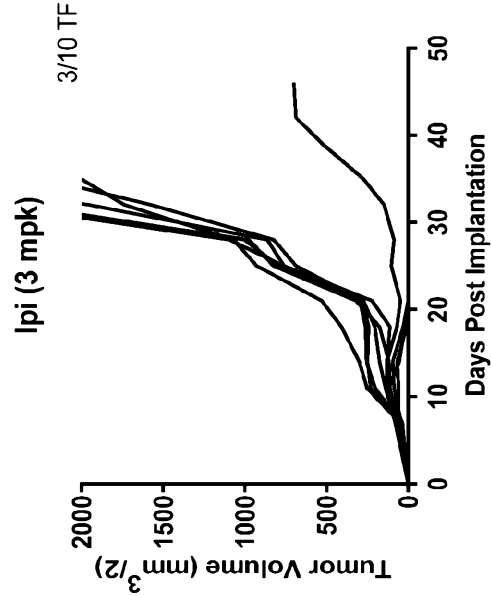
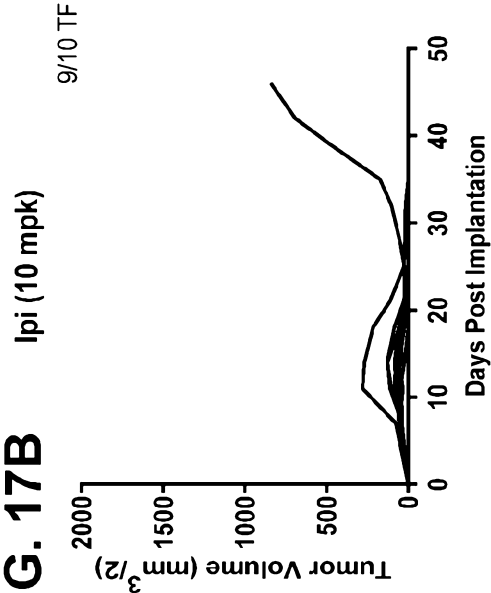


FIG. 17C

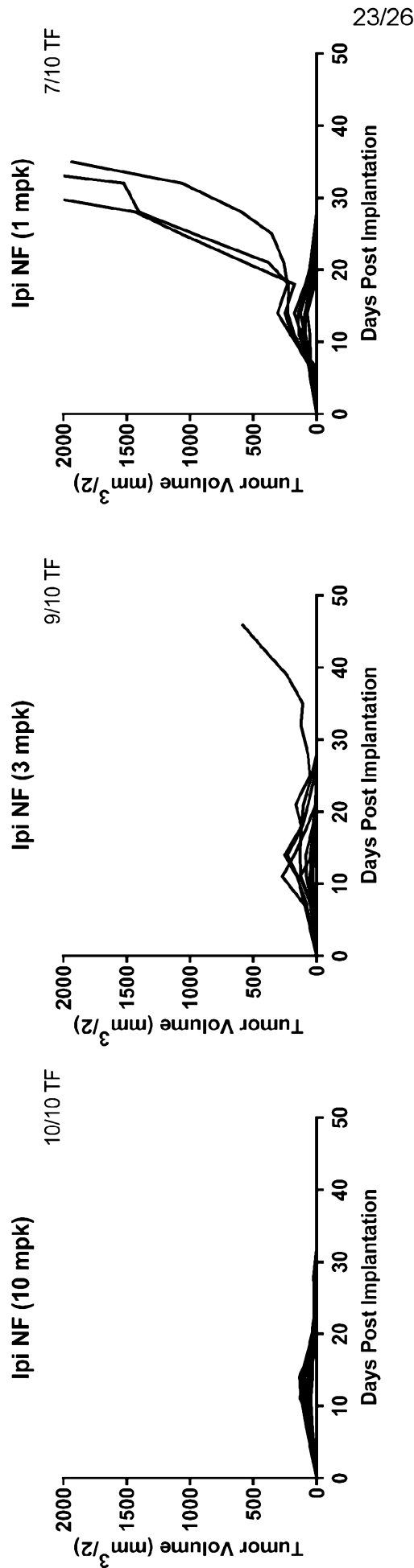


FIG. 17D

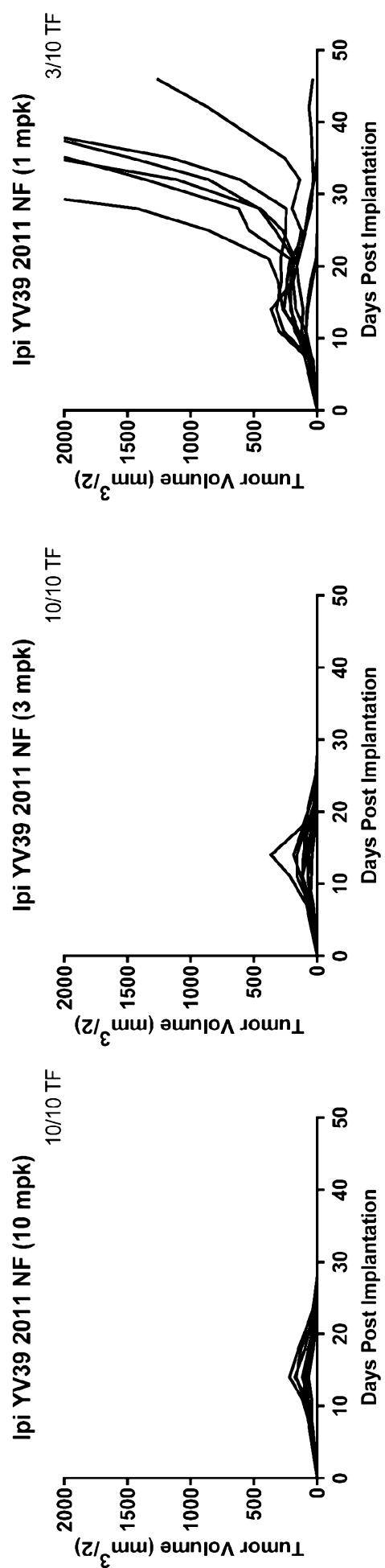


FIG. 18

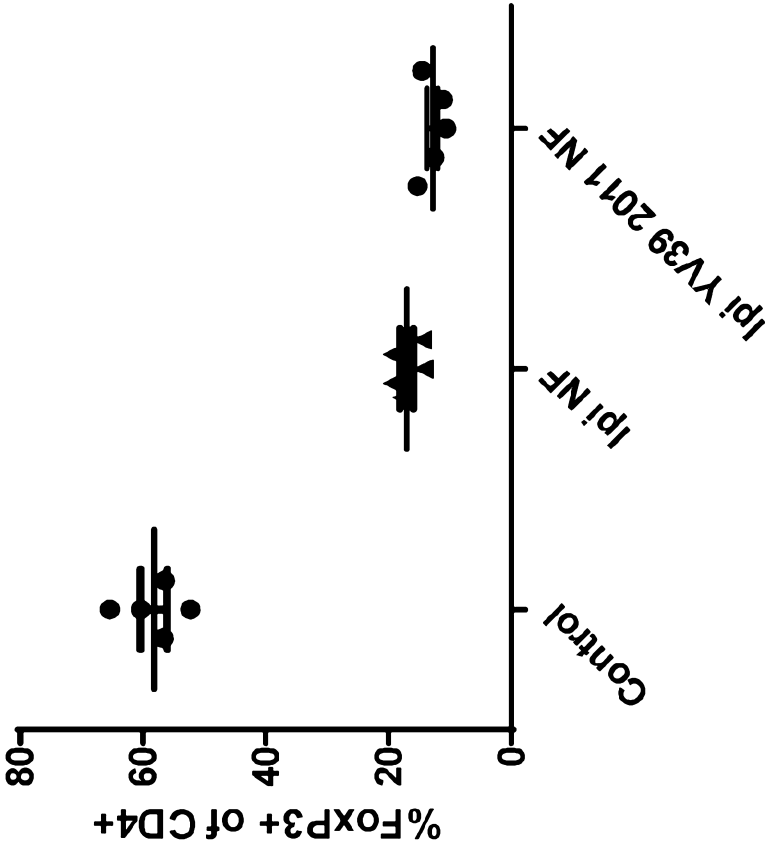
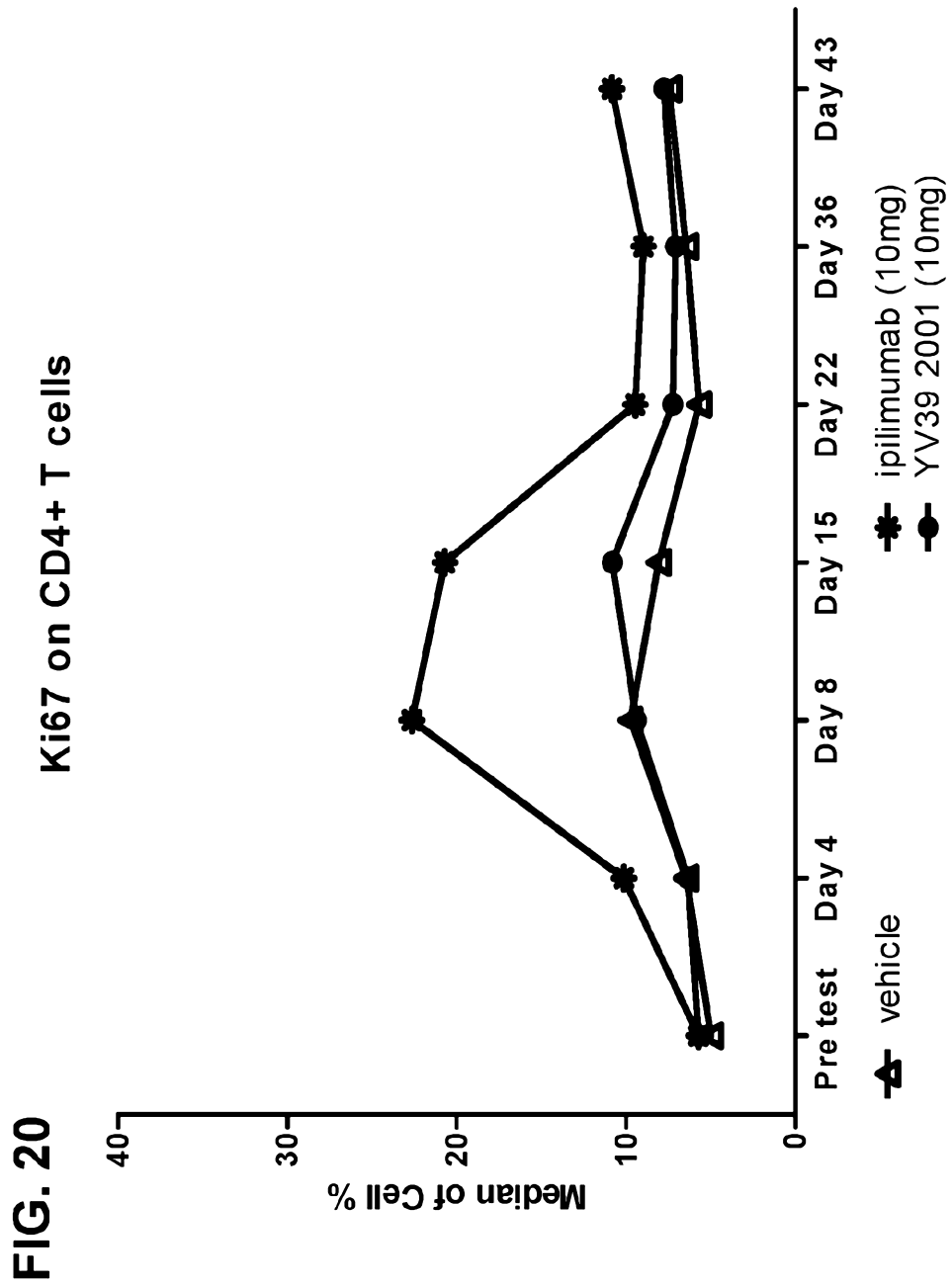


FIG. 19

	FcR	IgG1 Ipi (nM)	NF Ipi NF (nM)
Human FcR	hCD64	0.2	0.18
	hCD32a-H131	920	900
	hCD32a-R131	1100	730
	hCD32b	>5000	4200
	hCD16a-V158	310	9.5
	hCD16a-F158	4600	190
	hCD16B-NA1	>5000	1800
	hCD16B-NA2	4200	110
Cyno FcR	cyCD64	11	5.6
	cyCD32a	2700	2300
	cyCD32b	1900	2000
	cyCD16	370	7.5
Mouse FcR	mCD64	62	69
	mCD32	1300	1100
	mCD16	3100	2700
	mFcγRIV	29	6.3

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<170> PatentIn version 3.5

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<213> Artificial Sequence

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Asp	Phe	Ser	Cys	Leu	His	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 2

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV2

<400> 2

Gln	Pro	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10					15

<210> 3

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV3

<400> 3

Leu	His	Cys	Arg	Thr	Gln	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 4
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<400> 4

Leu	His	Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 5
 <211> 9
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 5

Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys
1				5				

<210> 6
 <211> 10
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<400> 6

Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys
1			5						10

<210> 7
 <211> 7
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 <213> Artificial Sequence

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<400> 7

Cys Pro Asn His Pro Met Cys
1 5

<210> 8

<211> 15

<212> PRT

<213> Artificial Sequence

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<223> Masking Moiety - YV8

<400> 8

Gly Thr Ala Cys Thr Tyr Ser Phe Phe Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 9

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<213> Artificial Sequence

<220>

<223> Masking Moiety - YV9

<400> 9

Phe Gly Thr Ala Cys Pro Asn His Pro Met Cys His Asp Trp Gln
1 5 10 15

<210> 10

<211> 15

<212> PRT

<213> Artificial Sequence

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<223> Masking Moiety - YV10

<400> 10

Ser Ala Cys Ala Tyr Trp Met Phe Gly Val Asn Leu Cys Pro Tyr
1 5 10 15

<210> 11

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<220>

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<400> 11

Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys
1				5					10	

<210> 12

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV12

<400> 12

Cys	Arg	Thr	Gln	Ile	Tyr	Gly	Tyr	Asn	Leu	Cys
1				5					10	

<210> 13

<211> 15

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<223> Masking Moiety - YV13

<400> 13

Leu	His	Cys	Arg	Thr	Gln	Ile	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 14

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV14

<400> 14

Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Trp	Gln
1				5					10	

<210> 15
<211> 14
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV15

<400> 15

Gly Thr Ala Cys Pro Asn His Pro Met Cys His Asp Trp Gln
1 5 10

<210> 16
<211> 13
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<220>
<223> Masking Moiety - YV16

<400> 16

Cys Ala Tyr Trp Met Phe Gly Val Asn Leu Cys Pro Tyr
1 5 10

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

<220>
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<400> 17

Gln Glu Cys His Leu Tyr Met Tyr Gly Val Asn Leu Cys Pro Tyr
1 5 10 15

<210> 18
<211> 13
<212> PRT
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<220>
<223> Masking Moiety - YV18

<400> 18

Cys His Leu Tyr Met Tyr Gly Val Asn Leu Cys Pro Tyr
1 5 10

<210> 19
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<400> 19

Gly Gln Cys Gln Phe Tyr Met Phe Gly Tyr Asn Leu Cys Pro Tyr
1 5 10 15

<210> 20
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<223> Masking Moiety - YV20

<400> 20

Leu Ser Thr Cys Met Tyr Ser Phe Phe Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 21
<211> 12
<212> PRT
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<220>
<223> Masking Moiety - YV21

<400> 21

Cys Leu His Ser Met Tyr Asn Val Cys Leu Asp Pro
1 5 10

<210> 22
<211> 9
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<220>

<223> Masking Moiety - YV22

<400> 22

Cys Leu His Ser Met Tyr Asn Val Cys
1 5

<210> 23

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV23

<400> 23

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro
1 5 10

<210> 24

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV24

<400> 24

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Asp Pro
1 5 10

<210> 25

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV25

<400> 25

Cys Met Tyr Ser Phe Phe Asn Val Cys Leu Asp Pro
1 5 10

<210> 26

<211> 9

<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV26

<400> 26

Cys Met Tyr Ser Phe Phe Asn Val Cys
1 5

<210> 27
<211> 12
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV27

<400> 27

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys
1 5 10

<210> 28
<211> 13
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV28

<400> 28

Cys Ala Gln Leu Tyr Gly Tyr Ser Met Cys Pro His Thr
1 5 10

<210> 29
<211> 13
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV29

<400> 29

Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Ala His Thr
1 5 10

<210> 30
 <211> 13
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<220>
 <223> Masking Moiety - YV30

<400> 30

Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	Ala	Thr
1				5					10			

<210> 31
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 <223> Masking Moiety - YV31

<400> 31

Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10			

<210> 32
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 <213> Artificial Sequence

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 <223> Masking Moiety - YV32

<400> 32

Cys	Pro	Asn	His	Pro	Leu	Cys	His	Asp	Trp	Gln
1				5					10	

<210> 33
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 <212> PRT
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<220>
 <223> Masking Moiety - YV33

<400> 33

Cys	Pro	Asn	His	Pro	Met	Cys	Ala	Asp	Trp	Gln
1				5					10	

<210> 34

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV34

<400> 34

Cys	Pro	Asn	His	Pro	Met	Cys	His	Ala	Trp	Gln
1				5					10	

<210> 35

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV35

<400> 35

Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Ala	Gln
1				5					10	

<210> 36

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV36

<400> 36

Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Trp	Ala
1				5					10	

<210> 37

<211> 10

<212> PRT

<213> Artificial Sequence

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<223> Masking Moiety - YV37

<400> 37

Gly	Thr	Ala	Cys	Pro	Asn	His	Pro	Met	Cys
1				5					10

<210> 38

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV38

<400> 38

Leu	His	Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys
1				5					10			

<210> 39

<211> 13

<212> PRT

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<220>

<223> Masking Moiety - YV39

<400> 39

Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10			

<210> 40

<211> 13

<212> PRT

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<223> Masking Moiety - YV40

<400> 40

Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys	Ala	Tyr
1				5					10			

<210> 41
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 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety - YV41

<400> 41

Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Ala
1				5					10			

<210> 42
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<220>
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<400> 42

Phe	Gly	Thr	Ala	Cys	Pro	Asn	His	Pro	Leu	Cys	His	Asp	Trp	Gln
1				5					10					15

<210> 43
 <211> 11
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety - YV43

<400> 43

Cys	Pro	Asn	His	Pro	Leu	Cys	His	Asp	Phe	Gln
1				5					10	

<210> 44
 <211> 11
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<220>
 <223> Masking Moiety - YV44

<400> 44

Cys Pro Asn His Pro Leu Cys His Asp Tyr Gln
1 5 10

<210> 45
<211> 9
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV45

<400> 45

Cys Pro Asn His Pro Leu Cys Pro Tyr
1 5

<210> 46
<211> 9
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV46

<400> 46

Cys Pro Asn His Pro Leu Cys Pro Ala
1 5

<210> 47
<211> 11
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety - YV47

<400> 47

Cys Met Tyr Ser Phe Phe Asn Val Cys Tyr Pro
1 5 10

<210> 48
<211> 11
<212> PRT
<213> Artificial Sequence

<220>

<223> Masking Moiety - YV48

<400> 48

Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Tyr	Ala
1				5				10		

<210> 49

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV49

<400> 49

Cys	Leu	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Tyr	Pro
1				5				10		

<210> 50

<211> 11

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV50

<400> 50

Cys	Leu	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Tyr	Ala
1				5				10		

<210> 51

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV51

<400> 51

Phe	Gly	Ala	Ala	Cys	Pro	Asn	His	Pro	Ile	Cys	His	Asp	Trp	Gln
1				5				10						15

<210> 52

<211> 15

<212> PRT
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<220>
 <223> Masking Moiety - YV52

<400> 52

Phe	Gly	Ala	Ala	Cys	Pro	Asn	His	Pro	Leu	Cys	His	Asp	Trp	Gln
1				5					10					15

<210> 53
 <211> 15
 <212> PRT
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<220>
 <223> Masking Moiety - YV53

<400> 53

Phe	Gly	Ala	Ala	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Ala	Gln
1				5					10					15

<210> 54
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<220>
 <223> Masking Moiety - YV54

<400> 54

Cys	Leu	His	Ser	Ala	Tyr	Asn	Ala	Cys	Leu	Asp	Pro
1				5					10		

<210> 55
 <211> 12
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 55

Cys	Ala	His	Ser	Ala	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10		

<210> 56
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 <212> PRT
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<220>
 <223> Masking Moiety - YV56

<400> 56

Cys	Leu	His	Ser	Ala	Tyr	Asn	Val	Cys	Ala	Asp	Pro
1				5					10		

<210> 57
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 <212> PRT
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<220>
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<400> 57

Cys	Leu	His	Ser	Ala	Tyr	Asn	Val	Cys	Leu	Ala	Pro
1				5					10		

<210> 58
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 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety - YV58

<400> 58

Cys	Leu	His	Ser	Ala	Tyr	Asn	Val	Cys	Leu	Asp	Ala
1				5					10		

<210> 59
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety - YV60

<400> 59

Lys	Asn	Thr	Cys	Thr	Tyr	Val	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 60

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV61

<400> 60

Tyr	Ile	Ser	Asp	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Gln
1				5					10					15

<210> 61

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV62

<400> 61

Phe	Arg	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Arg
1				5					10					15

<210> 62

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV63

<400> 62

Arg	Glu	Cys	His	Met	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 63

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV64

<400> 63

Ala	Val	Cys	His	Met	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Phe
1				5					10					15

<210> 64

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV65

<400> 64

Arg	Ser	Cys	Pro	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10					15

<210> 65

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - YV66

<400> 65

Gln	Pro	Cys	Ala	Gln	Met	Phe	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10					15

<210> 66

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 66

Thr	Ala	Lys	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 67
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 67

Asp	Phe	Ser	Cys	Leu	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10				15	

<210> 68
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 68

Asp	Val	Ser	Cys	Met	Tyr	Met	Met	Tyr	Asn	Phe	Cys	Leu	Asp	Pro
1				5					10				15	

<210> 69
 <211> 7
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 69

Cys	Pro	Asn	His	Pro	Met	Cys
1				5		

<210> 70
 <211> 11
 <212> PRT
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<220>
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<400> 70

Cys Met Tyr Ser Phe Phe Asn Val Cys Pro Tyr
1 5 10

<210> 71
<211> 11
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 71

Cys Met Tyr Ser Phe Phe Asn Val Cys Pro Ala
1 5 10

<210> 72
<211> 11
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 72

Cys Thr Tyr Ser Phe Phe Asn Val Cys Pro Tyr
1 5 10

<210> 73
<211> 11
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 73

Cys Thr Tyr Ser Phe Phe Asn Val Cys Pro Ala
1 5 10

<210> 74
<211> 15
<212> PRT
<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 74

Gly	Phe	Pro	Cys	Met	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 75

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 75

Gly	Leu	Ser	Cys	Met	Tyr	Ser	Met	Tyr	Gly	Tyr	Cys	Leu	Asp	Pro
1				5					10					15

<210> 76

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 76

Ile	Pro	Cys	Asp	Tyr	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 77

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 77

Gln	Val	Cys	His	Ala	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 78

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
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<400> 78

Arg	Met	Tyr	Cys	Thr	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 79
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 <213> Artificial Sequence

<220>
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<400> 79

Ala	Leu	Ser	Cys	Met	Tyr	Ile	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 80
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 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 80

Asp	Phe	Ser	Cys	Met	Tyr	Val	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 81
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 81

Asp	Phe	Ser	Cys	Val	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 82
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 <212> PRT
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<220>
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<400> 82

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Tyr	Asp	Tyr	Arg
1				5					10					15

<210> 83
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 <212> PRT
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<220>
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<400> 83

Asp	Met	Asn	Thr	Cys	Pro	Arg	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5					10					15

<210> 84
 <211> 15
 <212> PRT
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<220>
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<400> 84

Asp	Ser	Arg	Cys	Met	Tyr	Val	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 85
 <211> 15
 <212> PRT
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<220>
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<400> 85

Glu His Leu Cys Thr Tyr Ser Phe Tyr Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 86

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 86

Glu Leu Ser Cys Val Tyr Ser Met Phe Gly Phe Cys Leu Asp Pro
1 5 10 15

<210> 87

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 87

Phe Thr Asn Asn Cys Pro Tyr His Pro Met Cys His Asp Tyr Leu
1 5 10 15

<210> 88

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 88

Gly Phe Ser Cys Thr Tyr Ile Met Tyr Asp Val Cys Leu Asp Pro
1 5 10 15

<210> 89

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

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<400> 89

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

<210> 90

<211> 15

<212> PRT

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<220>

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<400> 90

His	Phe	Ser	Cys	Met	Tyr	Ile	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 91

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 91

Leu	His	Cys	Gly	Met	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Lys
1				5					10					15

<210> 92

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 92

Leu	Pro	Cys	Gln	Met	Trp	Met	Phe	Gly	His	Asn	Leu	Cys	Pro	His
1				5					10					15

<210> 93
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 93

Leu	Pro	Cys	Thr	Met	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5				10						15

<210> 94
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 94

Leu	Thr	Cys	His	His	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5				10						15

<210> 95
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<220>
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<400> 95

Asn	Phe	Ser	Cys	Met	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5				10						15

<210> 96
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 96

Asn Asn His Cys Met Tyr Ser Phe Phe Asn Ile Cys Leu Asp Pro
1 5 10 15

<210> 97
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 97

Asn Arg Ser Cys Met Tyr Ile Met Tyr Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 98
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 98

Asn Ser Cys Thr Met Phe Met Phe Gly Val Asn Leu Cys Pro Tyr
1 5 10 15

<210> 99
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 99

Asn Thr Cys Glu Leu Tyr Met Phe Gly Val Asn Leu Cys Pro Tyr
1 5 10 15

<210> 100
<211> 15
<212> PRT
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<220>

<223> Masking Moiety

<400> 100

Gln	His	Cys	Asp	Met	Trp	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 101

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 101

Gln	His	Cys	Pro	Met	Tyr	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Phe
1				5					10					15

<210> 102

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 102

Gln	Val	Cys	His	Ile	Gln	Met	Tyr	Gly	Phe	Asp	Leu	Cys	Pro	His
1				5					10					15

<210> 103

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 103

Arg	Ala	Cys	Asp	Tyr	Trp	Met	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 104

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 104

Arg	Gln	Cys	His	Met	Gln	Met	Phe	Gly	Tyr	Asp	Leu	Cys	Pro	Phe
1				5				10					15	

<210> 105
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 105

Ser	Gly	Ser	Cys	Leu	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5				10					15	

<210> 106
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 106

Ser	Asn	Gly	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5				10					15	

<210> 107
 <211> 14
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 107

Ser	Thr	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His
1				5			10						

<210> 108
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
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<400> 108

Ser	Tyr	Lys	Cys	Leu	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 109
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 109

Val	Leu	Tyr	Cys	Thr	Tyr	Val	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 110
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 110

Val	Asn	Cys	Gly	Met	Trp	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Lys
1				5					10					15

<210> 111
<211> 15
<212> PRT
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<220>
<223> Masking Moiety

<400> 111

Tyr	Gly	Ser	Cys	Leu	Tyr	Ser	Phe	Tyr	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 112

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 112

Tyr	Pro	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10					15

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<211> 15

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<400> 113

Ala	Ala	Cys	Asp	Leu	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 114

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<212> PRT

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<400> 114

Ala	Phe	Cys	Thr	Leu	Ala	Pro	Tyr	Asn	Gln	Ala	Cys	Ile	Ala	Asn
1				5					10					15

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<211> 15

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<400> 115

Ala	Gly	Ser	Cys	Leu	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
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<210> 116

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<400> 116

Ala	Leu	Cys	Glu	Asn	Thr	Met	Tyr	Gly	Tyr	His	Leu	Cys	Pro	Trp
1				5				10					15	

<210> 117

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<400> 117

Ala	Leu	Ser	Cys	Met	Tyr	Ile	Met	Tyr	Gly	Val	Cys	Leu	Asp	Pro
1				5				10					15	

<210> 118

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<400> 118

Ala	Pro	Val	Cys	Asp	Val	Leu	Met	Phe	Gly	Phe	Cys	Met	Gln	Pro
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<400> 119

Ala	Gln	Val	Cys	Ser	Ile	Met	Met	Tyr	Gly	Thr	Cys	Leu	Met	Pro
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<400> 120

Ala	Ser	Thr	Cys	Met	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10				15	

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<400> 121

Ala	Val	Cys	Glu	Phe	Trp	Met	Phe	Gly	Phe	Asn	Leu	Cys	Pro	Tyr
1				5					10				15	

<210> 122
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<400> 122

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Asp	Ala	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Tyr	Asp	Tyr	His
1				5					10					15

<210> 123
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 <212> PRT
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<400> 123

Asp	Phe	Ser	Cys	Ile	Tyr	Ile	Met	Phe	Asp	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 124
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 <212> PRT
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<220>
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<400> 124

Asp	Phe	Ser	Cys	Met	Tyr	Val	Met	Tyr	Gly	Phe	Cys	Leu	Asp	Pro
1				5					10					15

<210> 125
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 <212> PRT
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<220>
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<400> 125

Asp	Phe	Thr	Cys	Met	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 126
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<400> 126

Asp	Phe	Thr	Cys	Thr	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 127

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

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<400> 127

Asp	His	Tyr	Cys	Thr	Tyr	Ile	Met	Tyr	Ser	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 128

<211> 15

<212> PRT

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<400> 128

Asp	Ile	Cys	Thr	Asn	Phe	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 129

<211> 15

<212> PRT

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<400> 129

Asp	Ile	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5					10					15

<210> 130

<211> 15

<212> PRT
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<400> 130

Asp	Lys	Asn	Thr	Cys	Pro	Leu	His	Pro	Met	Cys	His	Asp	Tyr	Arg
1				5					10					15

<210> 131
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 <212> PRT
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<400> 131

Asp	Met	Asn	Met	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Trp	His
1				5					10					15

<210> 132
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 <212> PRT
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<220>
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<400> 132

Asp	Met	Asn	Ser	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5					10					15

<210> 133
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<400> 133

Asp	Met	Asn	Ser	Cys	Pro	Asn	His	Pro	Met	Cys	Tyr	Asp	Tyr	Arg
1				5					10					15

<210> 134
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<400> 134

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Phe	Asp	Tyr	Arg
1				5					10					15

<210> 135
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<400> 135

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Phe	Gln
1				5					10					15

<210> 136
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<400> 136

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Tyr	Arg
1				5					10					15

<210> 137
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<400> 137

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Tyr	Asp	Tyr	His
1				5					10					15

<210> 138

<211> 15

<212> PRT

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<400> 138

Asp	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Tyr	Asp	Tyr	Lys
1				5					10					15

<210> 139

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

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<400> 139

Asp	Met	Ser	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Tyr	Met
1				5					10					15

<210> 140

<211> 15

<212> PRT

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<400> 140

Asp	Arg	Asn	Met	Cys	Pro	Tyr	His	Pro	Met	Cys	Tyr	Asp	Tyr	Arg
1				5					10					15

<210> 141

<211> 15

<212> PRT

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<400> 141

Asp	Ser	Cys	Ala	Phe	Met	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 142

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 142

Asp	Ser	Cys	Arg	Ser	Val	Phe	Asp	Met	Val	Trp	Asn	Cys	Trp	Asn
1				5					10					15

<210> 143

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 143

Asp	Thr	Pro	Asn	Cys	Pro	His	His	Pro	Met	Cys	His	Asn	His	Met
1				5					10					15

<210> 144

<211> 15

<212> PRT

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<220>

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<400> 144

Asp	Val	Ser	Cys	Leu	Tyr	Val	Met	Tyr	Ser	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 145
 <211> 15
 <212> PRT
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<400> 145

Asp	Trp	Cys	Ala	Ser	Met	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 146
 <211> 15
 <212> PRT
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<220>
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<400> 146

Glu	Phe	Ser	Cys	Met	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5				10					15	

<210> 147
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 147

Glu	His	Cys	Asp	Val	Trp	Met	Phe	Gly	Phe	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 148
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 148

Glu Pro Cys Asp Tyr Trp Met Phe Gly Val Asn Leu Cys Pro Tyr
 1 5 10 15

<210> 149
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 149

Glu Gln Cys Thr Met Trp Met Tyr Gly Phe Asn Leu Cys Pro Tyr
 1 5 10 15

<210> 150
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 150

Glu Ser Ala Cys Ser Leu Arg Met Tyr Glu Val Cys Leu Gln Pro
 1 5 10 15

<210> 151
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 151

Glu Ser Cys Ala Ser Met Tyr Gly Tyr Ser Met Cys Pro Arg Thr
 1 5 10 15

<210> 152
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 152

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

<210> 153

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 153

Phe	Ser	Asn	Thr	Cys	Pro	His	His	Pro	Met	Cys	Tyr	Asp	Tyr	Arg
1				5					10					15

<210> 154

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 154

Phe	Trp	Asn	Thr	Cys	Pro	His	His	Pro	Met	Cys	His	Asp	Tyr	Lys
1				5					10					15

<210> 155

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 155

Phe	Tyr	Gln	Asn	Cys	Tyr	Pro	Pro	Thr	Trp	Cys	Ser	Met	Phe	Ser
1				5					10					15

<210> 156

<211> 15

<212> PRT
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<220>
 <223> Masking Moiety

<400> 156

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

<210> 157
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 157

Gly	Gly	Ser	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10				15	

<210> 158
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 158

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

<210> 159
 <211> 15
 <212> PRT
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<220>
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<400> 159

Gly	His	Cys	Leu	Met	His	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Lys
1				5					10				15	

<210> 160
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
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<400> 160

Gly	His	Cys	Arg	Met	Ser	Met	Tyr	Glu	Met	Thr	Leu	Cys	Pro	Arg
1				5				10					15	

<210> 161
<211> 15
<212> PRT
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<220>
<223> Masking Moiety

<400> 161

Gly	Ile	Ser	Cys	Val	His	Ile	Met	Phe	Asn	Phe	Cys	Leu	Asp	Pro
1				5				10					15	

<210> 162
<211> 15
<212> PRT
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<220>
<223> Masking Moiety

<400> 162

Gly	Leu	Cys	Val	Met	Tyr	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 163
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 163

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

<210> 164

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 164

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

<210> 165

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 165

Gly	Thr	Lys	Cys	Ile	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 166

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 166

Gly	Thr	Ser	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Arg
1				5					10					15

<210> 167

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 167

Gly	Thr	Thr	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 168

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 168

Gly	Val	Cys	His	Phe	Phe	Met	Tyr	Gly	Val	Ser	Met	Cys	Pro	Ala
1				5					10					15

<210> 169

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 169

Gly	Val	Pro	Cys	Trp	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 170

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 170

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

<210> 171
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 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 171

His	Ala	Lys	Cys	Val	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 172
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 172

His	Asp	Ser	Cys	Met	Tyr	Ser	Met	Tyr	Asn	Phe	Cys	Leu	Asp	Pro
1				5					10					15

<210> 173
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 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 173

His	Gly	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Tyr	Gln
1				5					10					15

<210> 174
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 <212> PRT
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<220>
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<400> 174

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His	Lys	Gly	Cys	Leu	Tyr	Ser	Phe	Tyr	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 175
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 175

His	Lys	Gly	Cys	Leu	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 176
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 176

His	Leu	Ser	Cys	Met	Tyr	Ile	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 177
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 177

His	Ser	Ser	Cys	Ile	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 178
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 <212> PRT
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<220>

<223> Masking Moiety

<400> 178

His	Thr	Asn	Met	Cys	Pro	Tyr	His	Pro	Met	Cys	Tyr	Asp	Tyr	Lys
1				5					10					15

<210> 179

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 179

His	Thr	Pro	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 180

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 180

Ile	Met	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Gln
1				5					10					15

<210> 181

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 181

Ile	Val	Pro	Cys	Thr	Tyr	Met	Met	Phe	Gly	Val	Cys	Leu	Gln	Pro
1				5					10					15

<210> 182

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
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<400> 182

Lys	Lys	Cys	Asp	Tyr	Trp	Phe	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 183
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 183

Lys	Asn	Thr	Cys	Val	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 184
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 184

Lys	Pro	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Pro
1				5					10					15

<210> 185
 <211> 15
 <212> PRT
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<220>
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<400> 185

Lys	Pro	Ser	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 186
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 186

Lys	Arg	Pro	Cys	Met	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 187
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 187

Lys	Thr	Ser	Cys	Met	Tyr	Ser	Phe	Tyr	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 188
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 188

Lys	Thr	Thr	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 189
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 189

Leu	Asp	Cys	Gln	Met	Tyr	Trp	Trp	Phe	Gly	Ala	Cys	Gly	Asp	Met
1				5					10					15

<210> 190

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 190

Leu	His	Cys	Ala	Ile	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Phe
1				5					10					15

<210> 191

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 191

Leu	His	Cys	Pro	Phe	Gln	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	His
1				5					10					15

<210> 192

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 192

Leu	His	Cys	Ser	Met	Tyr	Met	Tyr	Gly	Phe	Asn	Leu	Cys	Pro	Asn
1				5					10					15

<210> 193

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 193

Arg	Glu	Cys	Met	Ala	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 194

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 194

Arg	His	Cys	Gln	Met	His	Met	Phe	Gly	Tyr	Asp	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 195

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 195

Leu	Ile	His	Cys	Arg	Tyr	Val	Met	Tyr	Gly	Met	Cys	Leu	Glu	Pro
1				5					10					15

<210> 196

<211> 15

<212> PRT

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<223> Masking Moiety

<400> 196

Leu	Leu	Pro	Cys	Glu	Val	Met	Gly	Pro	Ser	Arg	Cys	Lys	His	Asp
1				5					10					15

<210> 197
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<400> 197

Leu	Pro	Cys	His	Ala	Tyr	Met	Tyr	Gly	Tyr	Ser	Leu	Cys	Pro	Tyr
1				5				10						15

<210> 198
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<400> 198

Leu	Pro	Cys	Leu	Ala	Tyr	Met	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Asn
1				5				10						15

<210> 199
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 <212> PRT
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<220>
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<400> 199

Leu	Pro	Cys	Met	Ala	Tyr	Met	Phe	Gly	Phe	Asn	Leu	Cys	Pro	His
1				5				10						15

<210> 200
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 <212> PRT
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<220>
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<400> 200

Leu Pro Cys Asn Phe His Met Phe Gly Phe Asn Leu Cys Pro Tyr
1 5 10 15

<210> 201
<211> 15
<212> PRT
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<220>
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<400> 201

Leu Gln Cys Ala Met Tyr Met Tyr Gly Tyr Asn Leu Cys Pro Tyr
1 5 10 15

<210> 202
<211> 15
<212> PRT
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<220>
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<400> 202

Leu Ser Ser Cys Thr Tyr Ser Phe Phe Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 203
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<400> 203

Leu Thr Cys Pro Phe Gln Met Tyr Gly Tyr Asn Leu Cys Pro Tyr
1 5 10 15

<210> 204
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<220>

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<400> 204

Leu	Thr	Ser	Gln	Cys	Ser	Pro	Trp	Tyr	Trp	Cys	Gln	Ile	Tyr	Asp
1				5					10					15

<210> 205

<211> 15

<212> PRT

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<400> 205

Leu	Tyr	Cys	Pro	Tyr	Met	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 206

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<212> PRT

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<400> 206

Leu	Tyr	His	Cys	Thr	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 207

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<212> PRT

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<400> 207

Leu	Tyr	Arg	Cys	Ile	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 208

<211> 15

<212> PRT
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<400> 208

Met	Gly	Cys	Ser	Met	Arg	Met	Trp	Gly	Met	Glu	Leu	Cys	Pro	Glu
1				5				10					15	

<210> 209
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<400> 209

Met	Lys	Cys	Asp	Tyr	Trp	Leu	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 210
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 <212> PRT
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<220>
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<400> 210

Met	Asn	His	Cys	Thr	Leu	His	Met	Tyr	Asn	Ile	Cys	Met	Asp	Pro
1				5				10					15	

<210> 211
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 <212> PRT
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<220>
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<400> 211

Met	Asn	Pro	Glu	Cys	Pro	His	His	Pro	Met	Cys	His	Asn	Ser	Asn
1				5				10					15	

<210> 212
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<212> PRT
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<220>
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<400> 212

Met	Pro	Ala	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 213
<211> 15
<212> PRT
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<220>
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<400> 213

Met	Pro	Gln	Cys	His	Val	Ile	Met	Tyr	Asn	Leu	Cys	Leu	Asp	Pro
1				5					10					15

<210> 214
<211> 15
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<220>
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<400> 214

Met	Ser	Thr	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 215
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<220>
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<400> 215

Met	Thr	Cys	Asn	Tyr	Trp	Phe	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 216

<211> 15

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<220>

<223> Masking Moiety

<400> 216

Met	Tyr	Cys	His	Gln	Ser	Met	Phe	Gly	Phe	Arg	Met	Cys	Pro	Asp
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<223> Masking Moiety

<400> 217

Asn	Ala	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5					10					15

<210> 218

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 218

Asn	Asp	Cys	Asp	Ile	Ser	Met	Phe	Asp	Gln	Ser	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 219

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 219

Asn	Phe	Ser	Cys	Val	Tyr	Val	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 220

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 220

Asn	Phe	Thr	Cys	Ala	Leu	Thr	Met	Tyr	Glu	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 221

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 221

Asn	Leu	Cys	His	Ala	Phe	Met	Phe	Gly	Phe	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 222

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 222

Asn	Leu	Asn	Asn	Cys	Pro	His	His	Pro	Met	Cys	His	Asp	Tyr	Gln
1				5					10					15

<210> 223
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 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 223

Asn	Pro	Pro	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

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

<220>
 <223> Masking Moiety

<400> 224

Asn	Ser	Ala	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 225
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 <212> PRT
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<220>
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<400> 225

Asn	Val	Cys	Thr	Val	Ser	Met	Phe	Gly	Val	Met	Leu	Cys	Pro	Ser
1				5					10					15

<210> 226
 <211> 15
 <212> PRT
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<220>
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<400> 226

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Pro Ala Cys Ala Thr Leu Met Tyr Ser Val Pro Leu Cys Pro Ala
1 5 10 15

<210> 227
<211> 15
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<213> Artificial Sequence

<220>
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<400> 227

Pro Ala Pro Cys Met Tyr Ser Phe Tyr Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 228
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
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<400> 228

Pro Leu Cys Ala Glu Met Tyr Gly Tyr Ser Met Cys Pro His Asn
1 5 10 15

<210> 229
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<213> Artificial Sequence

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<400> 229

Pro Gln Cys His Leu Tyr Met Tyr Gly Tyr Asn Leu Cys Pro Tyr
1 5 10 15

<210> 230
<211> 15
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<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 230

Pro	Arg	Pro	Cys	Met	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 231

<211> 15

<212> PRT

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<220>

<223> Masking Moiety

<400> 231

Gln	His	Cys	Pro	Phe	Gln	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 232

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 232

Gln	His	Cys	Gln	Met	His	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 233

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 233

Gln	His	Ser	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 234

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
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<400> 234

Gln	Lys	Cys	His	Ser	Tyr	Leu	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 235
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<400> 235

Gln	Lys	Cys	Asn	Met	Phe	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 236
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 <212> PRT
 <213> Artificial Sequence

<220>
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<400> 236

Gln	Met	Asn	Asp	Cys	Pro	Asn	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5				10					15	

<210> 237
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<220>
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<400> 237

Gln	Pro	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	Ala	Thr
1				5				10					15	

<210> 238
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<220>
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<400> 238

Gln	Pro	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	Arg	Thr
1				5					10					15

<210> 239
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<400> 239

Arg	Glu	Cys	His	Phe	Phe	Phe	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 240
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<220>
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<400> 240

Leu	Asn	Cys	Gly	Met	Phe	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 241
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<220>
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<400> 241

Arg	Leu	Cys	Thr	Ser	Tyr	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Gln
1				5					10					15

<210> 242

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 242

Arg	Leu	Ser	Cys	Met	Tyr	Ser	Met	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 243

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 243

Arg	Asn	Cys	Pro	Phe	Val	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 244

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 244

Arg	Asn	Gly	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 245

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 245

Arg	Asn	Gly	Cys	Val	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 246

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 246

Arg	Pro	Cys	His	Leu	Tyr	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Asp
1				5					10					15

<210> 247

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 247

Arg	Pro	Cys	His	Ser	Tyr	Met	Tyr	Gly	Ile	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 248

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 248

Arg	Ser	Cys	Asp	Met	Ile	Met	Phe	Gly	Phe	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 249
 <211> 15
 <212> PRT
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<220>
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<400> 249

Arg	Ser	Cys	Pro	Met	Trp	Phe	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 250
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<220>
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<400> 250

Arg	Ser	Thr	Val	Cys	Phe	Tyr	Asp	Phe	Cys	Gly	Pro	Trp	Glu	Arg
1				5					10					15

<210> 251
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<220>
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<400> 251

Arg	Thr	Cys	His	Phe	Tyr	Met	Tyr	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 252
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 <212> PRT
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<220>
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<400> 252

Arg	Thr	Cys	Ser	Met	Val	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 253
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 <212> PRT
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<220>
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<400> 253

Ser	Gly	Lys	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 254
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 <212> PRT
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<220>
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<400> 254

Ser	Ile	Val	Cys	Asp	Leu	Tyr	Trp	Glu	Ala	Thr	Cys	Leu	Arg	Pro
1				5					10					15

<210> 255
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<220>
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<400> 255

Ser	Leu	Ser	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 256
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>

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<400> 256

Ser	Met	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	Phe	Asp	Tyr	Lys
1				5					10					15

<210> 257

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 257

Ser	Gln	Cys	Trp	Met	Trp	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Lys
1				5					10					15

<210> 258

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 258

Ser	Ser	Ser	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 259

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 259

Ser	Thr	Ala	Cys	Thr	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 260

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
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<400> 260

Ser	Thr	Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Pro	His	Thr
1				5				10						15

<210> 261
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<220>
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<400> 261

Ser	Thr	Arg	Cys	Val	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5				10						15

<210> 262
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<400> 262

Thr	Ala	Cys	Gly	Ala	Trp	Met	Phe	Gly	Val	Asn	Leu	Cys	Pro	Tyr
1				5				10						15

<210> 263
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<220>
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<400> 263

Thr	Gly	Ala	Cys	Met	Tyr	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5				10						15

<210> 264
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<400> 264

Thr	Leu	Ser	Cys	Met	Tyr	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 265
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<400> 265

Thr	Ser	Cys	Thr	Val	Thr	Met	Tyr	Gln	Ile	Ser	Met	Cys	Pro	Tyr
1				5					10					15

<210> 266
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<400> 266

Val	Gly	Gly	Cys	Arg	His	Ser	Phe	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 267
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<400> 267

Val	His	Cys	Gln	Met	Tyr	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5				10					15	

<210> 268

<211> 15

<212> PRT

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<223> Masking Moiety

<400> 268

Val	His	Asn	Cys	Met	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 269

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 269

Val	Met	Cys	Lys	Leu	His	Met	Tyr	Gly	Ile	Pro	Val	Cys	Pro	Lys
1				5				10						15

<210> 270

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 270

Val	Asn	Phe	Cys	Asn	Tyr	Ser	Met	Tyr	Gly	Ile	Cys	Leu	Leu	Pro
1				5					10					15

<210> 271

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 271

Val	Asn	Phe	Cys	Tyr	Ala	Cys	Tyr	Cys	Met	Ser	Cys	Val	Phe	Ser
1				5					10					15

<210> 272

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 272

Val	Asn	Gln	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 273

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 273

Val	Pro	Cys	Pro	Phe	His	Met	Phe	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 274

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 274

Val	Arg	Cys	Gln	Met	Trp	Met	Tyr	Gly	Phe	Asn	Leu	Cys	Pro	His
1				5					10					15

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

<220>
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<400> 275

Val	Arg	Pro	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 276
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 <212> PRT
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<220>
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<400> 276

Val	Ser	Gly	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Ile	Cys	Leu	Asp	Pro
1				5					10					15

<210> 277
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<220>
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<400> 277

Tyr	Cys	Ser	Ser	Trp	Asp	Thr	Met	Thr	Ile	Pro	Ala	Cys	Asn	Asn
1				5					10					15

<210> 278
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<220>
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<400> 278

Tyr Asp Cys Asp Leu Ser Met Phe Gly Ile Glu Met Cys Pro Gln
1 5 10 15

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

<220>
<223> Masking Moiety

<400> 279

Tyr Gly Asn Thr Cys Pro Phe His Pro Met Cys His Asp Tyr Lys
1 5 10 15

<210> 280
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 280

Tyr Gly Tyr Cys Met Tyr Ser Phe Phe Asn Val Cys Leu Asp Pro
1 5 10 15

<210> 281
<211> 15
<212> PRT
<213> Artificial Sequence

<220>
<223> Masking Moiety

<400> 281

Tyr His Cys Thr Met His Met Phe Gly Tyr Asn Leu Cys Pro Phe
1 5 10 15

<210> 282
<211> 15
<212> PRT
<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 282

Tyr	Met	Asn	Thr	Cys	Pro	Asn	His	Pro	Met	Cys	Phe	Asp	Tyr	Gln
1				5					10					15

<210> 283

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 283

Tyr	Met	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Leu
1				5					10					15

<210> 284

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 284

Tyr	Met	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Arg
1				5					10					15

<210> 285

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety

<400> 285

Tyr	Asn	Asn	Cys	Thr	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 286

<211> 15

<212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 286

Tyr	Pro	Gly	Cys	Gln	Tyr	Ser	Phe	Phe	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 287
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 287

Tyr	Arg	Ser	Cys	Thr	His	Ile	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 288
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 288

Tyr	Ser	Phe	Cys	Asp	Met	Leu	Met	Tyr	Asp	Val	Cys	Leu	Val	Pro
1				5					10					15

<210> 289
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 289

Tyr	Ser	Ile	Asp	Cys	Gly	Leu	Ser	Trp	Trp	Cys	Gly	Gly	Met	Thr
1				5					10					15

<210> 290
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 290

Tyr	Ser	Thr	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5					10					15

<210> 291
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 291

Tyr	Val	Asn	Thr	Cys	Pro	His	His	Pro	Met	Cys	His	Asp	Tyr	His
1				5					10					15

<210> 292
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety

<400> 292

Tyr	Val	Asn	Thr	Cys	Pro	Tyr	His	Pro	Met	Cys	His	Asp	Tyr	Asn
1				5					10					15

<210> 293
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Masking Moiety - MY03

<400> 293

Met	Cys	Leu	Pro	Pro	Leu	Phe	Glu	Leu	Ala	Ser	Thr	Cys	Pro	Tyr
1				5					10					15

<210> 294

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - MY11

<400> 294

Leu	Pro	Asp	Cys	Gly	Met	Trp	Gly	Ile	Ser	Cys	Gly	Gly	Thr	Val
1				5					10					15

<210> 295

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - MY16

<400> 295

Arg	Asp	His	Thr	Cys	Asn	Pro	Arg	Asn	Cys	His	Pro	Asn	Met	Phe
1				5					10					15

<210> 296

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Masking Moiety - MY04

<400> 296

Trp	Arg	Cys	Met	Pro	Pro	Thr	Trp	Glu	Thr	Thr	Gln	Cys	His	Thr
1				5					10					15

<210> 297

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2001

<400> 297

Ile	Ser	Ser	Gly	Leu	Leu	Ser	Gly	Arg	Ser	Asp	Asn	His
1				5					10			

<210> 298

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2003

<400> 298

Ile	Ser	Ser	Gly	Leu	Leu	Ser	Gly	Arg	Ser	Ala	Asn	Pro	Arg	Gly
1				5					10					15

<210> 299

<211> 17

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2005

<400> 299

Ala	Val	Gly	Leu	Leu	Ala	Pro	Pro	Ser	Gly	Arg	Ser	Ala	Asn	Pro	Arg
1				5					10					15	

Gly

<210> 300

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2006

<400> 300

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asp His
1 5 10

<210> 301
<211> 13
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety - 2007

<400> 301

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Ile His
1 5 10

<210> 302
<211> 13
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety - 2008

<400> 302

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Gln His
1 5 10

<210> 303
<211> 13
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety - 2009

<400> 303

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Thr His
1 5 10

<210> 304
<211> 13
<212> PRT
<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2011

<400> 304

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro
1 5 10

<210> 305

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 2012

<400> 305

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro
1 5 10

<210> 306

<211> 18

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 3001

<400> 306

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp
1 5 10 15

Asn His

<210> 307

<211> 18

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety - 3006

<400> 307

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp
1 5 10 15

Asp His

<210> 308
 <211> 18
 <212> PRT
 <213> Artificial Sequence

 <220>
 <223> Cleavable Moiety - 3007

 <400> 308

Ala	Val	Gly	Leu	Leu	Ala	Pro	Pro	Gly	Gly	Leu	Ser	Gly	Arg	Ser	Asp
1				5					10					15	

Ile His

<210> 309
 <211> 18
 <212> PRT
 <213> Artificial Sequence

 <220>
 <223> Cleavable Moiety - 3008

 <400> 309

Ala	Val	Gly	Leu	Leu	Ala	Pro	Pro	Gly	Gly	Leu	Ser	Gly	Arg	Ser	Asp
1				5					10					15	

Gln His

<210> 310
 <211> 18
 <212> PRT
 <213> Artificial Sequence

 <220>
 <223> Cleavable Moiety - 3009

 <400> 310

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp
1 5 10 15

Thr His

<210> 311
<211> 18
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety - 3011

<400> 311

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp
1 5 10 15

Asn Pro

<210> 312
<211> 18
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety - 3012

<400> 312

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Ala
1 5 10 15

Asn Pro

<210> 313
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 313

Leu Ser Gly Arg Ser Asp Asn His
1 5

<210> 314

<211> 10

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 314

Leu Ser Gly Arg Ser Ala Asn Pro Arg Gly
1 5 10

<210> 315

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 315

Thr Gly Arg Gly Pro Ser Trp Val
1 5

<210> 316

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 316

Pro Leu Thr Gly Arg Ser Gly Gly
1 5

<210> 317

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 317

Thr Ala Arg Gly Pro Ser Phe Lys
 1 5

<210> 318

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 318

Asn Thr Leu Ser Gly Arg Ser Glu Asn His Ser Gly
 1 5 10

<210> 319

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 319

Asn Thr Leu Ser Gly Arg Ser Gly Asn His Gly Ser
 1 5 10

<210> 320

<211> 12

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 320

Thr Ser Thr Ser Gly Arg Ser Ala Asn Pro Arg Gly
 1 5 10

<210> 321
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 321

Thr Ser Gly Arg Ser Ala Asn Pro
1 5

<210> 322
<211> 10
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 322

Val His Met Pro Leu Gly Phe Leu Gly Pro
1 5 10

<210> 323
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 323

Ala Val Gly Leu Leu Ala Pro Pro
1 5

<210> 324
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 324

Ala Gln Asn Leu Leu Gly Met Val
1 5

<210> 325
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 325

Gln Asn Gln Ala Leu Arg Met Ala
1 5

<210> 326
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 326

Leu Ala Ala Pro Leu Gly Leu Leu
1 5

<210> 327
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Cleavable Moiety

<400> 327

Ser Thr Phe Pro Phe Gly Met Phe
1 5

<210> 328
<211> 8
<212> PRT
<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 328

Ile Ser Ser Gly Leu Leu Ser Ser
1 5

<210> 329

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 329

Pro Ala Gly Leu Trp Leu Asp Pro
1 5

<210> 330

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 330

Val Ala Gly Arg Ser Met Arg Pro
1 5

<210> 331

<211> 8

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 331

Val Val Pro Glu Gly Arg Arg Ser
1 5

<210> 332

<211> 8

<212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 332

Ile Leu Pro Arg Ser Pro Ala Phe
 1 5

<210> 333
 <211> 8
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 333

Met Val Leu Gly Arg Ser Leu Leu
 1 5

<210> 334
 <211> 8
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 334

Val Ala Gly Arg Ser Met Arg Pro
 1 5

<210> 335
 <211> 8
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 335

Gln Gly Arg Ala Ile Thr Phe Ile
 1 5

<210> 336
 <211> 8
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 336

Ser Pro Arg Ser Ile Met Leu Ala
 1 5

<210> 337
 <211> 8
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 337

Ser Met Leu Arg Ser Met Pro Leu
 1 5

<210> 338
 <211> 18
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Cleavable Moiety

<400> 338

Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp
 1 5 10 15

Tyr His

<210> 339
 <211> 18
 <212> PRT
 <213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 339

Ala	Val	Gly	Leu	Leu	Ala	Pro	Pro	Gly	Gly	Leu	Ser	Gly	Arg	Ser	Ala
1				5					10					15	

Asn Ile

<210> 340

<211> 18

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 340

Ala	Val	Gly	Leu	Leu	Ala	Pro	Pro	Gly	Gly	Leu	Ser	Gly	Arg	Ser	Asp
1				5					10					15	

Asn Ile

<210> 341

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 341

Ile	Ser	Ser	Gly	Leu	Leu	Ser	Gly	Arg	Ser	Asp	Tyr	His
1				5					10			

<210> 342

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 342

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Ile
1 5 10

<210> 343

<211> 13

<212> PRT

<213> Artificial Sequence

<220>

<223> Cleavable Moiety

<400> 343

Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Ile
1 5 10

<210> 344

<211> 108

<212> PRT

<213> Artificial Sequence

<220>

<223> Ipilimumab-VL

<400> 344

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

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

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

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

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

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 85 90 95

Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 100 105

<210> 345
 <211> 118
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Ipilimumab-VH

<400> 345

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

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

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

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

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

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

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

Leu Val Thr Val Ser Ser
 115

<210> 346
 <211> 107

<212> PRT

<213> Artificial Sequence

<220>

<223> Human Kappa constant LC

<400> 346

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
 1 5 10 15

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
 20 25 30

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
 35 40 45

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
 50 55 60

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
 65 70 75 80

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
 85 90 95

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 100 105

<210> 347

<211> 107

<212> PRT

<213> Artificial Sequence

<220>

<223> Mouse Kappa constant LC

<400> 347

Arg Ala Asp Ala Ala Pro Thr Val Ser Ile Phe Pro Pro Ser Ser Glu
 1 5 10 15

Gln Leu Thr Ser Gly Gly Ala Ser Val Val Cys Phe Leu Asn Asn Phe
 20 25 30

Tyr Pro Lys Asp Ile Asn Val Lys Trp Lys Ile Asp Gly Ser Glu Arg
35 40 45

Gln Asn Gly Val Leu Asn Ser Trp Thr Asp Gln Asp Ser Lys Asp Ser
50 55 60

Thr Tyr Ser Met Ser Ser Thr Leu Thr Leu Thr Lys Asp Glu Tyr Glu
65 70 75 80

Arg His Asn Ser Tyr Thr Cys Glu Ala Thr His Lys Thr Ser Thr Ser
85 90 95

Pro Ile Val Lys Ser Phe Asn Arg Asn Glu Cys
100 105

<210> 348
<211> 215
<212> PRT
<213> Artificial Sequence

<220>
<223> Ipilimumab-Human Kappa LC

<400> 348

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

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

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

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

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

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Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
85 90 95

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

Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
115 120 125

Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
130 135 140

Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
145 150 155 160

Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
165 170 175

Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
180 185 190

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

Ser Phe Asn Arg Gly Glu Cys
210 215

<210> 349
<211> 215
<212> PRT
<213> Artificial Sequence

<220>
<223> Ipilimumab-Mouse Kappa LC

<400> 349

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

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

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

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

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

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 85 90 95

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

Ala Pro Thr Val Ser Ile Phe Pro Pro Ser Ser Glu Gln Leu Thr Ser
 115 120 125

Gly Gly Ala Ser Val Val Cys Phe Leu Asn Asn Phe Tyr Pro Lys Asp
 130 135 140

Ile Asn Val Lys Trp Lys Ile Asp Gly Ser Glu Arg Gln Asn Gly Val
 145 150 155 160

Leu Asn Ser Trp Thr Asp Gln Asp Ser Lys Asp Ser Thr Tyr Ser Met
 165 170 175

Ser Ser Thr Leu Thr Leu Thr Lys Asp Glu Tyr Glu Arg His Asn Ser
 180 185 190

Tyr Thr Cys Glu Ala Thr His Lys Thr Ser Thr Ser Pro Ile Val Lys
 195 200 205

Ser Phe Asn Arg Asn Glu Cys
 210 215

<210> 350

<211> 329

<212> PRT

<213> Artificial Sequence

<220>

<223> Human IgG1 constant HC

<400> 350

Ala Ser Thr Lys Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys
 1 5 10 15

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

Phe Pro Glu Pro Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser
 35 40 45

Gly Val His Thr Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser
 50 55 60

Leu Ser Ser Val Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr
 65 70 75 80

Tyr Ile Cys Asn Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys
 85 90 95

Arg Val Glu Pro Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys
 100 105 110

Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro
 115 120 125

Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys
 130 135 140

Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp
 145 150 155 160

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

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Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu
180 185 190

His Gln Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn
195 200 205

Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly
210 215 220

Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Asp Glu
225 230 235 240

Leu Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr
245 250 255

Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn
260 265 270

Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe
275 280 285

Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn
290 295 300

Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr Thr
305 310 315 320

Gln Lys Ser Leu Ser Leu Ser Pro Gly
325

<210> 351
<211> 329
<212> PRT
<213> Artificial Sequence

<220>
<223> Mouse IgG1 constant HC

<400> 351

Ala Ser Thr Lys Gly Pro Ser Val Phe Pro Leu Ala Pro Ser Ser Lys
1 5 10 15

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

Phe Pro Glu Pro Val Thr Val Ser Trp Asn Ser Gly Ala Leu Thr Ser
 35 40 45

Gly Val His Thr Phe Pro Ala Val Leu Gln Ser Ser Gly Leu Tyr Ser
 50 55 60

Leu Ser Ser Val Val Thr Val Pro Ser Ser Ser Leu Gly Thr Gln Thr
 65 70 75 80

Tyr Ile Cys Asn Val Asn His Lys Pro Ser Asn Thr Lys Val Asp Lys
 85 90 95

Lys Val Glu Pro Lys Ser Cys Asp Lys Thr His Thr Cys Pro Pro Cys
 100 105 110

Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser Val Phe Leu Phe Pro Pro
 115 120 125

Lys Pro Lys Asp Thr Leu Met Ile Ser Arg Thr Pro Glu Val Thr Cys
 130 135 140

Val Val Val Asp Val Ser His Glu Asp Pro Glu Val Lys Phe Asn Trp
 145 150 155 160

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

Glu Gln Tyr Ala Ser Thr Tyr Arg Val Val Ser Val Leu Thr Val Leu
 180 185 190

His Gln Asp Trp Leu Asn Gly Lys Glu Tyr Lys Cys Lys Val Ser Asn
 195 200 205

Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr Ile Ser Lys Ala Lys Gly
 210 215 220

Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu Pro Pro Ser Arg Glu Glu
225 230 235 240

Met Thr Lys Asn Gln Val Ser Leu Thr Cys Leu Val Lys Gly Phe Tyr
245 250 255

Pro Ser Asp Ile Ala Val Glu Trp Glu Ser Asn Gly Gln Pro Glu Asn
260 265 270

Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp Ser Asp Gly Ser Phe Phe
275 280 285

Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser Arg Trp Gln Gln Gly Asn
290 295 300

Val Phe Ser Cys Ser Val Met His Glu Ala Leu His Asn His Tyr Thr
305 310 315 320

Gln Lys Ser Leu Ser Leu Ser Pro Gly
325

<210> 352
<211> 330
<212> PRT
<213> Artificial Sequence

<220>
<223> Mouse IgG2a constant HC

<400> 352

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

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

Phe Pro Glu Pro Val Thr Leu Thr Trp Asn Ser Gly Ser Leu Ser Ser
35 40 45

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

Ser Ser Ser Val Thr Val Thr Ser Ser Thr Trp Pro Ser Gln Ser Ile
 65 70 75 80

Thr Cys Asn Val Ala His Pro Ala Ser Ser Thr Lys Val Asp Lys Lys
 85 90 95

Ile Glu Pro Arg Gly Pro Thr Ile Lys Pro Cys Pro Pro Cys Lys Cys
 100 105 110

Pro Ala Pro Asn Leu Leu Gly Gly Pro Ser Val Phe Ile Phe Pro Pro
 115 120 125

Lys Ile Lys Asp Val Leu Met Ile Ser Leu Ser Pro Ile Val Thr Cys
 130 135 140

Val Val Val Asp Val Ser Glu Asp Asp Pro Asp Val Gln Ile Ser Trp
 145 150 155 160

Phe Val Asn Asn Val Glu Val His Thr Ala Gln Thr Gln Thr His Arg
 165 170 175

Glu Asp Tyr Asn Ser Thr Leu Arg Val Val Ser Ala Leu Pro Ile Gln
 180 185 190

His Gln Asp Trp Met Ser Gly Lys Glu Phe Lys Cys Lys Val Asn Asn
 195 200 205

Lys Asp Leu Pro Ala Pro Ile Glu Arg Thr Ile Ser Lys Pro Lys Gly
 210 215 220

Ser Val Arg Ala Pro Gln Val Tyr Val Leu Pro Pro Pro Glu Glu Glu
 225 230 235 240

Met Thr Lys Lys Gln Val Thr Leu Thr Cys Met Val Thr Asp Phe Met
 245 250 255

3338059PC02_ST25.txt

Pro Glu Asp Ile Tyr Val Glu Trp Thr Asn Asn Gly Lys Thr Glu Leu
260 265 270

Asn Tyr Lys Asn Thr Glu Pro Val Leu Asp Ser Asp Gly Ser Tyr Phe
275 280 285

Met Tyr Ser Lys Leu Arg Val Glu Lys Lys Asn Trp Val Glu Arg Asn
290 295 300

Ser Tyr Ser Cys Ser Val Val His Glu Gly Leu His Asn His His Thr
305 310 315 320

Thr Lys Ser Phe Ser Arg Thr Pro Gly Lys
325 330

<210> 353
<211> 447
<212> PRT
<213> Artificial Sequence

<220>
<223> Ipilimumab-VH-Human IgG1 constant HC

<400> 353

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

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

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

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

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

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

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

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

Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala Leu Gly
 130 135 140

Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Trp Asn
 145 150 155 160

Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val Leu Gln
 165 170 175

Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro Ser Ser
 180 185 190

Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys Pro Ser
 195 200 205

Asn Thr Lys Val Asp Lys Arg Val Glu Pro Lys Ser Cys Asp Lys Thr
 210 215 220

His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser
 225 230 235 240

Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg
 245 250 255

Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu Asp Pro
 260 265 270

Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His Asn Ala
 275 280 285

Lys Thr Lys Pro Arg Glu Glu Gln Tyr Asn Ser Thr Tyr Arg Val Val
 290 295 300

Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu Tyr
 305 310 315 320

Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr
 325 330 335

Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu
 340 345 350

Pro Pro Ser Arg Asp Glu Leu Thr Lys Asn Gln Val Ser Leu Thr Cys
 355 360 365

Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser
 370 375 380

Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp
 385 390 395 400

Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser
 405 410 415

Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala
 420 425 430

Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly
 435 440 445

<210> 354

<211> 447

<212> PRT

<213> Artificial Sequence

<220>

<223> Ipilimumab-VH-Mouse IgG1 constant HC

<400> 354

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

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

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

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

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

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

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

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

Leu Ala Pro Ser Ser Lys Ser Thr Ser Gly Gly Thr Ala Ala Leu Gly
 130 135 140

Cys Leu Val Lys Asp Tyr Phe Pro Glu Pro Val Thr Val Ser Trp Asn
 145 150 155 160

Ser Gly Ala Leu Thr Ser Gly Val His Thr Phe Pro Ala Val Leu Gln
 165 170 175

Ser Ser Gly Leu Tyr Ser Leu Ser Ser Val Val Thr Val Pro Ser Ser
 180 185 190

Ser Leu Gly Thr Gln Thr Tyr Ile Cys Asn Val Asn His Lys Pro Ser
 195 200 205

Asn Thr Lys Val Asp Lys Lys Val Glu Pro Lys Ser Cys Asp Lys Thr
 210 215 220

3338059PC02_ST25.txt

His Thr Cys Pro Pro Cys Pro Ala Pro Glu Leu Leu Gly Gly Pro Ser
 225 230 235 240

Val Phe Leu Phe Pro Pro Lys Pro Lys Asp Thr Leu Met Ile Ser Arg
 245 250 255

Thr Pro Glu Val Thr Cys Val Val Val Asp Val Ser His Glu Asp Pro
 260 265 270

Glu Val Lys Phe Asn Trp Tyr Val Asp Gly Val Glu Val His Asn Ala
 275 280 285

Lys Thr Lys Pro Arg Glu Glu Gln Tyr Ala Ser Thr Tyr Arg Val Val
 290 295 300

Ser Val Leu Thr Val Leu His Gln Asp Trp Leu Asn Gly Lys Glu Tyr
 305 310 315 320

Lys Cys Lys Val Ser Asn Lys Ala Leu Pro Ala Pro Ile Glu Lys Thr
 325 330 335

Ile Ser Lys Ala Lys Gly Gln Pro Arg Glu Pro Gln Val Tyr Thr Leu
 340 345 350

Pro Pro Ser Arg Glu Glu Met Thr Lys Asn Gln Val Ser Leu Thr Cys
 355 360 365

Leu Val Lys Gly Phe Tyr Pro Ser Asp Ile Ala Val Glu Trp Glu Ser
 370 375 380

Asn Gly Gln Pro Glu Asn Asn Tyr Lys Thr Thr Pro Pro Val Leu Asp
 385 390 395 400

Ser Asp Gly Ser Phe Phe Leu Tyr Ser Lys Leu Thr Val Asp Lys Ser
 405 410 415

Arg Trp Gln Gln Gly Asn Val Phe Ser Cys Ser Val Met His Glu Ala
 420 425 430

3338059PC02_ST25.txt

Leu His Asn His Tyr Thr Gln Lys Ser Leu Ser Leu Ser Pro Gly
 435 440 445

<210> 355

<211> 448

<212> PRT

<213> Artificial Sequence

<220>

<223> Ipilimumab-VH-Mouse IgG2a constant HC

<400> 355

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

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

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

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

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

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

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

Leu Val Thr Val Ser Ser Ala Lys Thr Thr Ala Pro Ser Val Tyr Pro
 115 120 125

Leu Ala Pro Val Cys Gly Asp Thr Thr Gly Ser Ser Val Thr Leu Gly
 130 135 140

Cys Leu Val Lys Gly Tyr Phe Pro Glu Pro Val Thr Leu Thr Trp Asn
 145 150 155 160

Ser Gly Ser Leu Ser Ser Gly Val His Thr Phe Pro Ala Val Leu Gln
 165 170 175

Ser Asp Leu Tyr Thr Leu Ser Ser Ser Val Thr Val Thr Ser Ser Thr
 180 185 190

Trp Pro Ser Gln Ser Ile Thr Cys Asn Val Ala His Pro Ala Ser Ser
 195 200 205

Thr Lys Val Asp Lys Lys Ile Glu Pro Arg Gly Pro Thr Ile Lys Pro
 210 215 220

Cys Pro Pro Cys Lys Cys Pro Ala Pro Asn Leu Leu Gly Gly Pro Ser
 225 230 235 240

Val Phe Ile Phe Pro Pro Lys Ile Lys Asp Val Leu Met Ile Ser Leu
 245 250 255

Ser Pro Ile Val Thr Cys Val Val Val Asp Val Ser Glu Asp Asp Pro
 260 265 270

Asp Val Gln Ile Ser Trp Phe Val Asn Asn Val Glu Val His Thr Ala
 275 280 285

Gln Thr Gln Thr His Arg Glu Asp Tyr Asn Ser Thr Leu Arg Val Val
 290 295 300

Ser Ala Leu Pro Ile Gln His Gln Asp Trp Met Ser Gly Lys Glu Phe
 305 310 315 320

Lys Cys Lys Val Asn Asn Lys Asp Leu Pro Ala Pro Ile Glu Arg Thr
 325 330 335

Ile Ser Lys Pro Lys Gly Ser Val Arg Ala Pro Gln Val Tyr Val Leu
 340 345 350

Pro Pro Pro Glu Glu Glu Met Thr Lys Lys Gln Val Thr Leu Thr Cys
 355 360 365

Met Val Thr Asp Phe Met Pro Glu Asp Ile Tyr Val Glu Trp Thr Asn
 370 375 380

Asn Gly Lys Thr Glu Leu Asn Tyr Lys Asn Thr Glu Pro Val Leu Asp
 385 390 395 400

Ser Asp Gly Ser Tyr Phe Met Tyr Ser Lys Leu Arg Val Glu Lys Lys
 405 410 415

Asn Trp Val Glu Arg Asn Ser Tyr Ser Cys Ser Val Val His Glu Gly
 420 425 430

Leu His Asn His His Thr Thr Lys Ser Phe Ser Arg Thr Pro Gly Lys
 435 440 445

<210> 356
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV01-2001 LC

<400> 356

Gln Gly Gln Ser Gly Gln Gly Asp Phe Ser Cys Leu His Ser Met Tyr
 1 5 10 15

Asn Val Cys Leu Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

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Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 357
<211> 255
<212> PRT
<213> Artificial Sequence

<220>

<223> YV01-2001 LC

<400> 357

Asp Phe Ser Cys Leu His Ser Met Tyr Asn Val Cys Leu Asp Pro Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 358
<211> 262
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV02-2001 LC

<400> 358

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Tyr Gly Tyr
1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 359
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV02-2001 LC

<400> 359

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 360
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV03-2001 LC

<400> 360

Gln Gly Gln Ser Gly Gln Gly Leu His Cys Arg Thr Gln Met Tyr Gly
 1 5 10 15

Tyr Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

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Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 361
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV03-2001 LC

<400> 361

Leu His Cys Arg Thr Gln Met Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 362
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV04-2001 LC

<400> 362

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

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Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 363
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2001 LC

<400> 363

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210>	364
<211>	256
<212>	PRT
<213>	Artificial Sequence

<220>
<223> Spacer - YV06-2001 LC

$\langle 400 \rangle$ 364

Gln Gly Gln Ser Gly Ser Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys
1 5 10 15

Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
130 135 140

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Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 365
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV06-2001 LC

<400> 365

Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Gly Gly Gly Ser Ser Gly
1 5 10 15

Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly
20 25 30

Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu
35 40 45

Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val
50 55 60

Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro
65 70 75 80

Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp
85 90 95

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

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 366

<211> 263

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV09-2001 LC

<400> 366

Gln Gly Gln Ser Gly Ser Phe Gly Tyr Val Thr Ala Cys Pro Asn His
 1 5 10 15

Pro Met Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile
 20 25 30

Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser
 35 40 45

Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
 50 55 60

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser
 65 70 75 80

Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
 85 90 95

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu
 115 120 125

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 130 135 140

Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
 145 150 155 160

Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
 165 170 175

3338059PC02_ST25.txt

Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
180 185 190

Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
195 200 205

Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
210 215 220

Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
225 230 235 240

Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
245 250 255

Ser Phe Asn Arg Gly Glu Cys
260

<210> 367
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV09-2001 LC

<400> 367

Phe Gly Thr Ala Cys Pro Asn His Pro Met Cys His Asp Trp Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 368
<211> 258
<212> PRT
<213> Artificial Sequence

<220>

<223> Spacer - YV23-2001 LC

<400> 368

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

3338059PC02_ST25.txt

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 369
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2001 LC

<400> 369

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 370
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV24-2001 LC

<400> 370

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

3338059PC02_ST25.txt

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
 210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
 225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
 245 250 255

Glu Cys

<210> 371
 <211> 252
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV24-2001 LC

<400> 371

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
 1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
 20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
 35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
 50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
 85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 372
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV27-2001 LC

<400> 372

Gln Gly Gln Ser Gly Ser Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser
 1 5 10 15

3338059PC02_ST25.txt

Met Cys Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
 210 215 220

3338059PC02_ST25.txt

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 373
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV27-2001 LC

<400> 373

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 374
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV29-2001 LC

<400> 374

Gln Gly Gln Ser Gly Ser Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys
1 5 10 15

Ala His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu
20 25 30

3338059PC02_ST25.txt

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
 225 230 235 240

3338059PC02_ST25.txt

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
 245 250 255

Gly Glu Cys

<210> 375
 <211> 253
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV29-2001 LC

<400> 375

Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Ala His Thr Gly Gly Gly
 1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
 20 25 30

Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 376
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV32-2001 LC

<400> 376

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Leu Cys His Asp Trp
1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

3338059PC02_ST25.txt

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
245 250 255

Cys

<210> 377
 <211> 251
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV32-2001 LC

<400> 377

Cys Pro Asn His Pro Leu Cys His Asp Trp Gln Gly Gly Gly Ser Ser
 1 5 10 15

Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His
 20 25 30

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 35 40 45

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
 50 55 60

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 65 70 75 80

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 85 90 95

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
 100 105 110

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
 115 120 125

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 130 135 140

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
 145 150 155 160

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
165 170 175

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
180 185 190

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
195 200 205

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
210 215 220

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
225 230 235 240

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 378
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV33-2001 LC

<400> 378

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Met Cys Ala Asp Trp
1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

3338059PC02_ST25.txt

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
245 250 255

Cys

<210> 379
 <211> 251
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV33-2001 LC

<400> 379

Cys Pro Asn His Pro Met Cys Ala Asp Trp Gln Gly Gly Gly Ser Ser
 1 5 10 15

Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His
 20 25 30

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 35 40 45

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
 50 55 60

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 65 70 75 80

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 85 90 95

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
 100 105 110

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
 115 120 125

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 130 135 140

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
 145 150 155 160

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
 165 170 175

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
180 185 190

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
195 200 205

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
210 215 220

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
225 230 235 240

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 380
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV35-2001 LC

<400> 380

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Met Cys His Asp Ala
1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

3338059PC02_ST25.txt

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
245 250 255

Cys

<210> 381
<211> 251
<212> PRT
<213> Artificial Sequence

<220>

<223> YV35-2001 LC

<400> 381

Cys Pro Asn His Pro Met Cys His Asp Ala Gln Gly Gly Gly Ser Ser
 1 5 10 15

Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His
 20 25 30

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 35 40 45

Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser
 50 55 60

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 65 70 75 80

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 85 90 95

Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
 100 105 110

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
 115 120 125

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
 130 135 140

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
 145 150 155 160

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
 165 170 175

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
 180 185 190

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
195 200 205

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
210 215 220

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
225 230 235 240

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 382
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV39-2001 LC

<400> 382

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
1 5 10 15

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

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

3338059PC02_ST25.txt

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 383
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2001 LC

<400> 383

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
 1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
 20 25 30

Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 384
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV41-2001 LC

<400> 384

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

Cys Pro Ala Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu
 20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

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Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 385
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV41-2001 LC

<400> 385

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Ala Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
 20 25 30

Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 386
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV51-2001 LC

<400> 386

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Ile
 1 5 10 15

Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

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Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 387
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV51-2001 LC

<400> 387

Phe Gly Ala Ala Cys Pro Asn His Pro Ile Cys His Asp Trp Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 388
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV52-2001 LC

<400> 388

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Leu
1 5 10 15

Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

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Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 389
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV52-2001 LC

<400> 389

Phe Gly Ala Ala Cys Pro Asn His Pro Leu Cys His Asp Trp Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 390
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV53-2001 LC

<400> 390

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Met
 1 5 10 15

Cys His Asp Ala Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

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Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 391
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV53-2001 LC

<400> 391

Phe Gly Ala Ala Cys Pro Asn His Pro Met Cys His Asp Ala Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 392

<211> 258

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV54-2001 LC

<400> 392

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Ala Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

3338059PC02_ST25.txt

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 393
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV54-2001 LC

<400> 393

Cys Leu His Ser Ala Tyr Asn Ala Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
 85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 394
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>

<223> Spacer - YV55-2001 LC

<400> 394

Gln Gly Gln Ser Gly Ser Cys Ala His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

3338059PC02_ST25.txt

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 395
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV55-2001 LC

<400> 395

Cys Ala His Ser Ala Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 396
<211> 258
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV56-2001 LC

<400> 396

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Ala
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

3338059PC02_ST25.txt

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 397
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV56-2001 LC

<400> 397

Cys Leu His Ser Ala Tyr Asn Val Cys Ala Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 398
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV57-2001 LC

<400> 398

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

3338059PC02_ST25.txt

Ala Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
 210 215 220

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Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 399
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV57-2001 LC

<400> 399

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Ala Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 400
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV58-2001 LC

<400> 400

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

Asp Ala Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

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Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
 210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
 225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
 245 250 255

Glu Cys

<210> 401
 <211> 252
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV58-2001 LC

<400> 401

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Asp Ala Gly Gly Gly Ser
 1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
 20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
 35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
 50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
 85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 402
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV61-2001 LC

<400> 402

Gln Gly Gln Ser Gly Ser Tyr Ile Ser Asp Cys Pro Tyr His Pro Met
1 5 10 15

Cys His Asp Tyr Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
35 40 45

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Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
 245 250 255

Asn Arg Gly Glu Cys
260

<210> 403
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV61-2001 LC

<400> 403

Tyr Ile Ser Asp Cys Pro Tyr His Pro Met Cys His Asp Tyr Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 404
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV62-2001 LC

<400> 404

Gln Gly Gln Ser Gly Ser Phe Arg Asn Thr Cys Pro Tyr His Pro Met
 1 5 10 15

Cys His Asp Tyr Arg Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

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Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 405
 <211> 255
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV62-2001 LC

<400> 405

Phe Arg Asn Thr Cys Pro Tyr His Pro Met Cys His Asp Tyr Arg Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 406
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV63-2001 LC

<400> 406

Gln Gly Gln Ser Gly Ser Arg Glu Cys His Met Trp Met Phe Gly Val
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

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Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 407
<211> 255
<212> PRT
<213> Artificial Sequence

<220>

<223> YV63-2001 LC

<400> 407

Arg Glu Cys His Met Trp Met Phe Gly Val Asn Leu Cys Pro Tyr Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 408
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV64-2001 LC

<400> 408

Gln Gly Gln Ser Gly Ser Ala Val Cys His Met Tyr Met Tyr Gly Tyr
1 5 10 15

Asn Leu Cys Pro Phe Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

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Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
 245 250 255

Asn Arg Gly Glu Cys
 260

<210> 409
 <211> 255
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV64-2001 LC

<400> 409

Ala Val Cys His Met Tyr Met Tyr Gly Tyr Asn Leu Cys Pro Phe Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 410
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV65-2001 LC

<400> 410

Gln Gly Gln Ser Gly Gln Gly Arg Ser Cys Pro Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

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Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 411
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV65-2001 LC

<400> 411

Arg Ser Cys Pro Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 412
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV66-2001 LC

<400> 412

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Phe Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

3338059PC02_ST25.txt

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 413
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV66-2001 LC

<400> 413

Gln Pro Cys Ala Gln Met Phe Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 414
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV04-2006 LC

<400> 414

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asp His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

3338059PC02_ST25.txt

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys Cys
260

<210> 415
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2006 LC

<400> 415

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asp His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 416
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV23-2006 LC

<400> 416

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asp His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

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Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 417
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2006 LC

<400> 417

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asp
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 418

<211> 259

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV39-2006 LC

<400> 418

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

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

Leu Ser Gly Arg Ser Asp Asp His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

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Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 419
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2006 LC

<400> 419

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Asp His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 420
<211> 261
<212> PRT
<213> Artificial Sequence

<220>

<223> Spacer - YV04-2007 LC

<400> 420

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Ile His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

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Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 421
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2007 LC

<400> 421

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Ile His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 422
<211> 258
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV23-2007 LC

<400> 422

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Ile His Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

3338059PC02_ST25.txt

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 423
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2007 LC

<400> 423

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Ile
20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 424
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV39-2007 LC

<400> 424

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

3338059PC02_ST25.txt

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

Leu Ser Gly Arg Ser Asp Ile His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

3338059PC02_ST25.txt

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 425
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2007 LC

<400> 425

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Ile His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 426
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV04-2008 LC

<400> 426

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

3338059PC02_ST25.txt

Gly Leu Leu Ser Gly Arg Ser Asp Gln His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

3338059PC02_ST25.txt

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 427
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2008 LC

<400> 427

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Gln His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 428
<211> 258
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV23-2008 LC

<400> 428

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
20 25 30

Ser Gly Arg Ser Asp Gln His Gly Gly Gly Ser Glu Ile Val Leu Thr
35 40 45

3338059PC02_ST25.txt

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
 210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
 225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
 245 250 255

Glu Cys

<210> 429
 <211> 252
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV23-2008 LC

<400> 429

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
 1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Gln
 20 25 30

His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
 35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
 50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
 65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
 85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
 100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 430
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV39-2008 LC

<400> 430

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
1 5 10 15

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

Leu Ser Gly Arg Ser Asp Gln His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

3338059PC02_ST25.txt

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
 225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
 245 250 255

Gly Glu Cys

<210> 431
 <211> 253
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV39-2008 LC

<400> 431

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
 1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
 20 25 30

Gln His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 432
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV04-2009 LC

<400> 432

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
1 5 10 15

Asn Leu Cys Pro Tyr Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu
20 25 30

Leu Ser Gly Arg Ser Asp Thr His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
65 70 75 80

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Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 433
<211> 253
<212> PRT
<213> Artificial Sequence

<220>

<223> YV04-2009 LC

<400> 433

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
 1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
 20 25 30

Thr His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 434
 <211> 256
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV23-2009 LC

<400> 434

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly
 20 25 30

Arg Ser Asp Thr His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
 35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
 50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
 65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
 85 90 95

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Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 435
<211> 250
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2009 LC

<400> 435

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Ser Ser Gly
1 5 10 15

Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Thr His Gly
 20 25 30

Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu
 35 40 45

Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val
 50 55 60

Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro
 65 70 75 80

Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp
 85 90 95

Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser
 100 105 110

Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly
 115 120 125

Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg
 130 135 140

Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln
 145 150 155 160

Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr
 165 170 175

Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser
 180 185 190

Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr
 195 200 205

Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys
 210 215 220

His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro
 225 230 235 240

Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 436
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV39-2009 LC

<400> 436

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

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

Leu Ser Gly Arg Ser Asp Thr His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

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Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 437
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2009 LC

<400> 437

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Thr His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 438
 <211> 263
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV01-2011 LC

<400> 438

Gln Gly Gln Ser Gly Gln Gly Asp Phe Ser Cys Leu His Ser Met Tyr
 1 5 10 15

Asn Val Cys Leu Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

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Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys Cys
260

<210> 439
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV01-2011 LC

<400> 439

Asp Phe Ser Cys Leu His Ser Met Tyr Asn Val Cys Leu Asp Pro Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 440
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV02-2011 LC

<400> 440

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

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Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 441
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV02-2011 LC

<400> 441

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 442

<211> 262

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV03-2011 LC

<400> 442

Gln Gly Gln Ser Gly Gln Gly Leu His Cys Arg Thr Gln Met Tyr Gly
 1 5 10 15

Tyr Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
 165 170 175

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Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 443
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV03-2011 LC

<400> 443

Leu His Cys Arg Thr Gln Met Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 444
<211> 261
<212> PRT
<213> Artificial Sequence

<220>

<223> Spacer - YV04-2011 LC

<400> 444

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

3338059PC02_ST25.txt

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 445
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2011 LC

<400> 445

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 446
<211> 258
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV23-2011 LC

<400> 446

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
 195 200 205

3338059PC02_ST25.txt

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 447
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2011 LC

<400> 447

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn
20 25 30

Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
 115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
 130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
 145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
 165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
 180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
 195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
 210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
 225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 448
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV39-2011 LC

<400> 448

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

3338059PC02_ST25.txt

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

Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

3338059PC02_ST25.txt

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 449
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2011 LC

<400> 449

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 450
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV61-2011 LC

<400> 450

Gln Gly Gln Ser Gly Ser Tyr Ile Ser Asp Cys Pro Tyr His Pro Met
 1 5 10 15

Cys His Asp Tyr Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

3338059PC02_ST25.txt

Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

3338059PC02_ST25.txt

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 451
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV61-2011 LC

<400> 451

Tyr Ile Ser Asp Cys Pro Tyr His Pro Met Cys His Asp Tyr Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 452
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV62-2011 LC

<400> 452

Gln Gly Gln Ser Gly Ser Phe Arg Asn Thr Cys Pro Tyr His Pro Met
1 5 10 15

Cys His Asp Tyr Arg Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile
35 40 45

3338059PC02_ST25.txt

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 453
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV62-2011 LC

<400> 453

Phe Arg Asn Thr Cys Pro Tyr His Pro Met Cys His Asp Tyr Arg Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 454
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV63-2011 LC

<400> 454

Gln Gly Gln Ser Gly Ser Arg Glu Cys His Met Trp Met Phe Gly Val
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

3338059PC02_ST25.txt

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 455
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV63-2011 LC

<400> 455

Arg Glu Cys His Met Trp Met Phe Gly Val Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 456
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV64-2011 LC

<400> 456

Gln Gly Gln Ser Gly Ser Ala Val Cys His Met Tyr Met Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Phe Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

3338059PC02_ST25.txt

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys Cys
260

<210> 457
<211> 255
<212> PRT
<213> Artificial Sequence

<220>

<223> YV64-2011 LC

<400> 457

Ala Val Cys His Met Tyr Met Tyr Gly Tyr Asn Leu Cys Pro Phe Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 458
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV65-2011 LC

<400> 458

Gln Gly Gln Ser Gly Gln Gly Arg Ser Cys Pro Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

3338059PC02_ST25.txt

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 459
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV65-2011 LC

<400> 459

Arg Ser Cys Pro Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 460
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV66-2011 LC

<400> 460

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Phe Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Asp Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

3338059PC02_ST25.txt

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 461
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV66-2011 LC

<400> 461

Gln Pro Cys Ala Gln Met Phe Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Asp Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 462
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV01-2012 LC

<400> 462

Gln Gly Gln Ser Gly Gln Gly Asp Phe Ser Cys Leu His Ser Met Tyr
 1 5 10 15

Asn Val Cys Leu Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

3338059PC02_ST25.txt

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 463
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV01-2012 LC

<400> 463

Asp Phe Ser Cys Leu His Ser Met Tyr Asn Val Cys Leu Asp Pro Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 464
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV02-2012 LC

<400> 464

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

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Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 465
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV02-2012 LC

<400> 465

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 466
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV03-2012 LC

<400> 466

Gln Gly Gln Ser Gly Gln Gly Leu His Cys Arg Thr Gln Met Tyr Gly
 1 5 10 15

Tyr Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

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Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 467
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV03-2012 LC

<400> 467

Leu His Cys Arg Thr Gln Met Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 468

<211> 261

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV04-2012 LC

<400> 468

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

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Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 469
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-2012 LC

<400> 469

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 470
 <211> 258
 <212> PRT
 <213> Artificial Sequence

<220>

<223> Spacer - YV23-2012 LC

<400> 470

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu
 20 25 30

Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr
 35 40 45

Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu
 50 55 60

Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr
 65 70 75 80

Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe
 85 90 95

Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly
 100 105 110

Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala
 115 120 125

Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln
 130 135 140

Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe
 145 150 155 160

Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val
 165 170 175

Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp
 180 185 190

3338059PC02_ST25.txt

Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr
195 200 205

Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr
210 215 220

Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val
225 230 235 240

Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly
245 250 255

Glu Cys

<210> 471
<211> 252
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-2012 LC

<400> 471

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn
20 25 30

Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu
35 40 45

Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln
50 55 60

Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln
65 70 75 80

Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile
85 90 95

Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr
100 105 110

Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln
115 120 125

Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile
130 135 140

Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp
145 150 155 160

Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn
165 170 175

Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu
180 185 190

Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp
195 200 205

Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr
210 215 220

Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser
225 230 235 240

Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250

<210> 472
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV39-2012 LC

<400> 472

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

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

Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

3338059PC02_ST25.txt

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 473
<211> 253
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2012 LC

<400> 473

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Ala
20 25 30

Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 474
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV61-2012 LC

<400> 474

Gln Gly Gln Ser Gly Ser Tyr Ile Ser Asp Cys Pro Tyr His Pro Met
 1 5 10 15

3338059PC02_ST25.txt

Cys His Asp Tyr Gln Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

3338059PC02_ST25.txt

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 475
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV61-2012 LC

<400> 475

Tyr Ile Ser Asp Cys Pro Tyr His Pro Met Cys His Asp Tyr Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 476
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV62-2012 LC

<400> 476

Gln Gly Gln Ser Gly Ser Phe Arg Asn Thr Cys Pro Tyr His Pro Met
 1 5 10 15

Cys His Asp Tyr Arg Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

3338059PC02_ST25.txt

Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

3338059PC02_ST25.txt

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 477
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV62-2012 LC

<400> 477

Phe Arg Asn Thr Cys Pro Tyr His Pro Met Cys His Asp Tyr Arg Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 478
<211> 261
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV63-2012 LC

<400> 478

Gln Gly Gln Ser Gly Ser Arg Glu Cys His Met Trp Met Phe Gly Val
1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
20 25 30

Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile
35 40 45

3338059PC02_ST25.txt

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
 195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
 210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
 225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
 245 250 255

Asn Arg Gly Glu Cys
260

<210> 479
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV63-2012 LC

<400> 479

Arg Glu Cys His Met Trp Met Phe Gly Val Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 480
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV64-2012 LC

<400> 480

Gln Gly Gln Ser Gly Ser Ala Val Cys His Met Tyr Met Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Phe Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser
 20 25 30

Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

3338059PC02_ST25.txt

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 481
 <211> 255
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV64-2012 LC

<400> 481

Ala Val Cys His Met Tyr Met Tyr Gly Tyr Asn Leu Cys Pro Phe Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 482
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV65-2012 LC

<400> 482

Gln Gly Gln Ser Gly Gln Gly Arg Ser Cys Pro Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

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Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 483
<211> 255
<212> PRT
<213> Artificial Sequence

<220>

<223> YV65-2012 LC

<400> 483

Arg Ser Cys Pro Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 484
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV66-2012 LC

<400> 484

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Phe Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ile Ser
 20 25 30

Ser Gly Leu Leu Ser Gly Arg Ser Ala Asn Pro Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 485
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV66-2012 LC

<400> 485

Gln Pro Cys Ala Gln Met Phe Gly Tyr Ser Met Cys Pro His Thr Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg
 20 25 30

Ser Ala Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
 35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
 50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
 65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 486
 <211> 267
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV01-3001 LC

<400> 486

Gln Gly Gln Ser Gly Gln Gly Asp Phe Ser Cys Leu His Ser Met Tyr
 1 5 10 15

Asn Val Cys Leu Asp Pro Gly Gly Gly Ser Ser Gly Gly Ser Ala Val
 20 25 30

Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His
 35 40 45

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 50 55 60

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

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 85 90 95

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 100 105 110

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Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
115 120 125

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
130 135 140

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
145 150 155 160

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
165 170 175

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
180 185 190

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
195 200 205

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
210 215 220

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
225 230 235 240

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
245 250 255

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 487
<211> 260
<212> PRT
<213> Artificial Sequence

<220>
<223> YV01-3001 LC

<400> 487

Asp Phe Ser Cys Leu His Ser Met Tyr Asn Val Cys Leu Asp Pro Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ala Val Gly Leu Leu Ala Pro Pro Gly
 20 25 30

Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val
 35 40 45

Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala
 50 55 60

Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala
 65 70 75 80

Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly
 85 90 95

Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly
 100 105 110

Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp
 115 120 125

Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe
 130 135 140

Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser
 145 150 155 160

Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala
 165 170 175

Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val
 180 185 190

Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser
 195 200 205

Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr
 210 215 220

Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys
 225 230 235 240

Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn
 245 250 255

Arg Gly Glu Cys
 260

<210> 488
 <211> 267
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV02-3001 LC

<400> 488

Gln Gly Gln Ser Gly Gln Gly Gln Pro Cys Ala Gln Met Tyr Gly Tyr
 1 5 10 15

Ser Met Cys Pro His Thr Gly Gly Gly Ser Ser Gly Gly Ser Ala Val
 20 25 30

Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His
 35 40 45

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 50 55 60

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

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 85 90 95

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 100 105 110

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Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
115 120 125

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
130 135 140

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
145 150 155 160

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
165 170 175

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
180 185 190

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
195 200 205

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
210 215 220

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
225 230 235 240

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
245 250 255

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 489
<211> 260
<212> PRT
<213> Artificial Sequence

<220>
<223> YV02-3001 LC

<400> 489

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Pro His Thr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ala Val Gly Leu Leu Ala Pro Pro Gly
 20 25 30

Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val
 35 40 45

Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala
 50 55 60

Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala
 65 70 75 80

Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly
 85 90 95

Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly
 100 105 110

Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp
 115 120 125

Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe
 130 135 140

Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser
 145 150 155 160

Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala
 165 170 175

Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val
 180 185 190

Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser
 195 200 205

Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr
 210 215 220

Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys
 225 230 235 240

Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn
 245 250 255

Arg Gly Glu Cys
 260

<210> 490
 <211> 267
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV03-3001 LC

<400> 490

Gln Gly Gln Ser Gly Gln Gly Leu His Cys Arg Thr Gln Met Tyr Gly
 1 5 10 15

Tyr Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Ala Val
 20 25 30

Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His
 35 40 45

Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser
 50 55 60

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

Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala
 85 90 95

Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro
 100 105 110

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Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile
115 120 125

Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr
130 135 140

Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys
145 150 155 160

Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu
165 170 175

Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe
180 185 190

Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln
195 200 205

Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser
210 215 220

Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu
225 230 235 240

Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser
245 250 255

Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 491
<211> 260
<212> PRT
<213> Artificial Sequence

<220>
<223> YV03-3001 LC

<400> 491

Leu His Cys Arg Thr Gln Met Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ser Ala Val Gly Leu Leu Ala Pro Pro Gly
 20 25 30

Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val
 35 40 45

Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala
 50 55 60

Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala
 65 70 75 80

Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly
 85 90 95

Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly
 100 105 110

Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp
 115 120 125

Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe
 130 135 140

Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser
 145 150 155 160

Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala
 165 170 175

Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val
 180 185 190

Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser
 195 200 205

Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr
 210 215 220

Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys
 225 230 235 240

Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn
 245 250 255

Arg Gly Glu Cys
 260

<210> 492
 <211> 265
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV04-3001 LC

<400> 492

Gln Gly Gln Ser Gly Ser Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr
 1 5 10 15

Asn Leu Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu
 20 25 30

Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly
 35 40 45

Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser
 50 55 60

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

Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg
 85 90 95

Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg
 100 105 110

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Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg
115 120 125

Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser
130 135 140

Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
145 150 155 160

Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu
165 170 175

Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro
180 185 190

Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly
195 200 205

Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr
210 215 220

Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His
225 230 235 240

Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val
245 250 255

Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 493
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> YV04-3001 LC

<400> 493

Leu His Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly
 20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
 225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
 245 250 255

Gly Glu Cys

<210> 494
 <211> 260
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV06-3001 LC

<400> 494

Gln Gly Gln Ser Gly Ser Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys
 1 5 10 15

Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly
 20 25 30

Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val
 35 40 45

Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala
 50 55 60

Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala
 65 70 75 80

Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly
 85 90 95

Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly
 100 105 110

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Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp
115 120 125

Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe
130 135 140

Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser
145 150 155 160

Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala
165 170 175

Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val
180 185 190

Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser
195 200 205

Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr
210 215 220

Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys
225 230 235 240

Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn
245 250 255

Arg Gly Glu Cys
260

<210> 495
<211> 254
<212> PRT
<213> Artificial Sequence

<220>
<223> YV06-3001 LC

<400> 495

Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Gly Gly Gly Ser Ser Gly
1 5 10 15

Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser
 20 25 30

Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly
 35 40 45

Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala
 50 55 60

Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro
 65 70 75 80

Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr
 85 90 95

Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr
 100 105 110

Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys
 115 120 125

Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val
 130 135 140

Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro
 145 150 155 160

Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu
 165 170 175

Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn
 180 185 190

Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser
 195 200 205

Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala
 210 215 220

Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly
 225 230 235 240

Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 496
 <211> 265
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV09-3001 LC

<400> 496

Gln Gly Gln Ser Gly Ser Phe Gly Thr Ala Cys Pro Asn His Pro Met
 1 5 10 15

Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu
 20 25 30

Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly
 35 40 45

Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser
 50 55 60

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

Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg
 85 90 95

Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg
 100 105 110

Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg
 115 120 125

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Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser
130 135 140

Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
145 150 155 160

Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu
165 170 175

Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro
180 185 190

Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly
195 200 205

Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr
210 215 220

Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His
225 230 235 240

Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val
245 250 255

Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 497
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> YV09-3001 LC

<400> 497

Phe Gly Thr Ala Cys Pro Asn His Pro Met Cys His Asp Trp Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly
20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
 225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
 245 250 255

Gly Glu Cys

<210> 498
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV23-3001 LC

<400> 498

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Leu Tyr Asn Val Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

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Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 499
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV23-3001 LC

<400> 499

Cys Leu His Ser Leu Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
 35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
 50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
 65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
 85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
 100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
 115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
 130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
 145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 500
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV27-3001 LC

<400> 500

Gln Gly Gln Ser Gly Ser Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser
 1 5 10 15

Met Cys Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

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Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 501
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV27-3001 LC

<400> 501

Gln Pro Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
 50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
 65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
 85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
 100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
 115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
 130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
 145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 502
 <211> 263
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV29-3001 LC

<400> 502

Gln Gly Gln Ser Gly Ser Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys
 1 5 10 15

Ala His Thr Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala
 20 25 30

Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser
 35 40 45

Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
 50 55 60

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser
 65 70 75 80

Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
 85 90 95

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu
 115 120 125

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 130 135 140

Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
 145 150 155 160

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Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
165 170 175

Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
180 185 190

Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
195 200 205

Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
210 215 220

Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
225 230 235 240

Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
245 250 255

Ser Phe Asn Arg Gly Glu Cys
260

<210> 503
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> YV29-3001 LC

<400> 503

Cys Ala Gln Met Tyr Gly Tyr Ser Met Cys Ala His Thr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
245 250 255

Cys

<210> 504
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV32-3001 LC

<400> 504

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Leu Cys His Asp Trp
 1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro
 20 25 30

Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

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Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 505
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV32-3001 LC

<400> 505

Cys Pro Asn His Pro Leu Cys His Asp Trp Gln Gly Gly Gly Ser Ser
1 5 10 15

Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 506

<211> 261

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer - YV33-3001 LC

<400> 506

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Met Cys Ala Asp Trp
 1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro
 20 25 30

Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

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Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
180 185 190

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 507
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV33-3001 LC

<400> 507

Cys Pro Asn His Pro Met Cys Ala Asp Trp Gln Gly Gly Gly Ser Ser
1 5 10 15

Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
 85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
 100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
 115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
 130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
 145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
 165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
 180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
 195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
 210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
 225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 508
 <211> 261
 <212> PRT
 <213> Artificial Sequence

<220>

<223> Spacer - YV35-3001 LC

<400> 508

Gln Gly Gln Ser Gly Ser Cys Pro Asn His Pro Met Cys His Asp Ala
 1 5 10 15

Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro
 20 25 30

Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile
 35 40 45

Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg
 50 55 60

Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu
 65 70 75 80

Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr
 85 90 95

Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser
 100 105 110

Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu
 115 120 125

Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr
 130 135 140

Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro
 145 150 155 160

Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr
 165 170 175

Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys
 180 185 190

3338059PC02_ST25.txt

Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu
195 200 205

Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser
210 215 220

Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala
225 230 235 240

Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe
245 250 255

Asn Arg Gly Glu Cys
260

<210> 509
<211> 255
<212> PRT
<213> Artificial Sequence

<220>
<223> YV35-3001 LC

<400> 509

Cys Pro Asn His Pro Met Cys His Asp Ala Gln Gly Gly Gly Ser Ser
1 5 10 15

Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg
20 25 30

Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro
35 40 45

Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg
50 55 60

Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys
65 70 75 80

Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala
85 90 95

Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe
100 105 110

Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr
115 120 125

Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys
130 135 140

Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro
145 150 155 160

Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu
165 170 175

Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp
180 185 190

Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp
195 200 205

Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys
210 215 220

Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln
225 230 235 240

Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 510
<211> 263
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV39-3001 LC

<400> 510

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

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

Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser
 35 40 45

Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
 50 55 60

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser
 65 70 75 80

Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
 85 90 95

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu
 115 120 125

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 130 135 140

Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
 145 150 155 160

Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
 165 170 175

Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
 180 185 190

Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
 195 200 205

3338059PC02_ST25.txt

Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
210 215 220

Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
225 230 235 240

Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
245 250 255

Ser Phe Asn Arg Gly Glu Cys
260

<210> 511
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-3001 LC

<400> 511

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
 115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
 130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
 145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
 165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
 180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
 195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
 210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
 225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
 245 250 255

Cys

<210> 512
 <211> 263
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV41-3001 LC

<400> 512

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

Cys Pro Ala Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala
 20 25 30

Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser
 35 40 45

Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly
 50 55 60

Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser
 65 70 75 80

Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu
 85 90 95

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

Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu
 115 120 125

Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro
 130 135 140

Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala
 145 150 155 160

Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser
 165 170 175

Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu
 180 185 190

Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser
 195 200 205

3338059PC02_ST25.txt

Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu
210 215 220

Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val
225 230 235 240

Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys
245 250 255

Ser Phe Asn Arg Gly Glu Cys
260

<210> 513
<211> 257
<212> PRT
<213> Artificial Sequence

<220>
<223> YV41-3001 LC

<400> 513

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Ala Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser
20 25 30

Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln
35 40 45

Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser
50 55 60

Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln
65 70 75 80

Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser
85 90 95

Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr
100 105 110

Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val
 115 120 125

Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly
 130 135 140

Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile
 145 150 155 160

Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val
 165 170 175

Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys
 180 185 190

Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu
 195 200 205

Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu
 210 215 220

Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr
 225 230 235 240

His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu
 245 250 255

Cys

<210> 514
 <211> 265
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV51-3001 LC

<400> 514

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Ile
 1 5 10 15

Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu
 20 25 30

Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly
 35 40 45

Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser
 50 55 60

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

Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg
 85 90 95

Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg
 100 105 110

Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg
 115 120 125

Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser
 130 135 140

Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
 145 150 155 160

Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu
 165 170 175

Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro
 180 185 190

Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly
 195 200 205

3338059PC02_ST25.txt

Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr
210 215 220

Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His
225 230 235 240

Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val
245 250 255

Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 515
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> YV51-3001 LC

<400> 515

Phe Gly Ala Ala Cys Pro Asn His Pro Ile Cys His Asp Trp Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly
20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
 115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
 130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
 145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
 165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
 180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
 195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
 210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
 225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
 245 250 255

Gly Glu Cys

<210> 516
 <211> 265
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV52-3001 LC

<400> 516

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Leu
 1 5 10 15

Cys His Asp Trp Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu
 20 25 30

Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly
 35 40 45

Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser
 50 55 60

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

Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg
 85 90 95

Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg
 100 105 110

Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg
 115 120 125

Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser
 130 135 140

Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
 145 150 155 160

Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu
 165 170 175

Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro
 180 185 190

Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly
 195 200 205

3338059PC02_ST25.txt

Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr
 210 215 220

Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His
 225 230 235 240

Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val
 245 250 255

Thr Lys Ser Phe Asn Arg Gly Glu Cys
 260 265

<210> 517
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV52-3001 LC

<400> 517

Phe Gly Ala Ala Cys Pro Asn His Pro Leu Cys His Asp Trp Gln Gly
 1 5 10 15

Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly
 20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
 85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
 100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 518
<211> 265
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV53-3001 LC

<400> 518

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Phe Gly Ala Ala Cys Pro Asn His Pro Met
 1 5 10 15

Cys His Asp Ala Gln Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu
 20 25 30

Leu Ala Pro Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly
 35 40 45

Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser
 50 55 60

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

Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg
 85 90 95

Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg
 100 105 110

Phe Ser Gly Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg
 115 120 125

Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser
 130 135 140

Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr
 145 150 155 160

Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu
 165 170 175

Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro
 180 185 190

Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly
 195 200 205

3338059PC02_ST25.txt

Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr
210 215 220

Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His
225 230 235 240

Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val
245 250 255

Thr Lys Ser Phe Asn Arg Gly Glu Cys
260 265

<210> 519
<211> 259
<212> PRT
<213> Artificial Sequence

<220>
<223> YV53-3001 LC

<400> 519

Phe Gly Ala Ala Cys Pro Asn His Pro Met Cys His Asp Ala Gln Gly
1 5 10 15

Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly
20 25 30

Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu
35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
65 70 75 80

Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 520
<211> 262
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV54-3001 LC

<400> 520

3338059PC02_ST25.txt

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Ala Cys Leu
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
 165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
 180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
 195 200 205

3338059PC02_ST25.txt

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 521
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV54-3001 LC

<400> 521

Cys Leu His Ser Ala Tyr Asn Ala Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
 115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
 130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
 145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 522
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV55-3001 LC

<400> 522

Gln Gly Gln Ser Gly Ser Cys Ala His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

3338059PC02_ST25.txt

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
 165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
 180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
 195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
 210 215 220

3338059PC02_ST25.txt

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 523
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV55-3001 LC

<400> 523

Cys Ala His Ser Ala Tyr Asn Val Cys Leu Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
 130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
 145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 524
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV56-3001 LC

<400> 524

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Ala
 1 5 10 15

Asp Pro Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

3338059PC02_ST25.txt

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
 65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
 85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
 115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
 130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
 145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
 165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
 180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
 195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
 210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
 225 230 235 240

3338059PC02_ST25.txt

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 525
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV56-3001 LC

<400> 525

Cys Leu His Ser Ala Tyr Asn Val Cys Ala Asp Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
245 250 255

<210> 526
<211> 262
<212> PRT
<213> Artificial Sequence

<220>
<223> Spacer - YV57-3001 LC

<400> 526

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Leu
1 5 10 15

Ala Pro Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
35 40 45

3338059PC02_ST25.txt

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
50 55 60

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 527
<211> 256
<212> PRT
<213> Artificial Sequence

<220>
<223> YV57-3001 LC

<400> 527

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Ala Pro Gly Gly Gly Ser
1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 528
 <211> 262
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV58-3001 LC

<400> 528

Gln Gly Gln Ser Gly Ser Cys Leu His Ser Ala Tyr Asn Val Cys Leu
 1 5 10 15

Asp Ala Gly Gly Gly Ser Ser Gly Gly Ala Val Gly Leu Leu Ala Pro
 20 25 30

Pro Gly Gly Leu Ser Gly Arg Ser Asp Asn His Gly Gly Gly Ser Glu
 35 40 45

Ile Val Leu Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu
 50 55 60

3338059PC02_ST25.txt

Arg Ala Thr Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr
65 70 75 80

Leu Ala Trp Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile
85 90 95

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

Ser Gly Ser Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro
115 120 125

Glu Asp Phe Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp
130 135 140

Thr Phe Gly Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala
145 150 155 160

Pro Ser Val Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly
165 170 175

Thr Ala Ser Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala
180 185 190

Lys Val Gln Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln
195 200 205

Glu Ser Val Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser
210 215 220

Ser Thr Leu Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr
225 230 235 240

Ala Cys Glu Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser
245 250 255

Phe Asn Arg Gly Glu Cys
260

<210> 529
 <211> 256
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> YV58-3001 LC

<400> 529

Cys Leu His Ser Ala Tyr Asn Val Cys Leu Asp Ala Gly Gly Gly Ser
 1 5 10 15

Ser Gly Gly Ala Val Gly Leu Leu Ala Pro Pro Gly Gly Leu Ser Gly
 20 25 30

Arg Ser Asp Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser
 35 40 45

Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys
 50 55 60

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln
 65 70 75 80

Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg
 85 90 95

Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser Gly Thr Asp
 100 105 110

Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr
 115 120 125

Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr
 130 135 140

Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe
 145 150 155 160

Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys
 165 170 175

Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val
 180 185 190

Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln
 195 200 205

Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser
 210 215 220

Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His
 225 230 235 240

Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250 255

<210> 530
 <211> 259
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer - YV39-NSUB LC

<400> 530

Gln Gly Gln Ser Gly Ser Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu
 1 5 10 15

Cys Pro Tyr Gly Gly Gly Ser Ser Gly Gly Ser Gly Gly Ser Gly Gly
 20 25 30

Ser Gly Gly Gly Ser Gly Gly Gly Ser Gly Gly Ser Glu Ile Val Leu
 35 40 45

Thr Gln Ser Pro Gly Thr Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr
 50 55 60

Leu Ser Cys Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp
 65 70 75 80

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Tyr Gln Gln Lys Pro Gly Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala
85 90 95

Phe Ser Arg Ala Thr Gly Ile Pro Asp Arg Phe Ser Gly Ser Gly Ser
100 105 110

Gly Thr Asp Phe Thr Leu Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe
115 120 125

Ala Val Tyr Tyr Cys Gln Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly
130 135 140

Gln Gly Thr Lys Val Glu Ile Lys Arg Thr Val Ala Ala Pro Ser Val
145 150 155 160

Phe Ile Phe Pro Pro Ser Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser
165 170 175

Val Val Cys Leu Leu Asn Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln
180 185 190

Trp Lys Val Asp Asn Ala Leu Gln Ser Gly Asn Ser Gln Glu Ser Val
195 200 205

Thr Glu Gln Asp Ser Lys Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu
210 215 220

Thr Leu Ser Lys Ala Asp Tyr Glu Lys His Lys Val Tyr Ala Cys Glu
225 230 235 240

Val Thr His Gln Gly Leu Ser Ser Pro Val Thr Lys Ser Phe Asn Arg
245 250 255

Gly Glu Cys

<210> 531
<211> 253
<212> PRT
<213> Artificial Sequence

<220>

<223> YV39-NSUB LC

<400> 531

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
 1 5 10 15

Ser Ser Gly Gly Ser Gly Gly Ser Gly Gly Ser Gly Gly Gly Ser Gly
 20 25 30

Gly Gly Ser Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
 35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
 50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
 65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
 85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
 115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
 130 135 140

Ile Lys Arg Thr Val Ala Ala Pro Ser Val Phe Ile Phe Pro Pro Ser
 145 150 155 160

Asp Glu Gln Leu Lys Ser Gly Thr Ala Ser Val Val Cys Leu Leu Asn
 165 170 175

Asn Phe Tyr Pro Arg Glu Ala Lys Val Gln Trp Lys Val Asp Asn Ala
 180 185 190

Leu Gln Ser Gly Asn Ser Gln Glu Ser Val Thr Glu Gln Asp Ser Lys
 195 200 205

Asp Ser Thr Tyr Ser Leu Ser Ser Thr Leu Thr Leu Ser Lys Ala Asp
 210 215 220

Tyr Glu Lys His Lys Val Tyr Ala Cys Glu Val Thr His Gln Gly Leu
 225 230 235 240

Ser Ser Pro Val Thr Lys Ser Phe Asn Arg Gly Glu Cys
 245 250

<210> 532
 <211> 2
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Linkers

<220>
 <221> MISC_FEATURE
 <222> (1)..(2)
 <223> sequence may be repeated any number of times

<400> 532

Gly Ser
 1

<210> 533
 <211> 3
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Linkers

<220>
 <221> MISC_FEATURE
 <222> (1)..(3)
 <223> sequence may be repeated any number of times

<400> 533

Gly Gly Ser
1

<210> 534

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<220>

<221> MISC_FEATURE

<222> (1)..(5)

<223> sequence may be repeated any number of times

<400> 534

Gly Ser Gly Gly Ser
1 5

<210> 535

<211> 4

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<220>

<221> MISC_FEATURE

<222> (1)..(4)

<223> sequence may be repeated any number of times

<400> 535

Gly Gly Gly Ser
1

<210> 536

<211> 4

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<400> 536

Gly Gly Ser Gly
1

<210> 537

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<400> 537

Gly Gly Ser Gly Gly
1 5

<210> 538

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<400> 538

Gly Ser Gly Ser Gly
1 5

<210> 539

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Linkers

<400> 539

Gly Ser Gly Gly Gly
1 5

<210> 540

<211> 5

<212> PRT
<213> Artificial Sequence

<220>
<223> Linkers

<400> 540

Gly Gly Gly Ser Gly
1 5

<210> 541
<211> 5
<212> PRT
<213> Artificial Sequence

<220>
<223> Linkers

<400> 541

Gly Ser Ser Ser Gly
1 5

<210> 542
<211> 8
<212> PRT
<213> Artificial Sequence

<220>
<223> Linkers

<400> 542

Gly Gly Gly Ser Ser Gly Gly Ser
1 5

<210> 543
<211> 4
<212> PRT
<213> Artificial Sequence

<220>
<223> Linkers

<400> 543

Gly Gly Gly Ser
1

<210> 544
 <211> 7
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer

<400> 544

Gln Gly Gln Ser Gly Gln Gly
 1 5

<210> 545
 <211> 6
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer

<400> 545

Gly Gln Ser Gly Gln Gly
 1 5

<210> 546
 <211> 6
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer

<400> 546

Gln Gly Gln Ser Gly Ser
 1 5

<210> 547
 <211> 6
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer

<400> 547

Gln Gly Gln Ser Gly Gln
1 5

<210> 548

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 548

Gln Ser Gly Gln Gly
1 5

<210> 549

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 549

Gly Gln Ser Gly Ser
1 5

<210> 550

<211> 5

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 550

Gln Gly Gln Ser Gly
1 5

<210> 551

<211> 4

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 551

Ser Gly Gln Gly
1

<210> 552

<211> 4

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 552

Gln Ser Gly Ser
1

<210> 553

<211> 4

<212> PRT

<213> Artificial Sequence

<220>

<223> Spacer

<400> 553

Gln Gly Gln Ser
1

<210> 554

<211> 15

<212> PRT

<213> Artificial Sequence

<220>

<223> Consensus Sequence

<220>

<221> MISC_FEATURE

<222> (1)..(2)

<223> Xaa can be any amino acid

<220>
 <221> MISC_FEATURE
 <222> (4)..(6)
 <223> Xaa can be any amino acid

<400> 554

Xaa	Xaa	Cys	Xaa	Xaa	Xaa	Met	Tyr	Gly	Tyr	Asn	Leu	Cys	Pro	Tyr
1				5					10					15

<210> 555
 <211> 15
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Consensus Sequence

<220>
 <221> MISC_FEATURE
 <222> (1)..(3)
 <223> Xaa can be any amino acid

<220>
 <221> MISC_FEATURE
 <222> (5)..(5)
 <223> Xaa can be any amino acid

<400> 555

Xaa	Xaa	Xaa	Cys	Xaa	His	Ser	Met	Tyr	Asn	Val	Cys	Leu	Asp	Pro
1				5					10					15

<210> 556
 <211> 19
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> Spacer-YV39

<400> 556

Gln	Gly	Gln	Ser	Gly	Ser	Cys	Arg	Thr	Gln	Leu	Tyr	Gly	Tyr	Asn	Leu
1				5					10					15	

Cys Pro Tyr

<210> 557
 <211> 5
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> CDR1 HC for Ipilimumab

<400> 557

Ser Tyr Thr Met His
 1 5

<210> 558
 <211> 17
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> CDR2 HC for Ipilimumab

<400> 558

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

Gly

<210> 559
 <211> 9
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> CDR3 HC for Ipilimumab

<400> 559

Thr Gly Trp Leu Gly Pro Phe Asp Tyr
 1 5

<210> 560
 <211> 12
 <212> PRT
 <213> Artificial Sequence

<220>

<223> CDR1 LC for Ipilimumab

<400> 560

Arg Ala Ser Gln Ser Val Gly Ser Ser Tyr Leu Ala
1 5 10

<210> 561

<211> 7

<212> PRT

<213> Artificial Sequence

<220>

<223> CDR2 LC for Ipilimumab

<400> 561

Gly Ala Phe Ser Arg Ala Thr
1 5

<210> 562

<211> 9

<212> PRT

<213> Artificial Sequence

<220>

<223> CDR3 LC for Ipilimumab

<400> 562

Gln Gln Tyr Gly Ser Ser Pro Trp Thr
1 5

<210> 563

<211> 146

<212> PRT

<213> Artificial Sequence

<220>

<223> YV39-2001 VL

<400> 563

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

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Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Asn His Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
130 135 140

Ile Lys
145

<210> 564
<211> 146
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2011 VL

<400> 564

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Asp
20 25 30

Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
130 135 140

Ile Lys
145

<210> 565
<211> 146
<212> PRT
<213> Artificial Sequence

<220>
<223> YV39-2012 VL

<400> 565

Cys Arg Thr Gln Leu Tyr Gly Tyr Asn Leu Cys Pro Tyr Gly Gly Gly
1 5 10 15

Ser Ser Gly Gly Ser Ile Ser Ser Gly Leu Leu Ser Gly Arg Ser Ala
20 25 30

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Asn Pro Gly Gly Gly Ser Glu Ile Val Leu Thr Gln Ser Pro Gly Thr
35 40 45

Leu Ser Leu Ser Pro Gly Glu Arg Ala Thr Leu Ser Cys Arg Ala Ser
50 55 60

Gln Ser Val Gly Ser Ser Tyr Leu Ala Trp Tyr Gln Gln Lys Pro Gly
65 70 75 80

Gln Ala Pro Arg Leu Leu Ile Tyr Gly Ala Phe Ser Arg Ala Thr Gly
85 90 95

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

Thr Ile Ser Arg Leu Glu Pro Glu Asp Phe Ala Val Tyr Tyr Cys Gln
115 120 125

Gln Tyr Gly Ser Ser Pro Trp Thr Phe Gly Gln Gly Thr Lys Val Glu
130 135 140

Ile Lys
145

<210> 566
<211> 12
<212> PRT
<213> Artificial Sequence

<220>
<223> MM Consensus Sequence 1

<220>
<221> MISC_FEATURE
<222> (2)..(2)
<223> Xaa can be L, M, V or T

<220>
<221> MISC_FEATURE
<222> (4)..(4)
<223> Xaa can be S, V or I

<220>

<221> MISC_FEATURE
 <222> (5)..(5)
 <223> Xaa can be F, L, M or A

<220>
 <221> MISC_FEATURE
 <222> (6)..(6)
 <223> Xaa can be Y or F

<220>
 <221> MISC_FEATURE
 <222> (8)..(8)
 <223> Xaa can be V or I

<400> 566

Cys	Xaa	Tyr	Xaa	Xaa	Xaa	Asn	Xaa	Cys	Leu	Asp	Pro
1			5						10		

<210> 567
 <211> 13
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> MM Consensus Sequence 2

<220>
 <221> MISC_FEATURE
 <222> (11)..(11)
 <223> Xaa can be P or A

<220>
 <221> MISC_FEATURE
 <222> (12)..(12)
 <223> Xaa can be H, R or A

<400> 567

Cys	Ala	Gln	Met	Tyr	Gly	Tyr	Ser	Met	Cys	Xaa	Xaa	Thr
1			5						10			

<210> 568
 <211> 13
 <212> PRT
 <213> Artificial Sequence

<220>
 <223> MM Consensus Sequence 3

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<220>
<221> MISC_FEATURE
<222> (2)..(2)
<223> Xaa can be any amino acid

<220>
<221> MISC_FEATURE
<222> (3)..(3)
<223> Xaa can be M, I, Y, L, N or F

<220>
<221> MISC_FEATURE
<222> (4)..(4)
<223> Xaa can be Y, W, F, Q or T

<220>
<221> MISC_FEATURE
<222> (5)..(5)
<223> Xaa can be M or Y

<220>
<221> MISC_FEATURE
<222> (8)..(8)
<223> Xaa can be Y, V or F

<220>
<221> MISC_FEATURE
<222> (9)..(9)
<223> Xaa can be N or D

<220>
<221> MISC_FEATURE
<222> (13)..(13)
<223> Xaa can be Y or F

<400> 568

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Cys Xaa Xaa Xaa Xaa Tyr Gly Xaa Xaa Leu Cys Pro Xaa
1           5           10

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<210> 569
<211> 12
<212> PRT
<213> Artificial Sequence

<220>
<223> MM Consensus Sequence 4

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<220>

<221> MISC_FEATURE

<222> (1)..(1)

<223> Xaa can be N or T

<220>

<221> MISC_FEATURE

<222> (2)..(2)

<223> Xaa can be S, T, M or A

<220>

<221> MISC_FEATURE

<222> (5)..(5)

<223> Xaa can be N or Y

<220>

<221> MISC_FEATURE

<222> (8)..(8)

<223> Xaa can be M or L

<220>

<221> MISC_FEATURE

<222> (10)..(10)

<223> Xaa can be H, F or Y

<220>

<221> MISC_FEATURE

<222> (12)..(12)

<223> Xaa can be Y, F or W

<400> 569

Xaa Xaa Cys Pro Xaa His Pro Xaa Cys Xaa Asp Xaa

1

5

10

<210> 570

<211> 14

<212> PRT

<213> Artificial Sequence

<220>

<223> Protease Resistant Linker

<400> 570

Gly Gly Ser Gly Gly Ser Gly Gly Gly Ser Gly Gly Gly Ser

1

5

10