

MESOTHELIN-TARGETED CD40 AGONISTIC MULTISPECIFIC ANTIBODY CONSTRUCTS FOR THE TREATMENT OF SOLID TUMORS

FIELD OF THE INVENTION

[0001] The present invention relates to the field of oncology, specifically cancer immunotherapy. The invention relates to a human agonistic CD40 antibody by engineering a molecule that specifically targets the CD40 pathway on tumor-associated APCs, without systemic CD40 activation.

BACKGROUND OF THE INVENTION

[0002] Treating cancer patients who do not respond to immune checkpoint inhibitors, such as anti-PD1, represent both the primary challenge and an exciting opportunity in the quickly changing landscape of immunotherapy. A major determinant of response to PD1 blockade in solid tumors is the degree of pre-treatment tumor-associated T cell infiltration, with patients having poorly infiltrated, ‘cold’ tumors showing minimum clinical benefit. While clinical data are still emerging, the clinical activity of other T cell-targeted immunotherapies beyond PD1 blockade, such as solid tumor BiTE® antibody constructs or CAR-T approaches, may also be limited in ‘cold’ tumors. Combining T cell-targeted approaches with new classes of therapeutics capable of enhancing T-cell infiltration into solid tumors may be critical for maximizing the number of patients benefiting from immunotherapy.

[0003] CD40 is a member of the TNF receptor (TNFR) superfamily that is preferentially expressed by antigen presenting cells (APCs), such as dendritic cells, B cells and macrophages. Interaction with its trimeric ligand on activated T helper cells results in APC activation that includes upregulation of cytokines/chemokines (such as interleukin-12 [IL-12] and CxCL10), proteins involved in antigen-presentation (such as MHC class I and II ligands), T cell costimulatory ligands (such as CD80 and CD86), and array of other immune modulatory factors (i.e., adhesion molecules and other TNFRs). These “licensed” APCs can then trigger a cascade of events leading to induction of robust adaptive immune responses.

[0004] Therapeutics capable of activating CD40 signaling have the potential to inflame solid tumors through their ability to enhance generation of anti-tumor T cells and enhance T cell recruitment directly into the tumor lesion. Preclinical studies with anti-CD40 agonists suggest that triggering CD40 with crosslinking antibodies on APCs can substitute for CD4 T cell help to license APCs and facilitate the activation as well as expansion of CD8 effector T cells. In addition, CD40-activated macrophages may also exert direct tumoricidal functions. These anti-CD40 agonist antibodies have been demonstrated to be efficacious in multiple syngeneic tumor models alone or in combination with other therapies. Based on these pre-clinical studies several CD40 agonistic antibodies are under investigation in phase I/II clinical trials of solid tumor patients. To date, these monoclonal anti-CD40 antibodies have shown some signs of clinical activity, but are often associated with immune-related

adverse effects, such as cytokine release syndrome and evidence of liver damage. These toxicities limit the dose of CD40 agonist that can be delivered, and thus may result in insufficient activation of the CD40 pathway in tumor-associated APC population, negatively impacting the efficacy of this therapeutic approach. Therapeutics capable of localizing activation of CD40 signaling in tumor tissue may improve anti-tumor effects while limiting systemic toxicities.

[0005] In order to achieve tumor localization, an agonistic multispecific antibody construct that binds both CD40 and a tumor-associated antigen (TAA), mesothelin (MSLN), was designed. Robust agonist activity of this multispecific antibody construct on CD40-expressing APCs is entirely dependent on the presence of neighboring MSLN-expressing cells. The multispecific antibody construct was specifically engineered to lack CD40 agonist activity upon binding to CD40 in the absence of MSLN-expressing cells. In addition, introduction of mutations into the IgG Fc domain limits binding to Fc receptors, thus preventing Fc receptor-mediated CD40 agonist activity. Thus, the MSLN-dependent CD40 agonist multispecific antibody construct described herein has the potential to drive robust CD40-mediated activation of APCs in a manner that is confined primarily to tumor tissue, thus minimizing induction of systemic toxicities.

SUMMARY OF THE INVENTION

[0006] The present invention is directed to a multispecific antibody construct comprising:

[0007] (i) a first antibody comprising two light chains and two heavy chains, wherein

[0008] the light chains comprise a first variable region (VL1) and a light chain constant region (CL);

[0009] the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and CH3 regions;

[0010] the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and

[0011] (ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,

[0012] wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and

[0013] wherein the first antibody specifically binds to and agonizes human CD40 (SEQ ID NO: 1) and the scFv specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2).

[0014] In one embodiment, the multispecific antibody construct specifically agonizes CD40 in a MSLN-dependent fashion.

[0015] In one embodiment, the multispecific antibody construct comprises mutations limiting Fc receptor binding and thus decreases Fc receptor-dependent CD40 agonism.

[0016] In one embodiment, the two light chains are identical and the two heavy chain fusion proteins are identical.

[0017] In one embodiment, the heavy chain comprises an amino acid substitution selected from the group consisting of:

[0018] (i) N297G or N297A;

[0019] (ii) L234A and L235A; and

[0020] (iii) R292C and V302C;

[0021] wherein the amino acid numbering is EU numbering according to Kabat.

[0022] In one embodiment, the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

[0023] In one embodiment, the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0024] SEQ ID NOs: 58, 59, and 60, respectively;

[0025] SEQ ID NOs: 64, 65, and 66, respectively;

[0026] SEQ ID NOs: 70, 71, and 72, respectively;

[0027] SEQ ID NOs: 76, 77, and 78, respectively;

[0028] SEQ ID NOs: 82, 83, and 84, respectively;

[0029] SEQ ID NOs: 88, 89, and 90, respectively;

[0030] SEQ ID NOs: 94, 95, and 96, respectively;

[0031] SEQ ID NOs: 100, 101, and 102, respectively;

[0032] SEQ ID NOs: 106, 107, and 108, respectively;

[0033] SEQ ID NOs: 112, 113, and 114, respectively;

[0034] SEQ ID NOs: 118, 119, and 120, respectively;

[0035] SEQ ID NOs: 124, 125, and 126, respectively; and

[0036] SEQ ID NOs: 130, 131, and 132, respectively;

[0037] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0038] SEQ ID NOs: 136, 137, and 138, respectively;

[0039] SEQ ID NOs: 142, 143, and 144, respectively;

[0040] SEQ ID NOs: 148, 149, and 150, respectively;

[0041] SEQ ID NOs: 154, 155, and 156, respectively;

[0042] SEQ ID NOs: 160, 161, and 162, respectively;

[0043] SEQ ID NOs: 166, 167, and 168, respectively;

[0044] SEQ ID NOs: 172, 173, and 174, respectively;

[0045] SEQ ID NOs: 178, 179, and 180, respectively;

[0046] SEQ ID NOs: 184, 185, and 186, respectively;

[0047] SEQ ID NOs: 190, 191, and 192, respectively;

[0048] SEQ ID NOs: 196, 197, and 198, respectively;

[0049] SEQ ID NOs: 202, 203, and 204, respectively; and

[0050] SEQ ID NOs: 208, 209, and 210, respectively;

[0051] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0052] SEQ ID NOs: 230, 231, and 232, respectively;

[0053] SEQ ID NOs: 236, 237, and 238, respectively;

[0054] SEQ ID NOs: 242, 243, and 244, respectively; and

[0055] SEQ ID NOs: 248, 249, and 250, respectively;

[0056] and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0057] SEQ ID NOs: 254, 255, and 256, respectively;

[0058] SEQ ID NOs: 260, 261, and 262, respectively;

[0059] SEQ ID NOs: 266, 267, and 268, respectively; and

[0060] SEQ ID NOs: 272, 273, and 274, respectively.

[0061] In one embodiment, 1) the VL1 and VH1 are selected from the group consisting of:

[0062] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

[0063] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

[0064] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

[0065] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

[0066] e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

[0067] f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

- [0068] g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [0069] h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;
- [0070] i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;
- [0071] j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;
- [0072] k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;
- [0073] l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and
- [0074] m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;
- [0075] and
- [0076] 2) the VL2 and VH2 are selected from the group consisting of:
- [0077] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;
- [0078] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;
- [0079] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and
- [0080] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

[0081] In one embodiment, the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

[0082] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

[0083] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[0084] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

[0085] In one embodiment,

[0086] 1) the VL1 and VH1 are selected from the group consisting of:

[0087] a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;

[0088] b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;

[0089] c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;

[0090] d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;

[0091] e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;

[0092] f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;

[0093] g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;

[0094] h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;

[0095] i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;

[0096] j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;

[0097] k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;

[0098] l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and

[0099] m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;

[0100] and

[0101] 2) the VL2 and VH2 are selected from the group consisting of:

[0102] a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;

[0103] b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;

[0104] c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and

[0105] d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.

[0106] In one embodiment, the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

[0107] In one embodiment, the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

[0108] In one embodiment, the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.

[0109] In one embodiment, the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.

[0110]

[0111] In one embodiment, the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.

[0112] In one embodiment,

[0113] the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 286, 290, 294, 298, 302, 306, 310, 314, 318, 322, 326, 330, 336, 342, 346, 350, 354, 358, 362, 366, 370, 374, and 378; and

[0114] the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID Nos: 285, 289, 293, 297, 301, 305, 309, 313, 317, 321, 325, 329, 333, 337, 341, 345, 349, 353, 357, 361, 365, 369, 373, and 377.

[0115] In one embodiment, the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:

[0116] SEQ ID NO: 286 and SEQ ID NO: 285, respectively;

[0117] SEQ ID NO: 290 and SEQ ID NO: 289, respectively;

[0118] SEQ ID NO: 294 and SEQ ID NO: 293, respectively;

[0119] SEQ ID NO: 298 and SEQ ID NO: 297, respectively;

[0120] SEQ ID NO: 302 and SEQ ID NO: 301, respectively;

[0121] SEQ ID NO: 306 and SEQ ID NO: 305, respectively;

[0122] SEQ ID NO: 310 and SEQ ID NO: 309, respectively;

[0123] SEQ ID NO: 314 and SEQ ID NO: 313, respectively;

[0124] SEQ ID NO: 318 and SEQ ID NO: 317, respectively;

[0125] SEQ ID NO: 322 and SEQ ID NO: 321, respectively;

[0126] SEQ ID NO: 326 and SEQ ID NO: 325, respectively;

[0127] SEQ ID NO: 330 and SEQ ID NO: 329, respectively;

[0128] SEQ ID NO: 334 and SEQ ID NO: 333, respectively;

[0129] SEQ ID NO: 338 and SEQ ID NO: 337, respectively;

[0130] SEQ ID NO: 342 and SEQ ID NO: 341, respectively;

[0131] SEQ ID NO: 346 and SEQ ID NO: 345, respectively;

[0132] SEQ ID NO: 350 and SEQ ID NO: 349, respectively;

[0133] SEQ ID NO: 354 and SEQ ID NO: 353, respectively;

[0134] SEQ ID NO: 358 and SEQ ID NO: 357, respectively;

[0135] SEQ ID NO: 362 and SEQ ID NO: 361, respectively;

[0136] SEQ ID NO: 366 and SEQ ID NO: 365, respectively;

[0137] SEQ ID NO: 370 and SEQ ID NO: 369, respectively;

[0138] SEQ ID NO: 374 and SEQ ID NO: 373, respectively; and

[0139] SEQ ID NO: 378 and SEQ ID NO: 377, respectively.

[0140] In one aspect, the present invention is directed to a polynucleotide encoding the light chain of the antibody construct of the present invention.

[0141] In one aspect, the present invention is directed to a polynucleotide encoding the heavy chain fusion protein of the antibody construct of the present invention.

[0142] In one aspect, the present invention is directed to a vector comprising the polynucleotide encoding the light chain of the antibody construct, the polynucleotide encoding the heavy chain of the antibody construct, or both.

[0143] In one aspect, the present invention is directed to a host cell transformed or transfected with the vector or the polynucleotide encoding the light chain of the antibody construct and the polynucleotide encoding the heavy chain of the antibody construct.

[0144] In one aspect, the present invention is directed to a process for producing the antibody construct of the present invention, the process comprising culturing a host cell comprising a polynucleotide encoding the light chain and also comprising a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

[0145] In one aspect, the present invention is directed to a pharmaceutical composition comprising the antibody construct according to the present invention and a carrier, stabilizer, excipient, diluent, solubilizer, surfactant, emulsifier, preservative or adjuvant.

[0146] In one aspect, the present invention is directed to a method for treating or ameliorating a solid tumor disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof an effective amount of the antibody construct according to the present invention.

[0147] In one embodiment, the solid tumor disease is selected from the group consisting of: ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer disease or a metastatic cancer disease derived from any of the foregoing.

[0148] In one aspect, the present invention is directed to a kit comprising the antibody construct according to the present invention and, optionally, directions for use.

[0149] In one aspect, the present invention is directed to a multispecific antibody construct comprising:

[0150] (i) a first antibody comprising two light chains and two heavy chains, wherein

[0151] the light chains comprise a first variable region (VL1) and a light chain constant region (CL);

[0152] the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and CH3 regions;

- [0153] the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and
- [0154] (ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,
- [0155] wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and
- [0156] wherein the first antibody specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2) and the scFv specifically binds to and agonizes human CD40 (SEQ ID NO: 1).
- [0157] In one embodiment, the multispecific antibody construct specifically agonizes CD40 in a MSLN-dependent fashion.
- [0158] In one embodiment, the multispecific antibody construct comprises mutations limiting Fc receptor binding and thus decreases Fc receptor-dependent CD40 agonism.
- [0159] In one embodiment, the two light chains are identical and the two heavy chain fusion proteins are identical.
- [0160] In one embodiment, the heavy chain comprises an amino acid substitution selected from the group consisting of:
- [0161] (i) N297G or N297A;
- [0162] (ii) L234A and L235A; and
- [0163] (iii) R292C and V302C;
- [0164] wherein the amino acid numbering is EU numbering according to Kabat.
- [0165] In one embodiment, the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.
- [0166] In one embodiment:
- [0167] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:
- [0168] SEQ ID NOs: 230, 231, and 232, respectively;
- [0169] SEQ ID NOs: 236, 237, and 238, respectively;
- [0170] SEQ ID NOs: 242, 243, and 244, respectively; and
- [0171] SEQ ID NOs: 248, 249, and 250, respectively;
- [0172] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [0173] SEQ ID NOs: 254, 255, and 256, respectively;
- [0174] SEQ ID NOs: 260, 261, and 262, respectively;
- [0175] SEQ ID NOs: 266, 267, and 268, respectively; and

- [0176] SEQ ID NOs: 272, 273, and 274, respectively;
- [0177] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:
- [0178] SEQ ID NOs: 58, 59, and 60, respectively;
- [0179] SEQ ID NOs: 64, 65, and 66, respectively;
- [0180] SEQ ID NOs: 70, 71, and 72, respectively;
- [0181] SEQ ID NOs: 76, 77, and 78, respectively;
- [0182] SEQ ID NOs: 82, 83, and 84, respectively;
- [0183] SEQ ID NOs: 88, 89, and 90, respectively;
- [0184] SEQ ID NOs: 94, 95, and 96, respectively;
- [0185] SEQ ID NOs: 100, 101, and 102, respectively;
- [0186] SEQ ID NOs: 106, 107, and 108, respectively;
- [0187] SEQ ID NOs: 112, 113, and 114, respectively;
- [0188] SEQ ID NOs: 118, 119, and 120, respectively;
- [0189] SEQ ID NOs: 124, 125, and 126, respectively; and
- [0190] SEQ ID NOs: 130, 131, and 132, respectively;
- [0191] and
- [0192] the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [0193] SEQ ID NOs: 136, 137, and 138, respectively;
- [0194] SEQ ID NOs: 142, 143, and 144, respectively;
- [0195] SEQ ID NOs: 148, 149, and 150, respectively;
- [0196] SEQ ID NOs: 154, 155, and 156, respectively;
- [0197] SEQ ID NOs: 160, 161, and 162, respectively;
- [0198] SEQ ID NOs: 166, 167, and 168, respectively;
- [0199] SEQ ID NOs: 172, 173, and 174, respectively;
- [0200] SEQ ID NOs: 178, 179, and 180, respectively;
- [0201] SEQ ID NOs: 184, 185, and 186, respectively;
- [0202] SEQ ID NOs: 190, 191, and 192, respectively;
- [0203] SEQ ID NOs: 196, 197, and 198, respectively;
- [0204] SEQ ID NOs: 202, 203, and 204, respectively; and
- [0205] SEQ ID NOs: 208, 209, and 210, respectively.
- [0206] In one embodiment,
- [0207] 1) the VL1 and VH1 are selected from the group consisting of:

[0208] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

[0209] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

[0210] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

[0211] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;

[0212] and

[0213] 2) the VL2 and VH2 are selected from the group consisting of:

[0214] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

[0215] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

[0216] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

[0217] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

[0218] e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

[0219] f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

[0220] g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

[0221] h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

[0222] i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[0223] j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[0224] k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[0225] l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[0226] m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

[0227] In one embodiment:

[0228] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[0229] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

[0230] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

[0231] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

[0232] In one embodiment:

[0233] 1) the VL1 and VH1 are selected from the group consisting of:

[0234] a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;

[0235] b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;

[0236] c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and

[0237] d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;

[0238] and

[0239] 2) the VL2 and VH2 are selected from the group consisting of:

[0240] a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;

- [0241] b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;
- [0242] c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;
- [0243] d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;
- [0244] e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;
- [0245] f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;
- [0246] g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;
- [0247] h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;
- [0248] i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;
- [0249] j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;
- [0250] k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;
- [0251] l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and
- [0252] m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.
- [0253] In one embodiment, the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.
- [0254] In one embodiment, the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.
- [0255] In one embodiment, the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.
- [0256] In one embodiment, the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.
- [0257] In one embodiment, the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.
- [0258] In one embodiment,
- [0259] the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 382, 386, 390, 394, 398, 402, 406, 410, 414, 418, 422, 426, 430, 434, 438, 442, 446, and 450; and
- [0260] the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID NOs: 381, 385, 389, 393, 397, 401, 405, 409, 413, 417, 421, 425, 429, 433, 437, 441, 445, and 449.
- [0261] In one embodiment, the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:
- [0262] SEQ ID NO: 382 and SEQ ID NO: 381, respectively;
- [0263] SEQ ID NO: 386 and SEQ ID NO: 385, respectively;
- [0264] SEQ ID NO: 390 and SEQ ID NO: 389, respectively;
- [0265] SEQ ID NO: 394 and SEQ ID NO: 393, respectively;
- [0266] SEQ ID NO: 398 and SEQ ID NO: 397, respectively;

[0267] SEQ ID NO: 402 and SEQ ID NO: 401, respectively;

[0268] SEQ ID NO: 406 and SEQ ID NO: 405, respectively;

[0269] SEQ ID NO: 410 and SEQ ID NO: 409, respectively;

[0270] SEQ ID NO: 414 and SEQ ID NO: 413, respectively;

[0271] SEQ ID NO: 418 and SEQ ID NO: 417, respectively;

[0272] SEQ ID NO: 422 and SEQ ID NO: 421, respectively;

[0273] SEQ ID NO: 426 and SEQ ID NO: 425, respectively;

[0274] SEQ ID NO: 430 and SEQ ID NO: 429, respectively;

[0275] SEQ ID NO: 434 and SEQ ID NO: 433, respectively;

[0276] SEQ ID NO: 438 and SEQ ID NO: 437, respectively;

[0277] SEQ ID NO: 442 and SEQ ID NO: 441, respectively;

[0278] SEQ ID NO: 446 and SEQ ID NO: 445, respectively; and

[0279] SEQ ID NO: 450 and SEQ ID NO: 449, respectively.

[0280] In one aspect, the present invention is directed to a multispecific antibody construct comprising:

[0281] a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

[0282] i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

[0283] ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

[0284] b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

[0285] c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

[0286] wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

[0287] wherein:

[0288] the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

[0289] the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

[0290] In one aspect, the present invention is directed to a multispecific antibody construct comprising:

[0291] a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

[0292] i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

[0293] ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

[0294] b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

[0295] c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

[0296] wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

[0297] wherein:

[0298] the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

[0299] the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

[0300] In one embodiment, the hinge-CH2-CH3 polypeptide is linked to the VH2 via a peptide linker.

[0301] In one embodiment, the peptide linker comprises a sequence selected from the group consisting of (Gly₃Ser)₂ (SEQ ID NO: 916), (Gly₄Ser)₂ (SEQ ID NO: 888), (Gly₃Ser)₃ (SEQ ID NO: 917), (Gly₄Ser)₃ (SEQ ID NO: 889), (Gly₃Ser)₄ (SEQ ID NO: 918), (Gly₄Ser)₄ (SEQ ID NO: 890), (Gly₃Ser)₅ (SEQ ID NO: 919), (Gly₄Ser)₅ (SEQ ID NO: 920), (Gly₃Ser)₆ (SEQ ID NO: 921), and (Gly₄Ser)₆ (SEQ ID NO: 922).

[0302] In one embodiment,

[0303] a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;

[0304] b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;

[0305] c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering; and

[0306] d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering.

[0307] In one embodiment,

[0308] a) the first CH1 domain comprises a S183K mutation using EU numbering;

[0309] b) the second CH1 domain comprises a S183E mutation using EU numbering;

[0310] c) the first CL domain comprises a S176E mutation using EU numbering; and

[0311] d) the second CL domain comprises a S176K mutation using EU numbering.

[0312] In one embodiment,

[0313] a) the VH1 comprises a Q39K mutation and the first CH1 domain comprises a S183K mutation using EU numbering;

[0314] b) the VH2 comprises a Q39E mutation and the second CH1 domain comprises a S183E mutation using EU numbering;

[0315] c) the VL1 comprises a Q38E mutation and the first CL domain comprises a S176E mutation using EU numbering; and

[0316] d) the VL2 comprises a Q38K mutation and the second CL domain comprises a S176K mutation using EU numbering.

[0317] In one embodiment,

[0318] a) the first CH1 domain comprises G44K and S183K mutations using EU numbering;

[0319] b) the second CH1 domain comprises G44E and S183E mutations using EU numbering;

[0320] c) the first CL domain comprises G100E and S176E mutations using EU numbering; and

[0321] d) the second CL domain comprises G100K and S176K mutations using EU numbering.

[0322] In one embodiment,

[0323] a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;

[0324] b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;

[0325] c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering; and

[0326] d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering.

[0327] In one embodiment,

[0328] a) the first CH1 domain comprises a S183E mutation using EU numbering;

[0329] b) the second CH1 domain comprises a S183K mutation using EU numbering;

[0330] c) the first CL domain comprises a S176K mutation using EU numbering; and

[0331] d) the second CL domain comprises a S176E mutation using EU numbering.

[0332] In one embodiment,

[0333] a) the VH1 comprises a Q39E mutation and the first CH1 domain comprises a S183E mutation using EU numbering;

[0334] b) the VH2 comprises a Q39K mutation and the second CH1 domain comprises a S183K mutation using EU numbering;

[0335] c) the VL1 comprises a Q38K mutation and the first CL domain comprises a S176K mutation using EU numbering; and

[0336] d) the VL2 comprises a Q38E mutation and the second CL domain comprises a S176E mutation using EU numbering.

[0337] In one embodiment,

[0338] a) the first CH1 domain comprises G44E and S183E mutations using EU numbering;

[0339] b) the second CH1 domain comprises G44K and S183K mutations using EU numbering;

[0340] c) the first CL domain comprises G100K and S176K mutations using EU numbering; and

[0341] d) the second CL domain comprises G100E and S176E mutations using EU numbering.

[0342] In one embodiment, the hinge-CH2-CH3 polypeptide comprises an amino acid substitution selected from the group consisting of:

[0343] (i) N297G or N297A;

[0344] (ii) L234A and L235A; and

[0345] (iii) R292C and V302C;

[0346] wherein the amino acid numbering is EU numbering according to Kabat.

[0347] In one embodiment, the hinge-CH2-CH3 polypeptide comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

[0348] In one embodiment, the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human MSLN (SEQ ID NO: 2); and

[0349] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0350] SEQ ID NOs: 58, 59, and 60, respectively;

[0351] SEQ ID NOs: 64, 65, and 66, respectively;

[0352] SEQ ID NOs: 70, 71, and 72, respectively;

[0353] SEQ ID NOs: 76, 77, and 78, respectively;

[0354] SEQ ID NOs: 82, 83, and 84, respectively;

[0355] SEQ ID NOs: 88, 89, and 90, respectively;

[0356] SEQ ID NOs: 94, 95, and 96, respectively;

[0357] SEQ ID NOs: 100, 101, and 102, respectively;

[0358] SEQ ID NOs: 106, 107, and 108, respectively;

[0359] SEQ ID NOs: 112, 113, and 114, respectively;

[0360] SEQ ID NOs: 118, 119, and 120, respectively;

[0361] SEQ ID NOs: 124, 125, and 126, respectively; and

[0362] SEQ ID NOs: 130, 131, and 132, respectively;

[0363] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0364] SEQ ID NOs: 136, 137, and 138, respectively;

[0365] SEQ ID NOs: 142, 143, and 144, respectively;

[0366] SEQ ID NOs: 148, 149, and 150, respectively;

[0367] SEQ ID NOs: 154, 155, and 156, respectively;

[0368] SEQ ID NOs: 160, 161, and 162, respectively;

[0369] SEQ ID NOs: 166, 167, and 168, respectively;

[0370] SEQ ID NOs: 172, 173, and 174, respectively;

[0371] SEQ ID NOs: 178, 179, and 180, respectively;

[0372] SEQ ID NOs: 184, 185, and 186, respectively;

[0373] SEQ ID NOs: 190, 191, and 192, respectively;

[0374] SEQ ID NOs: 196, 197, and 198, respectively;

[0375] SEQ ID NOs: 202, 203, and 204, respectively; and

[0376] SEQ ID NOs: 208, 209, and 210, respectively;

[0377] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0378] SEQ ID NOs: 230, 231, and 232, respectively;

[0379] SEQ ID NOs: 236, 237, and 238, respectively;

[0380] SEQ ID NOs: 242, 243, and 244, respectively; and

[0381] SEQ ID NOs: 248, 249, and 250, respectively;

[0382] and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0383] SEQ ID NOs: 254, 255, and 256, respectively;

[0384] SEQ ID NOs: 260, 261, and 262, respectively;

[0385] SEQ ID NOs: 266, 267, and 268, respectively; and

[0386] SEQ ID NOs: 272, 273, and 274, respectively.

[0387]

[0388] 64. The antibody construct of claim 63, wherein

[0389] 1) the VL1 and VH1 are selected from the group consisting of:

[0390] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

[0391] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

[0392] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

[0393] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

[0394] e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

[0395] f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

[0396] g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

[0397] h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

[0398] i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[0399] j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[0400] k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[0401] l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[0402] m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;

[0403] and

[0404] 2) the VL2 and VH2 are selected from the group consisting of:

[0405] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

[0406] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

[0407] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

[0408] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

[0409] In one embodiment:

[0410] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

[0411] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

[0412] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[0413] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

[0414] In one embodiment:

[0415] 1) the VL1 and VH1 are selected from the group consisting of:

[0416] a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;

[0417] b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;

[0418] c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;

[0419] d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;

[0420] e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;

[0421] f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;

[0422] g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;

[0423] h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;

[0424] i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;

[0425] j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;

[0426] k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;

[0427] l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and

[0428] m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;

[0429] and

[0430] 2) the VL2 and VH2 are selected from the group consisting of:

[0431] a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;

[0432] b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;

[0433] c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and

[0434] d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.

[0435] In one aspect, the present invention is directed to a polynucleotide encoding the first light chain of the antibody construct.

[0436] In one aspect, the present invention is directed to a polynucleotide encoding the second light chain of the antibody construct.

[0437] In one aspect, the present invention is directed to a polynucleotide encoding the heavy chain fusion protein of the antibody construct.

[0438] In one aspect, the present invention is directed to a vector comprising:

[0439] a) the polynucleotide encoding the first light chain of the antibody construct,

[0440] b) the polynucleotide encoding the second light chain of the antibody construct,

[0441] c) the polynucleotide encoding the heavy chain fusion protein of the antibody construct, or

[0442] d) any combination of a), b), and c).

[0443] In one aspect, the present invention is directed to a host cell transformed or transfected with the vector or polynucleotide according to the present invention.

[0444] In one aspect, the present invention is directed to a process for producing the antibody construct according to the present invention, the process comprising culturing a host cell comprising a polynucleotide encoding the first light chain, a polynucleotide encoding the second light chain and a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

[0445] In one aspect, the present invention is directed to a pharmaceutical composition comprising the antibody construct according to the present invention and a carrier, stabilizer, excipient, diluent, solubilizer, surfactant, emulsifier, preservative or adjuvant.

[0446] In one aspect, the present invention is directed to a method for treating or ameliorating a solid tumor disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof an effective amount of the antibody construct according to the present invention.

[0447] In one embodiment, the solid tumor disease is selected from the group consisting of: ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer disease or a metastatic cancer disease derived from any of the foregoing.

[0448] In one aspect, the present invention is directed to a kit comprising the antibody construct according to the present invention and, optionally, directions for use.

[0449] In one embodiment, the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1); wherein:

[0450] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0451] SEQ ID NOs: 230, 231, and 232, respectively;

[0452] SEQ ID NOs: 236, 237, and 238, respectively;

[0453] SEQ ID NOs: 242, 243, and 244, respectively; and

[0454] SEQ ID NOs: 248, 249, and 250, respectively;

[0455] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0456] SEQ ID NOs: 254, 255, and 256, respectively;

[0457] SEQ ID NOs: 260, 261, and 262, respectively;

[0458] SEQ ID NOs: 266, 267, and 268, respectively; and

[0459] SEQ ID NOs: 272, 273, and 274, respectively;

[0460] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0461] SEQ ID NOs: 58, 59, and 60, respectively;

[0462] SEQ ID NOs: 64, 65, and 66, respectively;

[0463] SEQ ID NOs: 70, 71, and 72, respectively;

[0464] SEQ ID NOs: 76, 77, and 78, respectively;

- [0465] SEQ ID NOs: 82, 83, and 84, respectively;
- [0466] SEQ ID NOs: 88, 89, and 90, respectively;
- [0467] SEQ ID NOs: 94, 95, and 96, respectively;
- [0468] SEQ ID NOs: 100, 101, and 102, respectively;
- [0469] SEQ ID NOs: 106, 107, and 108, respectively;
- [0470] SEQ ID NOs: 112, 113, and 114, respectively;
- [0471] SEQ ID NOs: 118, 119, and 120, respectively;
- [0472] SEQ ID NOs: 124, 125, and 126, respectively; and
- [0473] SEQ ID NOs: 130, 131, and 132, respectively;
- [0474] and
- [0475] the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [0476] SEQ ID NOs: 136, 137, and 138, respectively;
- [0477] SEQ ID NOs: 142, 143, and 144, respectively;
- [0478] SEQ ID NOs: 148, 149, and 150, respectively;
- [0479] SEQ ID NOs: 154, 155, and 156, respectively;
- [0480] SEQ ID NOs: 160, 161, and 162, respectively;
- [0481] SEQ ID NOs: 166, 167, and 168, respectively;
- [0482] SEQ ID NOs: 172, 173, and 174, respectively;
- [0483] SEQ ID NOs: 178, 179, and 180, respectively;
- [0484] SEQ ID NOs: 184, 185, and 186, respectively;
- [0485] SEQ ID NOs: 190, 191, and 192, respectively;
- [0486] SEQ ID NOs: 196, 197, and 198, respectively;
- [0487] SEQ ID NOs: 202, 203, and 204, respectively; and
- [0488] SEQ ID NOs: 208, 209, and 210, respectively.
- [0489] In one embodiment,
- [0490] 1) the VL1 and VH1 are selected from the group consisting of:
- [0491] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;
- [0492] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

- [0493] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and
- [0494] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;
- [0495] and
- [0496] 2) the VL2 and VH2 are selected from the group consisting of:
- [0497] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;
- [0498] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;
- [0499] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;
- [0500] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;
- [0501] e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;
- [0502] f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;
- [0503] g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [0504] h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;
- [0505] i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[0506] j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[0507] k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[0508] l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[0509] m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

[0510] In one embodiment,

[0511] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[0512] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

[0513] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

[0514] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

[0515] In one embodiment,

[0516] 1) the VL1 and VH1 are selected from the group consisting of:

[0517] a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;

[0518] b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;

[0519] c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and

[0520] d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;

[0521] and

[0522] 2) the VL2 and VH2 are selected from the group consisting of:

[0523] a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;

[0524] b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;

[0525] c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;

[0526] d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;

[0527] e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;

[0528] f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;

- [0529] g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;
- [0530] h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;
- [0531] i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;
- [0532] j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;
- [0533] k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;
- [0534] l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and
- [0535] m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.
- [0536] In one aspect, the present invention is directed to an antigen binding protein that specifically binds and agonizes human CD40 (SEQ ID NO: 1), the antigen binding protein comprising a light chain variable region (VL) and a heavy chain variable region (VH), wherein:
- [0537] the VL comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:
- [0538] SEQ ID NOs: 58, 59, and 60, respectively;
- [0539] SEQ ID NOs: 64, 65, and 66, respectively;
- [0540] SEQ ID NOs: 70, 71, and 72, respectively;
- [0541] SEQ ID NOs: 76, 77, and 78, respectively;
- [0542] SEQ ID NOs: 82, 83, and 84, respectively;
- [0543] SEQ ID NOs: 88, 89, and 90, respectively;
- [0544] SEQ ID NOs: 94, 95, and 96, respectively;
- [0545] SEQ ID NOs: 100, 101, and 102, respectively;
- [0546] SEQ ID NOs: 106, 107, and 108, respectively;
- [0547] SEQ ID NOs: 112, 113, and 114, respectively;
- [0548] SEQ ID NOs: 118, 119, and 120, respectively;
- [0549] SEQ ID NOs: 124, 125, and 126, respectively; and
- [0550] SEQ ID NOs: 130, 131, and 132, respectively;
- [0551] and
- [0552] the VH comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [0553] SEQ ID NOs: 136, 137, and 138, respectively;
- [0554] SEQ ID NOs: 142, 143, and 144, respectively;
- [0555] SEQ ID NOs: 148, 149, and 150, respectively;
- [0556] SEQ ID NOs: 154, 155, and 156, respectively;
- [0557] SEQ ID NOs: 160, 161, and 162, respectively;
- [0558] SEQ ID NOs: 166, 167, and 168, respectively;
- [0559] SEQ ID NOs: 172, 173, and 174, respectively;
- [0560] SEQ ID NOs: 178, 179, and 180, respectively;
- [0561] SEQ ID NOs: 184, 185, and 186, respectively;

- [0562] SEQ ID NOs: 190, 191, and 192, respectively;
- [0563] SEQ ID NOs: 196, 197, and 198, respectively;
- [0564] SEQ ID NOs: 202, 203, and 204, respectively; and
- [0565] SEQ ID NOs: 208, 209, and 210, respectively.
- [0566] In one embodiment, the multispecific antibody construct comprises mutations limiting Fc receptor binding and thus decreases Fc receptor-dependent CD40 agonism.
- [0567] In one embodiment,
- [0568] the VL and VH are selected from the group consisting of:
- [0569] a) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;
- [0570] b) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;
- [0571] c) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;
- [0572] d) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;
- [0573] e) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;
- [0574] f) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;
- [0575] g) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [0576] h) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;
- [0577] i) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

- [0578] j) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;
- [0579] k) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;
- [0580] l) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and
- [0581] m) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.
- [0582] In one embodiment,
- [0583] the VL comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and
- [0584] the VH comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.
- [0585] In one embodiment,
- [0586] the VL and VH are selected from the group consisting of:
- [0587] a) a VL comprising SEQ ID NO: 5 and a VH comprising SEQ ID NO: 6;
- [0588] b) a VL comprising SEQ ID NO: 9 and a VH comprising SEQ ID NO: 10;
- [0589] c) a VL comprising SEQ ID NO: 13 and a VH comprising SEQ ID NO: 14;
- [0590] d) a VL comprising SEQ ID NO: 17 and a VH comprising SEQ ID NO: 18;
- [0591] e) a VL comprising SEQ ID NO: 21 and a VH comprising SEQ ID NO: 22;
- [0592] f) a VL comprising SEQ ID NO: 25 and a VH comprising SEQ ID NO: 26;
- [0593] g) a VL comprising SEQ ID NO: 29 and a VH comprising SEQ ID NO: 30;
- [0594] h) a VL comprising SEQ ID NO: 33 and a VH comprising SEQ ID NO: 34;
- [0595] i) a VL comprising SEQ ID NO: 37 and a VH comprising SEQ ID NO: 38;
- [0596] j) a VL comprising SEQ ID NO: 41 and a VH comprising SEQ ID NO: 42;
- [0597] k) a VL comprising SEQ ID NO: 45 and a VH comprising SEQ ID NO: 46;
- [0598] l) a VL comprising SEQ ID NO: 49 and a VH comprising SEQ ID NO: 50; and
- [0599] m) a VL comprising SEQ ID NO: 53 and a VH comprising SEQ ID NO: 54.
- [0600] In one embodiment, the heavy chain comprises an amino acid substitution selected from the group consisting of:
- [0601] (i) N297G or N297A;

[0602] (ii) L234A and L235A; and

[0603] (iii) R292C and V302C;

[0604] wherein the amino acid numbering is EU numbering according to Kabat.

[0605] In one embodiment, the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

[0606] In one embodiment, the antigen binding protein further comprises a light chain CL polypeptide linked to the VL, wherein the CL polypeptide is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

[0607] In one embodiment, the antigen binding protein further comprises a heavy chain CH1-hinge-CH2-CH3 polypeptide wherein the CH1-hinge-CH2-CH3 polypeptide is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

[0608] In one embodiment, the antigen binding protein further comprises a second antigen binding portion that specifically binds a second antigen.

[0609] In one embodiment, the second antigen is a tumor associated antigen (TAA).

BRIEF DESCRIPTION OF THE DRAWINGS

[0610] Figure 1 depicts on cell binding of anti-MSLN antibodies to CHO cells expressing human MSLN.

[0611] Figure 2 depicts Activity of anti-CD40 antibodies with and without crosslinking in human B cell assay.

[0612] Figure 3 depicts Bivalent bispecific antibody formats.

[0613] Figure 4 depicts Generation of cell lines expressing varying levels of MSLN.

[0614] Figure 5 depicts Pharmacokinetic properties and stability of anti-CD40xMSLN bispecific molecules in mouse.

[0615] Figure 6 depicts Generation of MC38-human EPCAM and B16F10-human EPCAM expressing cell line.

[0616] Figure 7 depicts Activity of mouse surrogate anti-CD40xhuman EPCAM bispecific molecule.

[0617] Figure 8 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule has tumor-localized, TAA-mediated X-linking-dependent activity in vivo.

[0618] Figure 9 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule enhances CD8+ T cell anti-tumor responses.

[0619] Figure 10 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule has tumor-localized, TAA-mediated X-linking-dependent activity in vivo.

[0620] Figure 11 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule does not increase levels of systemic cytokines or increase liver inflammation.

[0621] Figure 12 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule induces regression of established MC38-human EPCAM tumors alone and in combination with PD1/PDL1 blockade.

[0622] Figure 13 depicts Mouse surrogate anti-CD40xhuman EPCAM bispecific molecule induces long-lasting immune memory that protects against tumor re-challenge.

DETAILED DESCRIPTION

[0623] Unless otherwise indicated, the term "at least" preceding a series of elements is to be understood to refer to every element in the series. Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the present invention.

[0624] The term "and/or" wherever used herein includes the meaning of "and", "or" and "all or any other combination of the elements connected by said term".

[0625] The term "about" or "approximately" as used herein means within $\pm 20\%$, preferably within $\pm 15\%$, more preferably within $\pm 10\%$, and most preferably within $\pm 5\%$ of a given value or range.

[0626] Throughout this specification and the claims which follow, unless the context requires otherwise, the word "comprise", and variations such as "comprises" and "comprising", will be understood to imply the inclusion of a stated integer or step or group of integers or steps but not the exclusion of any other integer or step or group of integer or step. When used herein the term "comprising" can be substituted with the term "containing" or "including" or sometimes when used herein with the term "having".

[0627] When used herein "consisting of" excludes any element, step, or ingredient not specified in the claim element. When used herein, "consisting essentially of" does not exclude materials or steps that do not materially affect the basic and novel characteristics of the claim.

[0628] In each instance herein any of the terms "comprising", "consisting essentially of" and "consisting of" may be replaced with either of the other two terms.

[0629] As used herein, the term "antigen binding protein" refers to a protein that specifically binds to one or more target antigens. An antigen binding protein can include an antibody and functional fragments thereof. A "functional antibody fragment" is a portion of an antibody that lacks at least some of the amino acids present in a full-length heavy chain and/or light chain, but which is still capable of specifically binding to an antigen. A functional antibody fragment includes, but is not limited to, a Fab fragment, a Fab' fragment, a F(ab')₂ fragment, a Fv fragment, a Fd fragment, and a complementarity determining region (CDR) fragment, and can be derived from any mammalian source, such as human, mouse, rat, rabbit, or camelid. Functional antibody fragments may compete for binding of a target antigen with an intact antibody and the fragments may be produced by the

modification of intact antibodies (e.g. enzymatic or chemical cleavage) or synthesized *de novo* using recombinant DNA technologies or peptide synthesis.

[0630] The term "antibody construct" refers to a molecule in which the structure and/or function is/are based on the structure and/or function of an antibody, e.g., of a full-length or whole immunoglobulin molecule. An antibody construct is hence capable of binding to its specific target or antigen. Furthermore, an antibody construct according to the invention comprises the minimum structural requirements of an antibody which allow for the target binding. This minimum requirement may e.g. be defined by the presence of at least the three light chain CDRs (i.e. CDR1, CDR2 and CDR3 of the VL region) and/or the three heavy chain CDRs (i.e. CDR1, CDR2 and CDR3 of the VH region), preferably of all six CDRs. The antibodies on which the constructs according to the invention are based include for example monoclonal, recombinant, chimeric, deimmunized, humanized and human antibodies.

[0631] In certain embodiments, the antigen binding proteins of the invention comprise antibodies. As used herein, the term "antibody" refers to a tetrameric immunoglobulin protein comprising two light chain polypeptides (about 25 kDa each) and two heavy chain polypeptides (about 50-70 kDa each). The term "light chain" or "immunoglobulin light chain" refers to a polypeptide comprising, from amino terminus to carboxyl terminus, a single immunoglobulin light chain variable region (VL) and a single immunoglobulin light chain constant domain (CL). The immunoglobulin light chain constant domain (CL) can be kappa (κ) or lambda (λ). The term "heavy chain" or "immunoglobulin heavy chain" refers to a polypeptide comprising, from amino terminus to carboxyl terminus, a single immunoglobulin heavy chain variable region (VH), an immunoglobulin heavy chain constant domain 1 (CH1), an immunoglobulin hinge region, an immunoglobulin heavy chain constant domain 2 (CH2), an immunoglobulin heavy chain constant domain 3 (CH3), and optionally an immunoglobulin heavy chain constant domain 4 (CH4). Heavy chains are classified as mu (μ), delta (Δ), gamma (γ), alpha (α), and epsilon (ϵ), and define the antibody's isotype as IgM, IgD, IgG, IgA, and IgE, respectively. The IgG-class and IgA-class antibodies are further divided into subclasses, namely, IgG1, IgG2, IgG3, and IgG4, and IgA1 and IgA2, respectively. The heavy chains in IgG, IgA, and IgD antibodies have three domains (CH1, CH2, and CH3), whereas the heavy chains in IgM and IgE antibodies have four domains (CH1, CH2, CH3, and CH4). The immunoglobulin heavy chain constant domains can be from any immunoglobulin isotype, including subtypes. The antibody chains are linked together via inter-polypeptide disulfide bonds between the CL domain and the CH1 domain (i.e. between the light and heavy chain) and between the hinge regions of the antibody heavy chains.

[0632] In a human antibody, CH1 means a region having the amino acid sequence at positions 118 to 215 of the EU index. A highly flexible amino acid region called a "hinge region" exists between CH1 and CH2. CH2 represents a region having the amino acid sequence at positions 231 to 340 of the EU

index, and CH3 represents a region having the amino acid sequence at positions 341 to 446 of the EU index.

[0633] “CL” represents a constant region of a light chain. In the case of a κ chain of a human antibody, CL represents a region having the amino acid sequence at positions 108 to 214 of the EU index. In a λ chain, CL represents a region having the amino acid sequence at positions 108 to 215.

[0634] Both the EU index as in Kabat *et al.*, Sequences of Proteins of Immunological Interest, 5th Ed. Public Health Service, National Institutes of Health, Bethesda, MD (1991) and AHo numbering schemes (Honegger A. and Plückthun A. *J Mol Biol.* 2001 Jun 8;309(3):657-70) can be used in the present invention. Amino acid positions and complementarity determining regions (CDRs) and framework regions (FR) of a given antibody may be identified using either system. For example, EU heavy chain positions of 39, 44, 183, 356, 357, 360, 370, 392, 399, and 409 are equivalent to AHo heavy chain positions 46, 51, 230, 484, 485, 491, 501, 528, 535, and 551, respectively.

[0635] In one aspect, the present invention is directed to an antigen binding protein that specifically binds and agonizes human CD40 (SEQ ID NO: 1), the antigen binding protein comprising a light chain variable region (VL) and a heavy chain variable region (VH), wherein:

[0636] the VL comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0637] SEQ ID NOs: 58, 59, and 60, respectively;

[0638] SEQ ID NOs: 64, 65, and 66, respectively;

[0639] SEQ ID NOs: 70, 71, and 72, respectively;

[0640] SEQ ID NOs: 76, 77, and 78, respectively;

[0641] SEQ ID NOs: 82, 83, and 84, respectively;

[0642] SEQ ID NOs: 88, 89, and 90, respectively;

[0643] SEQ ID NOs: 94, 95, and 96, respectively;

[0644] SEQ ID NOs: 100, 101, and 102, respectively;

[0645] SEQ ID NOs: 106, 107, and 108, respectively;

[0646] SEQ ID NOs: 112, 113, and 114, respectively;

[0647] SEQ ID NOs: 118, 119, and 120, respectively;

[0648] SEQ ID NOs: 124, 125, and 126, respectively; and

[0649] SEQ ID NOs: 130, 131, and 132, respectively;

[0650] and

[0651] the VH comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0652] SEQ ID NOs: 136, 137, and 138, respectively;

[0653] SEQ ID NOs: 142, 143, and 144, respectively;

[0654] SEQ ID NOs: 148, 149, and 150, respectively;

[0655] SEQ ID NOs: 154, 155, and 156, respectively;

- [0656] SEQ ID NOs: 160, 161, and 162, respectively;
- [0657] SEQ ID NOs: 166, 167, and 168, respectively;
- [0658] SEQ ID NOs: 172, 173, and 174, respectively;
- [0659] SEQ ID NOs: 178, 179, and 180, respectively;
- [0660] SEQ ID NOs: 184, 185, and 186, respectively;
- [0661] SEQ ID NOs: 190, 191, and 192, respectively;
- [0662] SEQ ID NOs: 196, 197, and 198, respectively;
- [0663] SEQ ID NOs: 202, 203, and 204, respectively; and
- [0664] SEQ ID NOs: 208, 209, and 210, respectively.
- [0665] In one embodiment,
- [0666] the VL and VH are selected from the group consisting of:
- [0667] a) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;
- [0668] b) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;
- [0669] c) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;
- [0670] d) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;
- [0671] e) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;
- [0672] f) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;
- [0673] g) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [0674] h) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

- [0675] i) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;
- [0676] j) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;
- [0677] k) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;
- [0678] l) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and
- [0679] m) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.
- [0680] In one embodiment,
- [0681] the VL comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and
- [0682] the VH comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.
- [0683] In one embodiment,
- [0684] the VL and VH are selected from the group consisting of:
- [0685] a) a VL comprising SEQ ID NO: 5 and a VH comprising SEQ ID NO: 6;
- [0686] b) a VL comprising SEQ ID NO: 9 and a VH comprising SEQ ID NO: 10;
- [0687] c) a VL comprising SEQ ID NO: 13 and a VH comprising SEQ ID NO: 14;
- [0688] d) a VL comprising SEQ ID NO: 17 and a VH comprising SEQ ID NO: 18;
- [0689] e) a VL comprising SEQ ID NO: 21 and a VH comprising SEQ ID NO: 22;
- [0690] f) a VL comprising SEQ ID NO: 25 and a VH comprising SEQ ID NO: 26;
- [0691] g) a VL comprising SEQ ID NO: 29 and a VH comprising SEQ ID NO: 30;
- [0692] h) a VL comprising SEQ ID NO: 33 and a VH comprising SEQ ID NO: 34;
- [0693] i) a VL comprising SEQ ID NO: 37 and a VH comprising SEQ ID NO: 38;
- [0694] j) a VL comprising SEQ ID NO: 41 and a VH comprising SEQ ID NO: 42;
- [0695] k) a VL comprising SEQ ID NO: 45 and a VH comprising SEQ ID NO: 46;
- [0696] l) a VL comprising SEQ ID NO: 49 and a VH comprising SEQ ID NO: 50; and
- [0697] m) a VL comprising SEQ ID NO: 53 and a VH comprising SEQ ID NO: 54.

[0698] In one embodiment, the antigen binding protein further comprises a second antigen binding portion that specifically binds a second antigen.

[0699] In one embodiment, the second antigen is a tumor associated antigen (TAA).

[0700] "Antibody constructs" according to the invention are fragments of full-length antibodies, such as VH, VHH, VL, (s)dAb, Fv, Fd, Fab, Fab', F(ab')₂ or "r IgG" ("half antibody"). Antibody constructs according to the invention may also be modified fragments of antibodies, also called antibody variants, such as scFv, di-scFv or bi(s)-scFv, scFv-Fc, scFv-zipper, scFab, Fab2, Fab3, diabodies, single chain diabodies, tandem diabodies (Tandab's), tandem di-scFv, tandem tri-scFv, „minibodies" exemplified by a structure which is as follows: (VH-VL-CH3)₂, (scFv-CH3)₂, ((scFv)₂-CH3 + CH3), ((scFv)₂-CH3) or (scFv-CH3-scFv)₂, multibodies such as triabodies or tetrabodies, and single domain antibodies such as nanobodies or single variable domain antibodies comprising merely one variable domain, which might be VHH, VH or VL, that specifically bind an antigen or epitope independently of other V regions or domains. Also, included within the definition of "antibody construct" are multispecific molecules that incorporate multiple types of antibodies and antibody constructs, for example, IgG-scFv which comprises a scFv linked to an IgG or an IgG-Fab, which comprises a Fab fragment linked to an IgG. In one embodiment, a scFv linked to an immunoglobulin heavy chain is called "a heavy chain fusion protein". In another embodiment, a VH-CH1 polypeptide linked to an immunoglobulin heavy chain is called "a heavy chain fusion protein".

[0701] A "binding domain", or "antigen binding domain", may typically comprise an antibody light chain variable region (VL) and an antibody heavy chain variable region (VH); however, it does not have to comprise both. Fd fragments, for example, have two VH regions and often retain some antigen-binding function of the intact antigen-binding domain. Additional examples for the format of antibody fragments, antibody variants or binding domains include (1) a Fab fragment, a monovalent fragment having the VL, VH, CL and CH1 domains; (2) a F(ab')₂ fragment, a bivalent fragment having two Fab fragments linked by a disulfide bridge at the hinge region; (3) an Fd fragment having the two VH and CH1 domains; (4) an Fv fragment having the VL and VH domains of a single arm of an antibody, (5) a dAb fragment (Ward et al., (1989) Nature 341 :544-546), which has a VH domain; (6) an isolated complementarity determining region (CDR), and (7) a single chain Fv (scFv), the latter being preferred (for example, derived from an scFv-library). Examples for embodiments of antibody constructs according to the invention are e.g. described in WO 00/006605, WO 2005/040220, WO 2008/1 19567, WO 2010/037838, WO 2013/026837, WO 2013/026833, US 2014/0308285, US 2014/0302037, W 02014/144722, WO 2014/151910, and WO 2015/048272.

[0702] Furthermore, the definition of the term "antibody construct" includes monovalent, bivalent and polyvalent / multivalent constructs and, thus, monospecific constructs, specifically binding to only one antigenic structure, as well as bispecific and multispecific constructs, which specifically bind more than one antigenic structure, e.g. two, three or more, through distinct binding domains.

Moreover, the definition of the term "antibody construct" includes molecules consisting of only one polypeptide chain as well as molecules consisting of more than one polypeptide chain, which chains can be either identical (homodimers, homotrimers or homo oligomers) or different (heterodimer, heterotrimer or heterooligomer). Examples for the above identified antibodies and variants or derivatives thereof are described inter alia in Harlow and Lane, *Antibodies a laboratory manual*, CSHL Press (1988) and *Using Antibodies: a laboratory manual*, CSHL Press (1999), Kontermann and Dubel, *Antibody Engineering*, Springer, 2nd ed. 2010 and Little, *Recombinant Antibodies for Immunotherapy*, Cambridge University Press 2009.

[0703] The antibody constructs of the present invention are preferably "in vitro generated antibody constructs". This term refers to an antibody construct according to the above definition where all or part of the variable region (e.g., at least one CDR) is generated in a non-immune cell selection, e.g., an in vitro phage display, protein chip or any other method in which candidate sequences can be tested for their ability to bind to an antigen. This term thus preferably excludes sequences generated solely by genomic rearrangement in an immune cell in an animal. A "recombinant antibody" is an antibody made through the use of recombinant DNA technology or genetic engineering.

[0704] In one aspect, the present invention is directed to a multispecific antibody construct comprising:

[0705] (i) a first antibody comprising two light chains and two heavy chains, wherein

[0706] the light chains comprise a first variable region (VL1) and a light chain constant region (CL);

[0707] the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and CH3 regions;

[0708] the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and

[0709] (ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,

[0710] wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and

[0711] wherein the first antibody specifically binds to and agonizes human CD40 (SEQ ID NO: 1) and the scFv specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2).

[0712] In one embodiment, the multispecific antibody construct specifically agonizes CD40 in a MSLN-dependent fashion.

[0713] In one embodiment, the multispecific antibody construct comprises mutations limiting Fc receptor binding and thus decreases Fc receptor-dependent CD40 agonism.

[0714] In one embodiment, the two light chains are identical and the two heavy chain fusion proteins are identical.

[0715] In one embodiment, the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0716] SEQ ID NOs: 58, 59, and 60, respectively;

[0717] SEQ ID NOs: 64, 65, and 66, respectively;

[0718] SEQ ID NOs: 70, 71, and 72, respectively;

[0719] SEQ ID NOs: 76, 77, and 78, respectively;

[0720] SEQ ID NOs: 82, 83, and 84, respectively;

[0721] SEQ ID NOs: 88, 89, and 90, respectively;

[0722] SEQ ID NOs: 94, 95, and 96, respectively;

[0723] SEQ ID NOs: 100, 101, and 102, respectively;

[0724] SEQ ID NOs: 106, 107, and 108, respectively;

[0725] SEQ ID NOs: 112, 113, and 114, respectively;

[0726] SEQ ID NOs: 118, 119, and 120, respectively;

[0727] SEQ ID NOs: 124, 125, and 126, respectively; and

[0728] SEQ ID NOs: 130, 131, and 132, respectively;

[0729] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0730] SEQ ID NOs: 136, 137, and 138, respectively;

[0731] SEQ ID NOs: 142, 143, and 144, respectively;

[0732] SEQ ID NOs: 148, 149, and 150, respectively;

[0733] SEQ ID NOs: 154, 155, and 156, respectively;

[0734] SEQ ID NOs: 160, 161, and 162, respectively;

[0735] SEQ ID NOs: 166, 167, and 168, respectively;

[0736] SEQ ID NOs: 172, 173, and 174, respectively;

[0737] SEQ ID NOs: 178, 179, and 180, respectively;

[0738] SEQ ID NOs: 184, 185, and 186, respectively;

[0739] SEQ ID NOs: 190, 191, and 192, respectively;

[0740] SEQ ID NOs: 196, 197, and 198, respectively;

[0741] SEQ ID NOs: 202, 203, and 204, respectively; and

[0742] SEQ ID NOs: 208, 209, and 210, respectively;

[0743] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0744] SEQ ID NOs: 230, 231, and 232, respectively;

[0745] SEQ ID NOs: 236, 237, and 238, respectively;

[0746] SEQ ID NOs: 242, 243, and 244, respectively; and

[0747] SEQ ID NOs: 248, 249, and 250, respectively;

[0748] and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0749] SEQ ID NOs: 254, 255, and 256, respectively;

[0750] SEQ ID NOs: 260, 261, and 262, respectively;

[0751] SEQ ID NOs: 266, 267, and 268, respectively; and

[0752] SEQ ID NOs: 272, 273, and 274, respectively.

[0753] In one embodiment, 1) the VL1 and VH1 are selected from the group consisting of:

[0754] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

[0755] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

[0756] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

[0757] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

[0758] e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

[0759] f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

[0760] g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

[0761] h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

[0762] i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[0763] j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[0764] k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[0765] l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[0766] m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;

[0767] and

[0768] 2) the VL2 and VH2 are selected from the group consisting of:

[0769] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

[0770] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

[0771] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

[0772] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

[0773] In one embodiment, the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

[0774] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

[0775] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[0776] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

[0777] In one embodiment,

[0778] 1) the VL1 and VH1 are selected from the group consisting of:

- [0779] a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;
- [0780] b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;
- [0781] c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;
- [0782] d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;
- [0783] e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;
- [0784] f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;
- [0785] g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;
- [0786] h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;
- [0787] i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;
- [0788] j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;
- [0789] k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;
- [0790] l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and
- [0791] m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;
- [0792] and
- [0793] 2) the VL2 and VH2 are selected from the group consisting of:
- [0794] a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;
- [0795] b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;
- [0796] c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and
- [0797] d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.
- [0798] In one embodiment,
- [0799] the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 286, 290, 294, 298, 302, 306, 310, 314, 318, 322, 326, 330, 336, 342, 346, 350, 354, 358, 362, 366, 370, 374, and 378; and
- [0800] the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID Nos: 285, 289, 293, 297, 301, 305, 309, 313, 317, 321, 325, 329, 333, 337, 341, 345, 349, 353, 357, 361, 365, 369, 373, and 377.
- [0801] In one embodiment, the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:
- [0802] SEQ ID NO: 286 and SEQ ID NO: 285, respectively;
- [0803] SEQ ID NO: 290 and SEQ ID NO: 289, respectively;
- [0804] SEQ ID NO: 294 and SEQ ID NO: 293, respectively;
- [0805] SEQ ID NO: 298 and SEQ ID NO: 297, respectively;
- [0806] SEQ ID NO: 302 and SEQ ID NO: 301, respectively;
- [0807] SEQ ID NO: 306 and SEQ ID NO: 305, respectively;
- [0808] SEQ ID NO: 310 and SEQ ID NO: 309, respectively;

- [0809] SEQ ID NO: 314 and SEQ ID NO: 313, respectively;
- [0810] SEQ ID NO: 318 and SEQ ID NO: 317, respectively;
- [0811] SEQ ID NO: 322 and SEQ ID NO: 321, respectively;
- [0812] SEQ ID NO: 326 and SEQ ID NO: 325, respectively;
- [0813] SEQ ID NO: 330 and SEQ ID NO: 329, respectively;
- [0814] SEQ ID NO: 334 and SEQ ID NO: 333, respectively;
- [0815] SEQ ID NO: 338 and SEQ ID NO: 337, respectively;
- [0816] SEQ ID NO: 342 and SEQ ID NO: 341, respectively;
- [0817] SEQ ID NO: 346 and SEQ ID NO: 345, respectively;
- [0818] SEQ ID NO: 350 and SEQ ID NO: 349, respectively;
- [0819] SEQ ID NO: 354 and SEQ ID NO: 353, respectively;
- [0820] SEQ ID NO: 358 and SEQ ID NO: 357, respectively;
- [0821] SEQ ID NO: 362 and SEQ ID NO: 361, respectively;
- [0822] SEQ ID NO: 366 and SEQ ID NO: 365, respectively;
- [0823] SEQ ID NO: 370 and SEQ ID NO: 369, respectively;
- [0824] SEQ ID NO: 374 and SEQ ID NO: 373, respectively; and
- [0825] SEQ ID NO: 378 and SEQ ID NO: 377, respectively.
- [0826] In another aspect, the present invention is directed to a multispecific antibody construct comprising:
- [0827] (i) a first antibody comprising two light chains and two heavy chains, wherein
- [0828] the light chains comprise a first variable region (VL1) and a light chain constant region (CL);
- [0829] the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and CH3 regions;
- [0830] the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and
- [0831] (ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,
- [0832] wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and
- [0833] wherein the first antibody specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2) and the scFv specifically binds to and agonizes human CD40 (SEQ ID NO: 1).
- [0834] In one embodiment, the multispecific antibody construct specifically agonizes CD40 in a MSLN-dependent fashion.

[0835] In one embodiment, the multispecific antibody construct comprises mutations limiting Fc receptor binding and thus decreases Fc receptor-dependent CD40 agonism.

[0836] In one embodiment, the two light chains are identical and the two heavy chain fusion proteins are identical.

[0837] In one embodiment:

[0838] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0839] SEQ ID NOs: 230, 231, and 232, respectively;

[0840] SEQ ID NOs: 236, 237, and 238, respectively;

[0841] SEQ ID NOs: 242, 243, and 244, respectively; and

[0842] SEQ ID NOs: 248, 249, and 250, respectively;

[0843] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0844] SEQ ID NOs: 254, 255, and 256, respectively;

[0845] SEQ ID NOs: 260, 261, and 262, respectively;

[0846] SEQ ID NOs: 266, 267, and 268, respectively; and

[0847] SEQ ID NOs: 272, 273, and 274, respectively;

[0848] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[0849] SEQ ID NOs: 58, 59, and 60, respectively;

[0850] SEQ ID NOs: 64, 65, and 66, respectively;

[0851] SEQ ID NOs: 70, 71, and 72, respectively;

[0852] SEQ ID NOs: 76, 77, and 78, respectively;

[0853] SEQ ID NOs: 82, 83, and 84, respectively;

[0854] SEQ ID NOs: 88, 89, and 90, respectively;

[0855] SEQ ID NOs: 94, 95, and 96, respectively;

[0856] SEQ ID NOs: 100, 101, and 102, respectively;

[0857] SEQ ID NOs: 106, 107, and 108, respectively;

[0858] SEQ ID NOs: 112, 113, and 114, respectively;

[0859] SEQ ID NOs: 118, 119, and 120, respectively;

[0860] SEQ ID NOs: 124, 125, and 126, respectively; and

[0861] SEQ ID NOs: 130, 131, and 132, respectively;

[0862] and

[0863] the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[0864] SEQ ID NOs: 136, 137, and 138, respectively;

[0865] SEQ ID NOs: 142, 143, and 144, respectively;

- [0866] SEQ ID NOs: 148, 149, and 150, respectively;
- [0867] SEQ ID NOs: 154, 155, and 156, respectively;
- [0868] SEQ ID NOs: 160, 161, and 162, respectively;
- [0869] SEQ ID NOs: 166, 167, and 168, respectively;
- [0870] SEQ ID NOs: 172, 173, and 174, respectively;
- [0871] SEQ ID NOs: 178, 179, and 180, respectively;
- [0872] SEQ ID NOs: 184, 185, and 186, respectively;
- [0873] SEQ ID NOs: 190, 191, and 192, respectively;
- [0874] SEQ ID NOs: 196, 197, and 198, respectively;
- [0875] SEQ ID NOs: 202, 203, and 204, respectively; and
- [0876] SEQ ID NOs: 208, 209, and 210, respectively.
- [0877] In one embodiment,
- [0878] 1) the VL1 and VH1 are selected from the group consisting of:
- [0879] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;
- [0880] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;
- [0881] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and
- [0882] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;
- [0883] and
- [0884] 2) the VL2 and VH2 are selected from the group consisting of:
- [0885] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;
- [0886] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;
- [0887] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

- [0888] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;
- [0889] e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;
- [0890] f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;
- [0891] g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [0892] h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;
- [0893] i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;
- [0894] j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;
- [0895] k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;
- [0896] l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and
- [0897] m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.
- [0898] In one embodiment:
- [0899] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;
- [0900] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

[0901] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

[0902] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

[0903] In one embodiment:

[0904] 1) the VL1 and VH1 are selected from the group consisting of:

[0905] a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;

[0906] b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;

[0907] c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and

[0908] d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;

[0909] and

[0910] 2) the VL2 and VH2 are selected from the group consisting of:

[0911] a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;

[0912] b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;

[0913] c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;

[0914] d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;

[0915] e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;

[0916] f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;

[0917] g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;

[0918] h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;

[0919] i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;

[0920] j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;

[0921] k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;

[0922] l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and

[0923] m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.

[0924] In one embodiment, the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

[0925] In one embodiment, the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

[0926] In one embodiment, the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.

[0927] In one embodiment, the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.

[0928] In one embodiment, the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.

[0929] In one embodiment,

[0930] the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 382, 386, 390, 394, 398, 402, 406, 410, 414, 418, 422, 426, 430, 434, 438, 442, 446, and 450; and

[0931] the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID NOs: 381, 385, 389, 393, 397, 401, 405, 409, 413, 417, 421, 425, 429, 433, 437, 441, 445, and 449.

[0932] In one embodiment, the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:

[0933] SEQ ID NO: 382 and SEQ ID NO: 381, respectively;

[0934] SEQ ID NO: 386 and SEQ ID NO: 385, respectively;

[0935] SEQ ID NO: 390 and SEQ ID NO: 389, respectively;

[0936] SEQ ID NO: 394 and SEQ ID NO: 393, respectively;

[0937] SEQ ID NO: 398 and SEQ ID NO: 397, respectively;

[0938] SEQ ID NO: 402 and SEQ ID NO: 401, respectively;

[0939] SEQ ID NO: 406 and SEQ ID NO: 405, respectively;

[0940] SEQ ID NO: 410 and SEQ ID NO: 409, respectively;

[0941] SEQ ID NO: 414 and SEQ ID NO: 413, respectively;

[0942] SEQ ID NO: 418 and SEQ ID NO: 417, respectively;

[0943] SEQ ID NO: 422 and SEQ ID NO: 421, respectively;

[0944] SEQ ID NO: 426 and SEQ ID NO: 425, respectively;

[0945] SEQ ID NO: 430 and SEQ ID NO: 429, respectively;

[0946] SEQ ID NO: 434 and SEQ ID NO: 433, respectively;

[0947] SEQ ID NO: 438 and SEQ ID NO: 437, respectively;

[0948] SEQ ID NO: 442 and SEQ ID NO: 441, respectively;

[0949] SEQ ID NO: 446 and SEQ ID NO: 445, respectively; and

[0950] SEQ ID NO: 450 and SEQ ID NO: 449, respectively.

[0951] In one embodiment, the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

[0952] In one embodiment, the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

[0953] In one embodiment, the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.

[0954] In one embodiment, the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.

[0955]

[0956] In one embodiment, the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.

[0957] The term "monoclonal antibody" (mAb) or monoclonal antibody construct as used herein refers to an antibody obtained from a population of substantially homogeneous antibodies, i.e., the individual antibodies comprising the population are identical except for possible naturally occurring mutations and/or post-translation modifications (e.g., isomerizations, amidations) that may be present in minor amounts. Monoclonal antibodies are highly specific, being directed against a single antigenic site or determinant on the antigen, in contrast to conventional (polyclonal) antibody preparations which typically include different antibodies directed against different determinants (or epitopes). In addition to their specificity, the monoclonal antibodies are advantageous in that they are synthesized by the hybridoma culture, hence uncontaminated by other immunoglobulins. The modifier "monoclonal" indicates the character of the antibody as being obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method.

[0958] For the preparation of monoclonal antibodies, any technique providing antibodies produced by continuous cell line cultures can be used. For example, monoclonal antibodies to be used may be made by the hybridoma method first described by Koehler et al., *Nature*, 256: 495 (1975), or may be made by recombinant DNA methods (see, e.g., U.S. Patent No. 4,816,567). Examples for further techniques to produce human monoclonal antibodies include the trioma technique, the human B-cell hybridoma technique (Kozbor, *Immunology Today* 4 (1983), 72) and the EBV- hybridoma technique (Cole et al., *Monoclonal Antibodies and Cancer Therapy*, Alan R. Liss, Inc. (1985), 77-96).

[0959] Hybridomas can then be screened using standard methods, such as enzyme-linked immunosorbent assay (ELISA) and surface plasmon resonance (BIAcore™) analysis, to identify one or more hybridomas that produce an antibody that specifically binds with a specified antigen. Any form of the relevant antigen may be used as the immunogen, e.g., recombinant antigen, naturally occurring forms, any variants or fragments thereof, as well as an antigenic peptide thereof. Surface plasmon resonance as employed in the BIAcore system can be used to increase the efficiency of phage antibodies which bind to an epitope of a target antigen, such as MSLN or CD40 (Schier, *Human Antibodies Hybridomas* 7 (1996), 97-105; Malmberg, *J. Immunol. Methods* 183 (1995), 7-13).

[0960] Another exemplary method of making monoclonal antibodies includes screening protein expression libraries, e.g., phage display or ribosome display libraries. Phage display is described, for example, in Ladner et al., U.S. Patent No. 5,223,409; Smith (1985) *Science* 228:1315-1317, Clackson et al., *Nature*, 352: 624-628 (1991) and Marks et al., *J. Mol. Biol.*, 222: 581 -597 (1991).

[0961] In addition to the use of display libraries, the relevant antigen can be used to immunize a non-human animal, e.g., a rodent (such as a mouse, hamster, rabbit or rat). In one embodiment, the non-human animal includes at least a part of a human immunoglobulin gene. For example, it is possible to

engineer mouse strains deficient in mouse antibody production with large fragments of the human Ig (immunoglobulin) loci. Using the hybridoma technology, antigen-specific monoclonal antibodies derived from the genes with the desired specificity may be produced and selected. See, e.g., XENOMOUSE™, Green et al. (1994) *Nature Genetics* 7:13-21, US 2003- 0070185, WO 96/34096, and WO 96/33735.

[0962] A monoclonal antibody can also be obtained from a non-human animal, and then modified, e.g., humanized, deimmunized, rendered chimeric etc., using recombinant DNA techniques known in the art. Examples of modified antibody constructs include humanized variants of non-human antibodies, "affinity matured" antibodies (see, e.g. Hawkins et al. *J. Mol. Biol.* 254, 889-896 (1992) and Lowman et al., *Biochemistry* 30, 10832- 10837 (1991)) and antibody mutants with altered effector function(s) (see, e.g., US Patent 5,648,260, Kontermann and Dubel (2010), loc. cit. and Little (2009), loc. cit).

[0963] In immunology, affinity maturation is the process by which B cells produce antibodies with increased affinity for antigen during the course of an immune response. With repeated exposures to the same antigen, a host will produce antibodies of successively greater affinities. Like the natural prototype, the in vitro affinity maturation is based on the principles of mutation and selection. The in vitro affinity maturation has successfully been used to optimize antibodies, antibody constructs, and antibody fragments. Random mutations inside the CDRs are introduced using radiation, chemical mutagens or error-prone PCR. In addition, the genetical diversity can be increased by chain shuffling. Two or three rounds of mutation and selection using display methods like phage display usually results in antibody fragments with affinities in the low nanomolar range.

[0964] A preferred type of an amino acid substitutional variation of the antibody constructs involves substituting one or more hypervariable region residues of a parent antibody (e. g. a humanized or human antibody). Generally, the resulting variant(s) selected for further development will have improved biological properties relative to the parent antibody from which they are generated. A convenient way for generating such substitutional variants involves affinity maturation using phage display. Briefly, several hypervariable region sites (e. g. 6-7 sites) are mutated to generate all possible amino acid substitutions at each site. The antibody variants thus generated are displayed in a monovalent fashion from filamentous phage particles as fusions to the gene III product of M13 packaged within each particle. The phage-displayed variants are then screened for their biological activity (e. g. binding affinity) as herein disclosed. In order to identify candidate hypervariable region sites for modification, alanine scanning mutagenesis can be performed to identify hypervariable region residues contributing significantly to antigen binding. Alternatively, or additionally, it may be beneficial to analyze a crystal structure of the antigen-antibody complex to identify contact points between the binding domain and, e.g., human MSLN. Such contact residues and neighboring residues are candidates for substitution according to the techniques elaborated herein. Once such variants are

generated, the panel of variants is subjected to screening as described herein and antibodies with superior properties in one or more relevant assays may be selected for further development.

[0965] The monoclonal antibodies and antibody constructs of the present invention specifically include "chimeric" antibodies (immunoglobulins) in which a portion of the heavy and/or light chain is identical with or homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular antibody class or subclass, while the remainder of the chain(s) is/are identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity (U.S. Patent No. 4,816,567; Morrison et al., Proc. Natl. Acad. Sci. USA, 81 : 6851 -6855 (1984)). Chimeric antibodies of interest herein include "primitized" antibodies comprising variable domain antigen- binding sequences derived from a non-human primate (e.g., Old World Monkey, Ape etc.) and human constant region sequences. A variety of approaches for making chimeric antibodies have been described. See e.g., Morrison et al., Proc. Natl. Acad. Sci. U.S.A. 81 :6851 , 1985; Takeda et al., Nature 314:452, 1985, Cabilly et al., U.S. Patent No. 4,816,567; Boss et al., U.S. Patent No. 4,816,397; Tanaguchi et al., EP 0171496; EP 0173494; and GB 2177096.

[0966] An antibody, antibody construct, antibody fragment or antibody variant may also be modified by specific deletion of human T cell epitopes (a method called "deimmunization") by the methods disclosed for example in WO 98/52976 or WO 00/34317. Briefly, the heavy and light chain variable domains of an antibody can be analyzed for peptides that bind to MHC class II; these peptides represent potential T cell epitopes (as defined in WO 98/52976 and WO 00/34317). For detection of potential T cell epitopes, a computer modeling approach termed "peptide threading" can be applied, and in addition a database of human MHC class II binding peptides can be searched for motifs present in the VH and VL sequences, as described in WO 98/52976 and WO 00/34317. These motifs bind to any of the 18 major MHC class II DR allotypes, and thus constitute potential T cell epitopes. Potential T cell epitopes detected can be eliminated by substituting small numbers of amino acid residues in the variable domains, or preferably, by single amino acid substitutions. Typically, conservative substitutions are made. Often, but not exclusively, an amino acid common to a position in human germline antibody sequences may be used. Human germline sequences are disclosed e.g. in Tomlinson, et al. (1992) J. Mol. Biol. 227:776-798; Cook, G.P. et al. (1995) Immunol. Today Vol. 16 (5): 237-242; and Tomlinson et al. (1995) EMBO J. 14: 14:4628-4638. The V BASE directory provides a comprehensive directory of human immunoglobulin variable region sequences (compiled by Tomlinson, L.A. et al. MRC Centre for Protein Engineering, Cambridge, UK). These sequences can be used as a source of human sequence, e.g., for framework regions and CDRs. Consensus human framework regions can also be used, for example as described in US Patent No. 6,300,064.

[0967] "Humanized" antibodies, antibody constructs, variants or fragments thereof (such as Fv, Fab, Fab', F(ab')₂ or other antigen-binding subsequences of antibodies) are antibodies or immunoglobulins of mostly human sequences, which contain (a) minimal sequence(s) derived from non-human immunoglobulin. For the most part, humanized antibodies are human immunoglobulins (recipient antibody) in which residues from a hypervariable region (also CDR) of the recipient are replaced by residues from a hypervariable region of a non-human (e.g., rodent) species (donor antibody) such as mouse, rat, hamster or rabbit having the desired specificity, affinity, and capacity. In some instances, Fv framework region (FR) residues of the human immunoglobulin are replaced by corresponding non-human residues. Furthermore, "humanized antibodies" as used herein may also comprise residues which are found neither in the recipient antibody nor the donor antibody. These modifications are made to further refine and optimize antibody performance. The humanized antibody may also comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin. For further details, see Jones et al., *Nature*, 321 : 522-525 (1986); Reichmann et al., *Nature*, 332: 323-329 (1988); and Presta, *Curr. Op. Struct. Biol.*, 2: 593-596 (1992).

[0968] Humanized antibodies or fragments thereof can be generated by replacing sequences of the Fv variable domain that are not directly involved in antigen binding with equivalent sequences from human Fv variable domains. Exemplary methods for generating humanized antibodies or fragments thereof are provided by Morrison (1985) *Science* 229:1202-1207; by Oi et al. (1986) *BioTechniques* 4:214; and by US 5,585,089; US 5,693,761 ; US 5,693,762; US 5,859,205; and US 6,407,213. Those methods include isolating, manipulating, and expressing the nucleic acid sequences that encode all or part of immunoglobulin Fv variable domains from at least one of a heavy or light chain. Such nucleic acids may be obtained from a hybridoma producing an antibody against a predetermined target, as described above, as well as from other sources. The recombinant DNA encoding the humanized antibody molecule can then be cloned into an appropriate expression vector.

[0969] Humanized antibodies may also be produced using transgenic animals such as mice that express human heavy and light chain genes, but are incapable of expressing the endogenous mouse immunoglobulin heavy and light chain genes. Winter describes an exemplary CDR grafting method that may be used to prepare the humanized antibodies described herein (U.S. Patent No. 5,225,539). All of the CDRs of a particular human antibody may be replaced with at least a portion of a non-human CDR, or only some of the CDRs may be replaced with non-human CDRs. It is only necessary to replace the number of CDRs required for binding of the humanized antibody to a predetermined antigen.

[0970] A humanized antibody can be optimized by the introduction of conservative substitutions, consensus sequence substitutions, germline substitutions and/or back mutations. Such altered immunoglobulin molecules can be made by any of several techniques known in the art, (e.g., Teng et

al., Proc. Natl. Acad. Sci. U.S.A., 80: 7308-7312, 1983; Kozbor et al., Immunology Today, 4: 7279, 1983; Olsson et al., Meth. Enzymol., 92: 3-16, 1982, and EP 239 400).

[0971] The term "human antibody", "human antibody construct" and "human binding domain" includes antibodies, antibody constructs and binding domains having antibody regions such as variable and constant regions or domains which correspond substantially to human germline immunoglobulin sequences known in the art, including, for example, those described by Kabat et al. (1991) (loc. cit.). The human antibodies, antibody constructs or binding domains of the invention may include amino acid residues not encoded by human germline immunoglobulin sequences (e.g., mutations introduced by random or site-specific mutagenesis in vitro or by somatic mutation in vivo), for example in the CDRs, and in particular, in CDR3. The human antibodies, antibody constructs or binding domains can have at least one, two, three, four, five, or more positions replaced with an amino acid residue that is not encoded by the human germline immunoglobulin sequence. The definition of human antibodies, antibody constructs and binding domains as used herein also contemplates fully human antibodies, which include only non-artificially and/or genetically altered human sequences of antibodies as those can be derived by using technologies or systems such as the Xenomouse.

[0972] In some embodiments, the antibody constructs of the invention are "isolated" or "substantially pure" antibody constructs. "Isolated" or "substantially pure", when used to describe the antibody constructs disclosed herein, means an antibody construct that has been identified, separated and/or recovered from a component of its production environment. Preferably, the antibody construct is free or substantially free of association with all other components from its production environment. Contaminant components of its production environment, such as that resulting from recombinant transfected cells, are materials that would typically interfere with diagnostic or therapeutic uses for the polypeptide, and may include enzymes, hormones, and other proteinaceous or non-proteinaceous solutes. The antibody constructs may e.g. constitute at least about 5%, or at least about 50% by weight of the total protein in a given sample. It is understood that the isolated protein may constitute from 5% to 99.9% by weight of the total protein content, depending on the circumstances. The polypeptide may be made at a significantly higher concentration through the use of an inducible promoter or high expression promoter, such that it is made at increased concentration levels. The definition includes the production of an antibody construct in a wide variety of organisms and/or host cells that are known in the art. In preferred embodiments, the antibody construct will be purified (1) to a degree sufficient to obtain at least 15 residues of N-terminal or internal amino acid sequence by use of a spinning cup sequenator, or (2) to homogeneity by SDS-PAGE under non-reducing or reducing conditions using Coomassie blue or, preferably, silver stain. Ordinarily, however, an isolated antibody construct will be prepared by at least one purification step.

[0973] The term "binding domain" characterizes in connection with the present invention a domain which (specifically) binds to / interacts with / recognizes a given target epitope or a given target site on the target molecules (antigens), here: MSLN and CD40, respectively. The structure and function of the one binding domain (recognizing MSLN), and preferably also the structure and/or function of another binding domain (recognizing CD40), is/are based on the structure and/or function of an antibody, e.g. of a full-length or whole immunoglobulin molecule. According to the invention, the binding domains are characterized by the presence of three light chain CDRs (i.e. CDRL1, CDRL2 and CDRL3 of the VL region) and/or three heavy chain CDRs (i.e. CDRH1, CDRH2 and CDRH3 of the VH region). It is envisaged that the binding domains are produced by or obtainable by phage-display or library screening methods rather than by grafting CDR sequences from a pre-existing (monoclonal) antibody into a scaffold. According to the present invention, binding domains are in the form of one or more polypeptides. Such polypeptides may include proteinaceous parts and non-proteinaceous parts (e.g. chemical linkers or chemical cross-linking agents such as glutaraldehyde). Proteins (including fragments thereof, preferably biologically active fragments, and peptides, usually having less than 30 amino acids) comprise two or more amino acids coupled to each other via a covalent peptide bond (resulting in a chain of amino acids). The term "polypeptide" as used herein describes a group of molecules, which usually consist of more than 30 amino acids. Polypeptides may further form multimers such as dimers, trimers and higher oligomers, i.e., consisting of more than one polypeptide molecule. Polypeptide molecules forming such dimers, trimers etc. may be identical or non-identical. The corresponding higher order structures of such multimers are, consequently, termed homo- or heterodimers, homo- or heterotrimers etc. An example for a heteromultimer is an antibody molecule, which, in its naturally occurring form, consists of two identical light polypeptide chains and two identical heavy polypeptide chains. The terms "peptide", "polypeptide" and "protein" also refer to naturally modified peptides / polypeptides / proteins wherein the modification is effected e.g. by post-translational modifications like glycosylation, acetylation, phosphorylation and the like. A "peptide", "polypeptide" or "protein" when referred to herein may also be chemically modified such as pegylated. Such modifications are well known in the art and described herein below.

[0974] Preferably the binding domain which binds to MSLN and/or the binding domain which binds to CD40 is/are human binding domains. Antibodies and antibody constructs comprising at least one human binding domain avoid some of the problems associated with antibodies or antibody constructs that possess non-human such as rodent (e.g. murine, rat, hamster or rabbit) variable and/or constant regions. The presence of such rodent derived proteins can lead to the rapid clearance of the antibodies or antibody constructs or can lead to the generation of an immune response against the antibody or antibody construct by a patient. In order to avoid the use of rodent derived antibodies or antibody constructs, human or fully human antibodies / antibody constructs can be generated through the introduction of human antibody function into a rodent so that the rodent produces fully human antibodies.

[0975] The ability to clone and reconstruct megabase-sized human loci in YACs and to introduce them into the mouse germline provides a powerful approach to elucidating the functional components of very large or crudely mapped loci as well as generating useful models of human disease. Furthermore, the use of such technology for substitution of mouse loci with their human equivalents could provide unique insights into the expression and regulation of human gene products during development, their communication with other systems, and their involvement in disease induction and progression.

[0976] An important practical application of such a strategy is the "humanization" of the mouse humoral immune system. Introduction of human immunoglobulin (Ig) loci into mice in which the endogenous Ig genes have been inactivated offers the opportunity to study the mechanisms underlying programmed expression and assembly of antibodies as well as their role in B-cell development. Furthermore, such a strategy could provide an ideal source for production of fully human monoclonal antibodies (mAbs) - an important milestone towards fulfilling the promise of antibody therapy in human disease. Fully human antibodies or antibody constructs are expected to minimize the immunogenic and allergic responses intrinsic to mouse or mouse-derivatized mAbs and thus to increase the efficacy and safety of the administered antibodies / antibody constructs. The use of fully human antibodies or antibody constructs can be expected to provide a substantial advantage in the treatment of chronic and recurring human diseases, such as inflammation, autoimmunity, and cancer, which require repeated compound administrations.

[0977] One approach towards this goal was to engineer mouse strains deficient in mouse antibody production with large fragments of the human Ig loci in anticipation that such mice would produce a large repertoire of human antibodies in the absence of mouse antibodies. Large human Ig fragments would preserve the large variable gene diversity as well as the proper regulation of antibody production and expression. By exploiting the mouse machinery for antibody diversification and selection and the lack of immunological tolerance to human proteins, the reproduced human antibody repertoire in these mouse strains should yield high affinity antibodies against any antigen of interest, including human antigens. Using the hybridoma technology, antigen-specific human mAbs with the desired specificity could be readily produced and selected. This general strategy was demonstrated in connection with the generation of the first XenoMouse mouse strains (see Green et al. *Nature Genetics* 7:13-21 (1994)). The XenoMouse strains were engineered with yeast artificial chromosomes (YACs) containing 245 kb and 190 kb-sized germline configuration fragments of the human heavy chain locus and kappa light chain locus, respectively, which contained core variable and constant region sequences. The human Ig containing YACs proved to be compatible with the mouse system for both rearrangement and expression of antibodies and were capable of substituting for the inactivated mouse Ig genes. This was demonstrated by their ability to induce B cell development, to produce an adult-like human repertoire of fully human antibodies, and to generate antigen-specific human mAbs.

These results also suggested that introduction of larger portions of the human Ig loci containing greater numbers of V genes, additional regulatory elements, and human Ig constant regions might recapitulate substantially the full repertoire that is characteristic of the human humoral response to infection and immunization. The work of Green et al. was recently extended to the introduction of greater than approximately 80% of the human antibody repertoire through introduction of megabase sized, germline configuration YAC fragments of the human heavy chain loci and kappa light chain loci, respectively. See Mendez et al. *Nature Genetics* 15:146-156 (1997) and U.S. patent application Ser. No. 08/759,620.

[0978] The production of the XenoMouse mice is further discussed and delineated in U.S. patent applications Ser. No. 07/466,008, Ser. No. 07/610,515, Ser. No. 07/919,297, Ser. No. 07/922,649, Ser. No. 08/031,801, Ser. No. 08/112,848, Ser. No. 08/234,145, Ser. No. 08/376,279, Ser. No. 08/430,938, Ser. No. 08/464,584, Ser. No. 08/464,582, Ser. No. 08/463,191, Ser. No. 08/462,837, Ser. No. 08/486,853, Ser. No. 08/486,857, Ser. No. 08/486,859, Ser. No. 08/462,513, Ser. No. 08/724,752, and Ser. No. 08/759,620; and U.S. Pat. Nos. 6,162,963; 6,150,584; 6,114,598; 6,075,181, and 5,939,598 and Japanese Patent Nos. 3 068 180 B2, 3 068 506 B2, and 3 068 507 B2. See also Mendez et al. *Nature Genetics* 15:146-156 (1997) and Green and Jakobovits *J. Exp. Med.* 188:483-495 (1998), EP 0 463 151 B1, WO 94/02602, WO 96/34096, WO 98/24893, WO 00/76310, and WO 03/47336.

[0979] In an alternative approach, others, including GenPharm International, Inc., have utilized a "minilocus" approach. In the minilocus approach, an exogenous Ig locus is mimicked through the inclusion of pieces (individual genes) from the Ig locus. Thus, one or more VH genes, one or more DH genes, one or more JH genes, a mu constant region, and a second constant region (preferably a gamma constant region) are formed into a construct for insertion into an animal. This approach is described in U.S. Pat. No. 5,545,807 to Surani et al. and U.S. Pat. Nos. 5,545,806; 5,625,825; 5,625,126; 5,633,425; 5,661,016; 5,770,429; 5,789,650; 5,814,318; 5,877,397; 5,874,299; and 6,255,458 each to Lonberg and Kay, U.S. Pat. Nos. 5,591,669 and 6,023,010 to Krimpenfort and Berns, U.S. Pat. Nos. 5,612,205; 5,721,367; and 5,789,215 to Berns et al., and U.S. Pat. No. 5,643,763 to Choi and Dunn, and GenPharm International U.S. patent application Ser. No. 07/574,748, Ser. No. 07/575,962, Ser. No. 07/810,279, Ser. No. 07/853,408, Ser. No. 07/904,068, Ser. No. 07/990,860, Ser. No. 08/053,131, Ser. No. 08/096,762, Ser. No. 08/155,301, Ser. No. 08/161,739, Ser. No. 08/165,699, Ser. No. 08/209,741. See also EP 0 546 073 B1, WO 92/03918, WO 92/22645, WO 92/22647, WO 92/22670, WO 93/12227, WO 94/00569, WO 94/25585, WO 96/14436, WO 97/13852, and WO 98/24884 and U.S. Pat. No. 5,981,175. See further Taylor et al. (1992), Chen et al. (1993), Tuaillon et al. (1993), Choi et al. (1993), Lonberg et al. (1994), Taylor et al. (1994), and Tuaillon et al. (1995), Fishwild et al. (1996).

[0980] Kirin has also demonstrated the generation of human antibodies from mice in which, through microcell fusion, large pieces of chromosomes, or entire chromosomes, have been introduced. See European Patent Application Nos. 773 288 and 843 961 . Xenerex Biosciences is developing a technology for the potential generation of human antibodies. In this technology, SCID mice are reconstituted with human lymphatic cells, e.g., B and/or T cells. Mice are then immunized with an antigen and can generate an immune response against the antigen. See U.S. Pat. Nos. 5,476,996; 5,698,767; and 5,958,765.

[0981] Human anti-mouse antibody (HAMA) responses have led the industry to prepare chimeric or otherwise humanized antibodies. It is however expected that certain human anti-chimeric antibody (HACA) responses will be observed, particularly in chronic or multi-dose utilizations of the antibody. Thus, it would be desirable to provide antibody constructs comprising a human binding domain against MSLN and a human binding domain against CD40 in order to vitiate concerns and/or effects of HAMA or HACA response.

[0982] The terms "(specifically) binds to", "(specifically) recognizes", "is (specifically) directed to", and "(specifically) reacts with" mean in accordance with this invention that a binding domain interacts or specifically interacts with a given epitope or a given target site on the target molecules (antigens), here: MSLN and CD40, respectively.

[0983] The term "epitope" refers to a site on an antigen to which a binding domain, such as an antibody or immunoglobulin, or a derivative, fragment or variant of an antibody or an immunoglobulin, specifically binds. An "epitope" is antigenic and thus the term epitope is sometimes also referred to herein as "antigenic structure" or "antigenic determinant". Thus, the binding domain is an "antigen interaction site". Said binding/interaction is also understood to define a "specific recognition".

[0984] "Epitopes" can be formed both by contiguous amino acids or non-contiguous amino acids juxtaposed by tertiary folding of a protein. A "linear epitope" is an epitope where an amino acid primary sequence comprises the recognized epitope. A linear epitope typically includes at least 3 or at least 4, and more usually, at least 5 or at least 6 or at least 7, for example, about 8 to about 10 amino acids in a unique sequence.

[0985] A "conformational epitope", in contrast to a linear epitope, is an epitope wherein the primary sequence of the amino acids comprising the epitope is not the sole defining component of the epitope recognized (e.g., an epitope wherein the primary sequence of amino acids is not necessarily recognized by the binding domain). Typically a conformational epitope comprises an increased number of amino acids relative to a linear epitope. With regard to recognition of conformational epitopes, the binding domain recognizes a three-dimensional structure of the antigen, preferably a peptide or protein or fragment thereof (in the context of the present invention, the antigenic structure for one of the binding domains is comprised within the MSLN protein). For example, when a protein

molecule folds to form a three-dimensional structure, certain amino acids and/or the polypeptide backbone forming the conformational epitope become juxtaposed enabling the antibody to recognize the epitope. Methods of determining the conformation of epitopes include, but are not limited to, x-ray crystallography, two-dimensional nuclear magnetic resonance (2D-NMR) spectroscopy and site-directed spin labelling and electron paramagnetic resonance (EPR) spectroscopy.

[0986] A method for epitope mapping is described in the following: When a region (a contiguous amino acid stretch) in the human MSLN protein is exchanged / replaced with its corresponding region of a non-human and non-primate MSLN antigen (e.g., mouse MSLN, but others like chicken, rat, hamster, rabbit etc. might also be conceivable), a decrease in the binding of the binding domain is expected to occur, unless the binding domain is cross-reactive for the non-human, non-primate MSLN used. Said decrease is preferably at least 10%, 20%, 30%, 40%, or 50%; more preferably at least 60%, 70%, or 80%, and most preferably 90%, 95% or even 100% in comparison to the binding to the respective region in the human MSLN protein, whereby binding to the respective region in the human MSLN protein is set to be 100%. It is envisaged that the aforementioned human MSLN / non-human MSLN chimeras are expressed in CHO cells. The human MSLN / non-human MSLN chimeras may also be fused with a transmembrane domain and/or cytoplasmic domain of a different membrane-bound protein such as EpCAM, although such technique was not necessary for the method described in Examples 1 and 2.

[0987] In an alternative or additional method for epitope mapping, several truncated versions of the human MSLN extracellular domain can be generated in order to determine a specific region that is recognized by a binding domain. In these truncated versions, the different extracellular MSLN domains / sub-domains or regions are stepwise deleted, starting from the N-terminus. The truncated MSLN versions that may be expressed in CHO cells. It is also envisaged that the truncated MSLN versions may be fused with a transmembrane domain and/or cytoplasmic domain of a different membrane-bound protein such as EpCAM. It is also envisaged that the truncated MSLN versions may encompass a signal peptide domain at their N-terminus, for example a signal peptide derived from mouse IgG heavy chain signal peptide. It is furtherore envisaged that the truncated MSLN versions may encompass a v5 domain at their N-terminus (following the signal peptide) which allows verifying their correct expression on the cell surface. A decrease or a loss of binding is expected to occur with those truncated MSLN versions which do not encompass any more the MSLN region that is recognized by the binding domain. The decrease of binding is preferably at least 10%, 20%, 30%, 40%, 50%; more preferably at least 60%, 70%, 80%, and most preferably 90%, 95% or even 100%, whereby binding to the entire human MSLN protein (or its extracellular region or domain) is set to be 100%.

[0988] A further method to determine the contribution of a specific residue of a target antigen to the recognition by an antibody construct or binding domain is alanine scanning (see e.g. Morrison KL &

Weiss GA. *Cur Opin Chem Biol.* 2001 Jun;5(3):302-7), where each residue to be analyzed is replaced by alanine, e.g. via site-directed mutagenesis. Alanine is used because of its non- bulky, chemically inert, methyl functional group that nevertheless mimics the secondary structure references that many of the other amino acids possess. Sometimes bulky amino acids such as valine or leucine can be used in cases where conservation of the size of mutated residues is desired. Alanine scanning is a mature technology which has been used for a long period of time.

[0989] The interaction between the binding domain and the epitope or the region comprising the epitope implies that a binding domain exhibits appreciable affinity for the epitope / the region comprising the epitope on a particular protein or antigen (here: MSLN and CD40, respectively) and, generally, does not exhibit significant reactivity with proteins or antigens other than MSLN or CD40. "Appreciable affinity" includes binding with an affinity of about 10^{-6} M (KD) or stronger. Preferably, binding is considered specific when the binding affinity is about 10^{-12} to 10^{-8} M, 10^{-12} to 10^{-9} M, 10^{-12} to 10^{-10} M, 10^{-11} to 10^{-8} M, preferably of about 10^{-11} to 10^{-9} M. Whether a binding domain specifically reacts with or binds to a target can be tested readily by, inter alia, comparing the reaction of said binding domain with a target protein or antigen with the reaction of said binding domain with proteins or antigens other than MSLN or CD40. Preferably, a binding domain of the invention does not essentially or substantially bind to proteins or antigens other than MSLN or CD40 (e.g., one binding domain is not capable of binding to proteins other than MSLN and the other binding domain is not capable of binding to proteins other than CD40).

[0990] The term "does not essentially / substantially bind" or "is not capable of binding" means that a binding domain of the present invention does not bind a protein or antigen other than MSLN or CD40, i.e., does not show reactivity of more than 30%, preferably not more than 20%, more preferably not more than 10%, particularly preferably not more than 9%, 8%, 7%, 6% or 5% with proteins or antigens other than MSLN or CD40, whereby binding to MSLN or CD40, respectively, is set to be 100%.

[0991] Specific binding is believed to be effected by specific motifs in the amino acid sequence of the binding domain and the antigen. Thus, binding is achieved as a result of their primary, secondary and/or tertiary structure as well as the result of secondary modifications of said structures. The specific interaction of the antigen-interaction-site with its specific antigen may result in a simple binding of said site to the antigen. Moreover, the specific interaction of the antigen-interaction-site with its specific antigen may alternatively or additionally result in the initiation of a signal, e.g. due to the induction of a change of the conformation of the antigen, an oligomerization of the antigen, etc.

[0992] The Fc region of an antibody interacts with a number of Fc receptors and ligands, imparting an array of important functional capabilities referred to as effector functions. For IgG the Fc region comprises Ig domains CH2 and CH3. An important family of Fc receptors for the IgG isotype are the Fc gamma receptors (FcγRs). These receptors mediate communication between antibodies and the

cellular arm of the immune system. In humans this protein family includes Fc γ RI (CD64), including isoforms Fc γ RIa, Fc γ RIb, and Fc γ RIc; Fc γ RII (CD32), including isoforms Fc γ RIIa (including allotypes H131 and R131), Fc γ RIIb (including Fc γ RIIb-1 and Fc γ RIIb-2), and Fc γ RIIc; and Fc γ RIII (CD16), including isoforms Fc γ RIIIa (including allotypes V158 and F158) and Fc γ RIIIb (including allotypes Fc γ RIIIb-NA1 and Fc γ RIIIb-NA2) (Jefferis et al., 2002, *Immunol Lett* 82:57-65). These receptors typically have an extracellular domain that mediates binding to Fc, a membrane spanning region, and an intracellular domain that may mediate some signaling event within the cell. These receptors are expressed in a variety of immune cells including monocytes, macrophages, neutrophils, dendritic cells, eosinophils, mast cells, platelets, B cells, large granular lymphocytes, Langerhans' cells, natural killer (NK) cells, and $\gamma\delta$ T cells. Formation of the Fc/Fc γ R complex recruits these effector cells to sites of bound antigen, typically resulting in signaling events within the cells and important subsequent immune responses such as release of inflammation mediators, B cell activation, endocytosis, phagocytosis, and cytotoxic attack. The ability to mediate cytotoxic and phagocytic effector functions is a potential mechanism by which antibodies destroy targeted cells. The cell-mediated reaction wherein nonspecific cytotoxic cells that express Fc γ Rs recognize bound antibody on a target cell and subsequently cause lysis of the target cell is referred to as antibody dependent cell-mediated cytotoxicity (ADCC). The cell-mediated reaction wherein nonspecific cytotoxic cells that express Fc γ Rs recognize bound antibody on a target cell and subsequently cause phagocytosis of the target cell is referred to as antibody dependent cell-mediated phagocytosis (ADCP).

[0993] The different IgG subclasses have different affinities for the Fc γ Rs, with IgG1 and IgG3 typically binding substantially better to the receptors than IgG2 and IgG4. All Fc γ Rs bind the same region on IgG Fc, yet with different affinities: the high affinity binder Fc γ RI has a K_d for IgG1 of 10⁻⁸ M-1, whereas the low affinity receptors Fc γ RII and Fc γ RIII generally bind at 10⁻⁶ and 10⁻⁵ respectively.

[0994] By "Fc gamma receptor" or "Fc γ R" as used herein is meant any member of the family of proteins that bind the IgG antibody Fc region and are substantially encoded by the Fc γ R genes. In humans this family includes but is not limited to Fc γ RI (CD64), including isoforms Fc γ RIa, Fc γ RIb, and Fc γ RIc; Fc γ RII (CD32), including isoforms Fc γ RIIa (including allotypes H131 and R131), Fc γ RIIb (including Fc γ RIIb-1 and Fc γ RIIb-2), and Fc γ RIIc; and Fc γ RIII (CD16), including isoforms Fc γ RIIIa (including allotypes V158 and F158) and Fc γ RIIIb (including allotypes Fc γ RIIIb-NA1 and Fc γ RIIIb-NA2) (Jefferis et al., 2002, *Immunol Lett* 82:57-65), as well as any undiscovered human Fc γ Rs or Fc γ R isoforms or allotypes. An Fc γ R may be from any organism, including but not limited to humans, mice, rats, rabbits, and monkeys. Mouse Fc γ Rs include but are not limited to Fc γ RI (CD64), Fc γ RII (CD32), Fc γ RIII (CD16), and Fc γ RIII-2 (CD16- 2), as well as any undiscovered mouse Fc γ Rs or Fc γ R isoforms or allotypes.

[0995] Binding to FcγRs can be performed directly by determining the affinity of an antibody for the FcγR. Or binding can be determined indirectly by measuring cytokine release in a cell-based assay.

[0996] To facilitate the association of a particular heavy chain with its cognate light chain, both the heavy and light chains may contain complimentary amino acid substitutions. As used herein, “complimentary amino acid substitutions” refer to a substitution to a positively-charged amino acid in one chain paired with a negatively-charged amino acid substitution in the other chain. For example, in some embodiments, the heavy chain comprises at least one amino acid substitution to introduce a charged amino acid and the corresponding light chain comprises at least one amino acid substitution to introduce a charged amino acid, wherein the charged amino acid introduced into the heavy chain has the opposite charge of the amino acid introduced into the light chain. In certain embodiments, one or more positively-charged residues (e.g., lysine, histidine or arginine) can be introduced into a first light chain (LC1) and one or more negatively-charged residues (e.g., aspartic acid or glutamic acid) can be introduced into the companion heavy chain (HC1) at the binding interface of LC1/HC1, whereas one or more negatively-charged residues (e.g., aspartic acid or glutamic acid) can be introduced into a second light chain (LC2) and one or more positively-charged residues (e.g., lysine, histidine or arginine) can be introduced into the companion heavy chain (HC2) at the binding interface of LC2/HC2. The electrostatic interactions will direct the LC1 to pair with HC1 and LC2 to pair with HC2, as the opposite charged residues (polarity) at the interface attract. The heavy/light chain pairs having the same charged residues (polarity) at an interface (e.g. LC1/HC2 and LC2/HC1) will repel, resulting in suppression of the unwanted HC/LC pairings.

[0997] In these and other embodiments, the CH1 domain of the heavy chain or the CL domain of the light chain comprises an amino acid sequence differing from wild-type IgG amino acid sequence such that one or more positively-charged amino acids in wild-type IgG amino acid sequence is replaced with one or more negatively-charged amino acids. Alternatively, the CH1 domain of the heavy chain or the CL domain of the light chain comprises an amino acid sequence differing from wild-type IgG amino acid sequence such that one or more negatively-charged amino acids in wild-type IgG amino acid sequence is replaced with one or more positively-charged amino acids. In some embodiments, one or more amino acids in the CH1 domain of the first and/or second heavy chain in the multispecific antibody construct at an EU position selected from F126, P127, L128, A141, L145, K147, D148, H168, F170, P171, V173, Q175, S176, S183, V185 and K213 is replaced with a charged amino acid. In certain embodiments, a heavy chain residue for substitution with a negatively- or positively- charged amino acid is S183 (EU numbering system). In some embodiments, S183 is substituted with a positively-charged amino acid. In alternative embodiments, S183 is substituted with a negatively-charged amino acid. For instance, in one embodiment, S183 is substituted with a negatively-charged amino acid (e.g. S183E) in the first heavy chain, and S183 is substituted with a positively-charged amino acid (e.g. S183K) in the second heavy chain.

[0998] In embodiments in which the light chain is a kappa light chain, one or more amino acids in the CL domain of the first and/or second light chain in the multimeric antibody construct at a position (EU numbering in a kappa light chain) selected from F116, F118, S121, D122, E123, Q124, S131, V133, L135, N137, N138, Q160, S162, T164, S174 and S176 is replaced with a charged amino acid. In embodiments in which the light chain is a lambda light chain, one or more amino acids in the CL domain of the first and/or second light chain in the multispecific antibody construct at a position (EU numbering in a lambda chain) selected from T116, F118, S121, E123, E124, K129, T131, V133, L135, S137, E160, T162, S165, Q167, A174, S176 and Y178 is replaced with a charged amino acid. In some embodiments, a residue for substitution with a negatively- or positively- charged amino acid is S176 (EU numbering system) of the CL domain of either a kappa or lambda light chain. In certain embodiments, S176 of the CL domain is replaced with a positively-charged amino acid. In alternative embodiments, S176 of the CL domain is replaced with a negatively-charged amino acid. In one embodiment, S176 is substituted with a positively-charged amino acid (e.g. S176K) in the first light chain, and S176 is substituted with a negatively-charged amino acid (e.g. S176E) in the second light chain.

[0999] In addition to or as an alternative to the complimentary amino acid substitutions in the CH1 and CL domains, the variable regions of the light and heavy chains in the multispecific antibody construct may contain one or more complimentary amino acid substitutions to introduce charged amino acids. For instance, in some embodiments, the VH region of the heavy chain or the VL region of the light chain of a multispecific antibody construct comprises an amino acid sequence differing from wild-type IgG amino acid sequence such that one or more positively-charged amino acids in wild-type IgG amino acid sequence is replaced with one or more negatively-charged amino acids. Alternatively, the VH region of the heavy chain or the VL region of the light chain comprises an amino acid sequence differing from wild-type IgG amino acid sequence such that one or more negatively-charged amino acids in wild-type IgG amino acid sequence is replaced with one or more positively-charged amino acids.

[1000] V region interface residues (i.e., amino acid residues that mediate assembly of the VH and VL regions) within the VH region include EU positions 1, 3, 35, 37, 39, 43, 44, 45, 46, 47, 50, 59, 89, 91, and 93. One or more of these interface residues in the VH region can be substituted with a charged (positively- or negatively-charged) amino acid. In certain embodiments, the amino acid at EU position 39 in the VH region of the first and/or second heavy chain is substituted for a positively-charged amino acid, e.g., lysine. In alternative embodiments, the amino acid at EU position 39 in the VH region of the first and/or second heavy chain is substituted for a negatively-charged amino acid, e.g., glutamic acid. In some embodiments, the amino acid at EU position 39 in the VH region of the first heavy chain is substituted for a negatively-charged amino acid (e.g. G39E), and the amino acid at EU position 39 in the VH region of the second heavy chain is substituted for a positively-charged amino

acid (e.g. G39K). In some embodiments, the amino acid at EU position 44 in the VH region of the first and/or second heavy chain is substituted for a positively-charged amino acid, e.g., lysine. In alternative embodiments, the amino acid at EU position 44 in the VH region of the first and/or second heavy chain is substituted for a negatively-charged amino acid, e.g., glutamic acid. In certain embodiments, the amino acid at EU position 44 in the VH region of the first heavy chain is substituted for a negatively-charged amino acid (e.g. G44E), and the amino acid at EU position 44 in the VH region of the second heavy chain is substituted for a positively-charged amino acid (e.g. G44K).

[1001] V region interface residues (i.e., amino acid residues that mediate assembly of the VH and VL regions) within the VL region include EU positions 32, 34, 35, 36, 38, 41, 42, 43, 44, 45, 46, 48, 49, 50, 51, 53, 54, 55, 56, 57, 58, 85, 87, 89, 90, 91, and 100. One or more interface residues in the VL region can be substituted with a charged amino acid, preferably an amino acid that has an opposite charge to those introduced into the VH region of the cognate heavy chain. In some embodiments, the amino acid at EU position 100 in the VL region of the first and/or second light chain is substituted for a positively-charged amino acid, e.g., lysine. In alternative embodiments, the amino acid at EU position 100 in the VL region of the first and/or second light chain is substituted for a negatively-charged amino acid, e.g., glutamic acid. In certain embodiments, the amino acid at EU position 100 in the VL region of the first light chain is substituted for a positively-charged amino acid (e.g. G100K), and the amino acid at EU position 100 in the VL region of the second light chain is substituted for a negatively-charged amino acid (e.g. G100E).

[1002] Any of the constant domains can be modified to contain one or more of the charge pair mutations described above to facilitate correct assembly of a multispecific antibody construct.

[1003] As used herein, the term “Fc region” refers to the C-terminal region of an immunoglobulin heavy chain which may be generated by papain digestion of an intact antibody. The Fc region of an immunoglobulin generally comprises two constant domains, a CH2 domain and a CH3 domain, and optionally comprises a CH4 domain. In certain embodiments, the Fc region is an Fc region from an IgG1, IgG2, IgG3, or IgG4 immunoglobulin. In some embodiments, the Fc region comprises CH2 and CH3 domains from a human IgG1 or human IgG2 immunoglobulin. The Fc region may retain effector function, such as C1q binding, complement dependent cytotoxicity (CDC), Fc receptor binding, antibody-dependent cell-mediated cytotoxicity (ADCC), and phagocytosis. In other embodiments, the Fc region may be modified to reduce or eliminate effector function as described in further detail herein.

[1004] The heavy chain constant regions or the Fc regions of the multispecific antibody constructs described herein may comprise one or more amino acid substitutions that affect the glycosylation and/or effector function of the antigen binding protein. One of the functions of the Fc region of an immunoglobulin is to communicate to the immune system when the immunoglobulin binds its target. This is commonly referred to as “effector function.” Communication leads to antibody-dependent

cellular cytotoxicity (ADCC), antibody-dependent cellular phagocytosis (ADCP), and/or complement dependent cytotoxicity (CDC). ADCC and ADCP are mediated through the binding of the Fc region to Fc receptors on the surface of cells of the immune system. CDC is mediated through the binding of the Fc with proteins of the complement system, e.g., C1q. In some embodiments, the multispecific antibody constructs of the invention comprise one or more amino acid substitutions in the constant region to enhance effector function, including ADCC activity, CDC activity, ADCP activity, and/or the clearance or half-life of the antigen binding protein. Exemplary amino acid substitutions (EU numbering) that can enhance effector function include, but are not limited to, E233L, L234I, L234Y, L235S, G236A, S239D, F243L, F243V, P247I, D280H, K290S, K290E, K290N, K290Y, R292P, E294L, Y296W, S298A, S298D, S298V, S298G, S298T, T299A, Y300L, V305I, Q311M, K326A, K326E, K326W, A330S, A330L, A330M, A330F, I332E, D333A, E333S, E333A, K334A, K334V, A339D, A339Q, P396L, or combinations of any of the foregoing.

[1005] In other embodiments, the multispecific antibody constructs of the invention comprise one or more amino acid substitutions in the constant region to reduce effector function. Exemplary amino acid substitutions (EU numbering) that can reduce effector function include, but are not limited to, C220S, C226S, C229S, E233P, L234A, L234V, V234A, L234F, L235A, L235E, G237A, P238S, S267E, H268Q, N297A, N297G, V309L, E318A, L328F, A330S, A331S, P331S or combinations of any of the foregoing.

[1006] Glycosylation can contribute to the effector function of antibodies, particularly IgG1 antibodies. Thus, in some embodiments, the multispecific antibody constructs of the invention may comprise one or more amino acid substitutions that affect the level or type of glycosylation of the binding proteins. Glycosylation of polypeptides is typically either N-linked or O-linked. N-linked refers to the attachment of the carbohydrate moiety to the side chain of an asparagine residue. The tri-peptide sequences asparagine-X-serine and asparagine-X-threonine, where X is any amino acid except proline, are the recognition sequences for enzymatic attachment of the carbohydrate moiety to the asparagine side chain. Thus, the presence of either of these tri-peptide sequences in a polypeptide creates a potential glycosylation site. O-linked glycosylation refers to the attachment of one of the sugars N-acetylgalactosamine, galactose, or xylose, to a hydroxyamino acid, most commonly serine or threonine, although 5-hydroxyproline or 5-hydroxylysine may also be used.

[1007] In certain embodiments, glycosylation of the multispecific antibody constructs described herein is increased by adding one or more glycosylation sites, e.g., to the Fc region of the binding protein. Addition of glycosylation sites to the antigen binding protein can be conveniently accomplished by altering the amino acid sequence such that it contains one or more of the above-described tri-peptide sequences (for N-linked glycosylation sites). The alteration may also be made by the addition of, or substitution by, one or more serine or threonine residues to the starting sequence (for O-linked glycosylation sites). For ease, the antigen binding protein amino acid sequence may be

altered through changes at the DNA level, particularly by mutating the DNA encoding the target polypeptide at preselected bases such that codons are generated that will translate into the desired amino acids.

[1008] The invention also encompasses production of multispecific antibody construct molecules with altered carbohydrate structure resulting in altered effector activity, including antigen binding proteins with absent or reduced fucosylation that exhibit improved ADCC activity. Various methods are known in the art to reduce or eliminate fucosylation. For example, ADCC effector activity is mediated by binding of the antibody molecule to the FcγRIII receptor, which has been shown to be dependent on the carbohydrate structure of the N-linked glycosylation at the N297 residue of the CH2 domain. Non-fucosylated antibodies bind this receptor with increased affinity and trigger FcγRIII-mediated effector functions more efficiently than native, fucosylated antibodies. For example, recombinant production of non-fucosylated antibody in CHO cells in which the alpha-1,6-fucosyl transferase enzyme has been knocked out results in antibody with 100-fold increased ADCC activity (*see Yamane-Ohnuki et al.*, *Biotechnol Bioeng.* 87(5):614-22, 2004). Similar effects can be accomplished through decreasing the activity of alpha-1,6-fucosyl transferase enzyme or other enzymes in the fucosylation pathway, e.g., through siRNA or antisense RNA treatment, engineering cell lines to knockout the enzyme(s), or culturing with selective glycosylation inhibitors (*see Rothman et al.*, *Mol Immunol.* 26(12):1113-23, 1989). Some host cell strains, e.g. Lec13 or rat hybridoma YB2/0 cell line naturally produce antibodies with lower fucosylation levels (*see Shields et al.*, *J Biol Chem.* 277(30):26733-40, 2002 and Shinkawa *et al.*, *J Biol Chem.* 278(5):3466-73, 2003). An increase in the level of bisected carbohydrate, e.g. through recombinantly producing antibody in cells that overexpress GnTIII enzyme, has also been determined to increase ADCC activity (*see Umana et al.*, *Nat Biotechnol.* 17(2):176-80, 1999).

[1009] In other embodiments, glycosylation of the multispecific antibody constructs described herein is decreased or eliminated by removing one or more glycosylation sites, e.g., from the Fc region of the binding protein. Amino acid substitutions that eliminate or alter N-linked glycosylation sites can reduce or eliminate N-linked glycosylation of the antigen binding protein. In certain embodiments, the multispecific antibody constructs described herein comprise a mutation at position N297 (EU numbering), such as N297Q, N297A, or N297G. In certain embodiments, the multispecific antibody constructs described herein comprise a mutation at positions L234 and L235 (EU numbering), such as L234A and L235A. In one particular embodiment, the multispecific antibody constructs of the invention comprise a Fc region from a human IgG1 antibody with a N297G mutation. To improve the stability of molecules comprising a N297 mutation, the Fc region of the molecules may be further engineered. For instance, in some embodiments, one or more amino acids in the Fc region are substituted with cysteine to promote disulfide bond formation in the dimeric state. Residues corresponding to V259, A287, R292, V302, L306, V323, or I332 (EU numbering) of an IgG1 Fc region may thus be substituted with cysteine. In one embodiment, specific pairs of residues are

substituted with cysteine such that they preferentially form a disulfide bond with each other, thus limiting or preventing disulfide bond scrambling. In certain embodiments pairs include, but are not limited to, A287C and L306C, V259C and L306C, R292C and V302C, and V323C and I332C. In particular embodiments, the multispecific antibody constructs described herein comprise a Fc region from a human IgG1 antibody with mutations at R292C and V302C. In such embodiments, the Fc region may also comprise a N297G mutation.

[1010] In one embodiment, the heavy chain comprises an amino acid substitution selected from the group consisting of:

[1011] (i) N297G or N297A;

[1012] (ii) L234A and L235A; and

[1013] (iii) R292C and V302C;

[1014] wherein the amino acid numbering is EU numbering according to Kabat.

[1015] In one embodiment, the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

[1016] Modifications of the multispecific antibody constructs of the invention to increase serum half-life also may be desirable, for example, by incorporation of or addition of a salvage receptor binding epitope (e.g., by mutation of the appropriate region or by incorporating the epitope into a peptide tag that is then fused to the antigen binding protein at either end or in the middle, e.g., by DNA or peptide synthesis; *see, e.g.*, WO96/32478) or adding molecules such as PEG or other water soluble polymers, including polysaccharide polymers. The salvage receptor binding epitope preferably constitutes a region wherein any one or more amino acid residues from one or two loops of a Fc region are transferred to an analogous position in the antigen binding protein. In one embodiment, three or more residues from one or two loops of the Fc region are transferred. In one embodiment, the epitope is taken from the CH2 domain of the Fc region (e.g., an IgG Fc region) and transferred to the CH1, CH3, or VH region, or more than one such region, of the antigen binding protein. Alternatively, the epitope is taken from the CH2 domain of the Fc region and transferred to the CL region or VL region, or both, of the antigen binding protein. *See* International applications WO 97/34631 and WO 96/32478 for a description of Fc variants and their interaction with the salvage receptor.

[1017] In certain embodiments of the multispecific antibody construct of the invention, the binding domain positioned at the amino terminus of the Fc region (i.e. the amino-terminal binding domain) is a Fab fragment fused to the amino terminus of the Fc region through a peptide linker described herein or through an immunoglobulin hinge region. An “immunoglobulin hinge region” refers to the amino acid sequence connecting the CH1 domain and the CH2 domain of an immunoglobulin heavy chain. The hinge region of human IgG1 is generally defined as the amino acid sequence from about Glu216 or about Cys226, to about Pro230. Hinge regions of other IgG isotypes may be aligned with the IgG1 sequence by placing the first and last cysteine residues forming inter-heavy chain disulfide bonds in

the same positions and are determinable to those of skill in the art. In some embodiments, the amino-terminal binding domain is joined to the amino terminus of the Fc region through a human IgG1 hinge region. In other embodiments, the amino-terminal binding domain is joined to the amino terminus of the Fc region through a human IgG2 hinge region. In one embodiment, the amino-terminal binding domain (e.g. Fab fragment) is fused to the Fc region through the carboxyl terminus of the CH1 region of the Fab.

[1018] In some embodiments of the heavy chain fusion proteins of the invention, the binding domain positioned at the carboxyl terminus of the Fc region (i.e. the carboxyl-terminal binding domain) is a Fab fragment. In such embodiments, the Fab is fused or otherwise connected to the carboxyl terminus of the Fc region (e.g. the carboxyl terminus of the CH3 domain) through a peptide linker through the amino terminus of the VH region of the Fab fragment. Thus, in one embodiment, the Fab is fused to an Fc region through the amino terminus of the VH region of the Fab such that the resulting fusion protein comprises, from N-terminus to C-terminus, a CH2 domain, a CH3 domain, a peptide linker, a VH region, and a CH1 region.

[1019] The peptide linker joining the Fc region to the carboxyl-terminal Fab can be any of the peptide linkers described herein. In particular embodiments, the peptide linker joining the Fc region to the carboxyl-terminal Fab fragment is at least 5 amino acids in length. In other embodiments, the peptide linker joining the Fc region to the carboxyl-terminal Fab fragment is at least 8 amino acids in length. Particularly suitable peptide linkers for joining the Fc region to the carboxyl-terminal Fab fragment are glycine-serine linkers, such as $(\text{Gly}_x\text{Ser})_n$ where $x=3$ or 4 and $n=2, 3, 4, 5$ or 6 (SEQ ID NO: 923). In one embodiment, the peptide linker connecting the Fc region to the carboxyl-terminal Fab fragment is a L10 $(\text{G}_4\text{S})_2$ linker (SEQ ID NO: 888). In another embodiment, the peptide linker connecting the Fc region to the carboxyl-terminal Fab fragment is a L9 or $\text{G}_3\text{SG}_4\text{S}$ linker (SEQ ID NO: 924).

[1020] In some embodiments of the antigen binding proteins of the invention in which the carboxyl-terminal binding domain is a Fab fragment, the binding domain positioned at the amino terminus of the Fc region (i.e. the amino-terminal binding domain) is also a Fab fragment. The amino-terminal Fab fragment can be fused to the amino terminus of the Fc region through a peptide linker or an immunoglobulin hinge region described herein. In some embodiments, the amino-terminal Fab fragment is joined to the amino terminus of the Fc region through a human IgG1 hinge region. In other embodiments, the amino-terminal Fab fragment is joined to the amino terminus of the Fc region through a human IgG2 hinge region. In one embodiment, the amino-terminal Fab fragment is fused to the Fc region through the carboxyl terminus of the CH1 region of the Fab.

[1021] In some embodiments, the multispecific antibody construct of the invention comprises a first antibody that specifically binds to a first target where one polypeptide chain (e.g. the heavy chain (VH2-CH1)) of a Fab fragment from a second antibody that specifically binds to a second target is

fused to the carboxyl terminus of the heavy chain of the first antibody. The multispecific antibody construct in such embodiments also comprises a polypeptide chain containing the other half of the Fab fragment from the second antibody (e.g., the light chain (VL2-CL)). This format is referred to herein as the “IgG-Fab” format, and one embodiment of this type of molecule is shown schematically in Figure 1. Thus, in certain embodiments, the present invention includes a bispecific, multivalent antigen binding protein comprising: (i) a light chain from a first antibody, (ii) a heavy chain from the first antibody, wherein the heavy chain is fused at its carboxyl terminus through a peptide linker to a first polypeptide comprising VH-CH1 domains of a second antibody to form a modified heavy chain, and (iii) a second polypeptide comprising VL-CL domains of the second antibody. When dimerized, the multispecific antibody construct is a homohexamer comprising two modified heavy chains, two light chains from the first antibody, and two polypeptide chains containing the other half of the Fab fragment from the second antibody (the Fd fragment). In one embodiment, the first polypeptide, which is fused to the carboxyl terminus of the heavy chain, comprises VH and CH1 domains from the second antibody, and the second polypeptide comprises VL and CL domains from the second antibody.

[1022] Charge pair mutations or complimentary amino acid substitutions as described herein can be introduced into the Fab regions of the first antibody (Fab 1) or second antibody (Fab 2) to promote correct heavy chain-light chain pairing. For instance, in some embodiments, the amino acid at EU position 38 of the VL domain in Fab 1 is replaced with a negatively-charged amino acid (e.g. glutamic acid) and the amino acid at EU position 39 of the VH domain in Fab 1 is replaced with a positively-charged amino acid (e.g. lysine). In other embodiments, the amino acid at EU position 38 of the VL domain in Fab 1 is replaced with a positively-charged amino acid (e.g. lysine) and the amino acid at EU position 39 of the VH domain in Fab 1 is replaced with a negatively-charged amino acid (e.g. glutamic acid). In certain embodiments, the amino acid at EU position 38 of the VL domain in Fab 2 is replaced with a negatively-charged amino acid (e.g. glutamic acid) and the amino acid at EU position 39 of the VH domain in Fab 2 is replaced with a positively-charged amino acid (e.g. lysine). In other embodiments, the amino acid at EU position 38 of the VL domain in Fab 2 is replaced with a positively-charged amino acid (e.g. lysine) and the amino acid at EU position 39 of the VH domain in Fab 2 is replaced with a negatively-charged amino acid (e.g. glutamic acid).

[1023] In embodiments in which the VH-CH1 region (i.e. Fd fragment) from the second antibody is fused to the heavy chain of the first antibody, the heavy chain from the first antibody comprises a S183E mutation (EU numbering), the light chain from the first antibody comprises a S176K mutation (EU numbering), the light chain from the second antibody comprises a S176E mutation (EU numbering), and the Fd region from the second antibody (which is fused to the C-terminus of the heavy chain from the first antibody) comprises a S183K mutation (EU numbering). In other embodiments, the heavy chain from the first antibody comprises a G44E mutation (EU) and S183E mutation (EU numbering), the light chain from the first antibody comprises a G100K mutation (EU)

and S176K mutation (EU numbering), the light chain from the second antibody comprises a G100E mutation (EU) and S176E mutation (EU numbering), and the Fd region from the second antibody (which is fused to the C-terminus of the heavy chain from the first antibody) comprises a G44K mutation (EU) and S183K mutation (EU numbering). The charges in the foregoing examples may be reversed so long as the charge on the corresponding light or heavy chain is also reversed so that the correct heavy/light chain pairs have opposite charges.

[1024] “Corresponds to” as it pertains to the VH2 and second CH1 domain means that the amino acid residues of the VH2 and second CH1 domain are counted from the C-terminus of the first heavy chain if there is no linker. If there is a peptide linker, the amino acid residues of the VH2 and second CH1 domain are counted from the C-terminus of the peptide linker. In neither case are the amino acid residues counted from the N-terminus of the first heavy chain. Rather, for the VH2 and second CH1 domain, counting begins at the first amino acid residue of the VH2 domain. The counting of amino acid residues is performed using the EU or AHo convention.

[1025] In certain embodiments: a) the VH1 comprises a Q39E mutation and the first CH1 domain comprises a S183K mutation using EU numbering; b) the VH2 comprises a Q39K mutation and the second CH1 domain comprises a S183E mutation using EU numbering; c) the VL1 comprises a Q38K mutation and the first CL domain comprises a S176E mutation using EU numbering; and d) the VL2 comprises a Q38E mutation and the second CL domain comprises a S176K mutation using EU numbering.

[1026] In certain embodiments: a) the first CH1 domain comprises G44E and S183K mutations using EU numbering; b) the second CH1 domain comprises G44K and S183E mutations using EU numbering; c) the first CL domain comprises G100K and S176E mutations using EU numbering; and d) the second CL domain comprises G100E and S176K mutations using EU numbering.

[1027] In certain embodiments: a) the VH1 comprises a Q39K mutation and the first CH1 domain comprises a S183E mutation using EU numbering; b) the VH2 comprises a Q39E mutation and the second CH1 domain comprises a S183K mutation using EU numbering; c) the VL1 comprises a Q38E mutation and the first CL domain comprises a S176K mutation using EU numbering; and d) the VL2 comprises a Q38K mutation and the second CL domain comprises a S176E mutation using EU numbering.

[1028] In certain embodiments: a) the first CH1 domain comprises G44K and S183E mutations using EU numbering; b) the second CH1 domain comprises G44E and S183K mutations using EU numbering; c) the first CL domain comprises G100E and S176K mutations using EU numbering; and d) the second CL domain comprises G100K and S176E mutations using EU numbering.

[1029] In certain embodiments the first heavy chain is fused to the VH2 via a peptide linker. In certain embodiments the peptide linker comprises a sequence selected from the group consisting of (Gly₃Ser)₂ (SEQ ID NO: 916), (Gly₄Ser)₂ (SEQ ID NO: 888), (Gly₃Ser)₃ (SEQ ID NO: 917),

(Gly₄Ser)₃ (SEQ ID NO: 889), (Gly₃Ser)₄ (SEQ ID NO: 918), (Gly₄Ser)₄ (SEQ ID NO: 890), (Gly₃Ser)₅ (SEQ ID NO: 919), (Gly₄Ser)₅ (SEQ ID NO: 920), (Gly₃Ser)₆ (SEQ ID NO: 921), and (Gly₄Ser)₆ (SEQ ID NO: 922). These sequences can also be written as GGGSGGGGS (SEQ ID NO: 916), GGGGSGGGGS (SEQ ID NO: 888), GGGSGGGSGGGGS (SEQ ID NO: 917), GGGGSGGGGSGGGGS (SEQ ID NO: 889), GGGSGGGSGGGSGGGGS (SEQ ID NO: 918), GGGGSGGGGSGGGGSGGGGS (SEQ ID NO: 890), GGGSGGGSGGGSGGGGSGGGGS (SEQ ID NO: 919), GGGGSGGGGSGGGGSGGGGSGGGGS (SEQ ID NO: 920), GGGSGGGSGGGSGGGGSGGGGS (SEQ ID NO: 921), and GGGGSGGGGSGGGGSGGGGSGGGGSGGGGS (SEQ ID NO: 922).

[1030] Additionally or alternatively, correct heavy-light chain pairing may be facilitated by swapping the CH1 and CL domains in the carboxyl-terminal Fab binding domain. By way of example, the first polypeptide, which is fused to the carboxyl terminus of the heavy chain, may comprise a VL domain and CH1 domain from the second antibody, and the second polypeptide may comprise a VH domain and CL domain from the second antibody. In another embodiment, the first polypeptide, which is fused to the carboxyl terminus of the heavy chain, may comprise a VH domain and a CL domain from the second antibody, and the second polypeptide may comprise a VL domain and CH1 domain from the second antibody.

[1031] In another aspect, the present invention is directed to a multispecific antibody construct comprising:

[1032] a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

[1033] i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

[1034] ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

[1035] b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

[1036] c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a positively charged

amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

[1037] wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

[1038] wherein:

[1039] the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

[1040] the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

[1041] In one aspect, the present invention is directed to a multispecific antibody construct comprising:

[1042] a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

[1043] i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

[1044] ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

[1045] b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

[1046] c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

[1047] wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

[1048] wherein:

[1049] the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

[1050] the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

[1051] In one embodiment, the hinge-CH2-CH3 polypeptide is linked to the VH2 via a peptide linker.

[1052] In one embodiment, the peptide linker comprises a sequence selected from the group consisting of (Gly₃Ser)₂ (SEQ ID NO: 916), (Gly₄Ser)₂ (SEQ ID NO: 888), (Gly₃Ser)₃ (SEQ ID NO: 917), (Gly₄Ser)₃ (SEQ ID NO: 889), (Gly₃Ser)₄ (SEQ ID NO: 918), (Gly₄Ser)₄ (SEQ ID NO: 890), (Gly₃Ser)₅ (SEQ ID NO: 919), (Gly₄Ser)₅ (SEQ ID NO: 920), (Gly₃Ser)₆ (SEQ ID NO: 921), and (Gly₄Ser)₆ (SEQ ID NO: 922).

[1053] In one embodiment,

[1054] a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;

[1055] b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;

[1056] c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering; and

[1057] d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering.

[1058] In one embodiment,

[1059] a) the first CH1 domain comprises a S183K mutation using EU numbering;

[1060] b) the second CH1 domain comprises a S183E mutation using EU numbering;

[1061] c) the first CL domain comprises a S176E mutation using EU numbering; and

[1062] d) the second CL domain comprises a S176K mutation using EU numbering.

[1063] In one embodiment,

[1064] a) the VH1 comprises a Q39K mutation and the first CH1 domain comprises a S183K mutation using EU numbering;

[1065] b) the VH2 comprises a Q39E mutation and the second CH1 domain comprises a S183E mutation using EU numbering;

[1066] c) the VL1 comprises a Q38E mutation and the first CL domain comprises a S176E mutation using EU numbering; and

[1067] d) the VL2 comprises a Q38K mutation and the second CL domain comprises a S176K mutation using EU numbering.

[1068] In one embodiment,

[1069] a) the first CH1 domain comprises G44K and S183K mutations using EU numbering;

[1070] b) the second CH1 domain comprises G44E and S183E mutations using EU numbering;

[1071] c) the first CL domain comprises G100E and S176E mutations using EU numbering; and

- [1072] d) the second CL domain comprises G100K and S176K mutations using EU numbering.
- [1073] In one embodiment,
- [1074] a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;
- [1075] b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;
- [1076] c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering; and
- [1077] d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering.
- [1078] In one embodiment,
- [1079] a) the first CH1 domain comprises a S183E mutation using EU numbering;
- [1080] b) the second CH1 domain comprises a S183K mutation using EU numbering;
- [1081] c) the first CL domain comprises a S176K mutation using EU numbering; and
- [1082] d) the second CL domain comprises a S176E mutation using EU numbering.
- [1083] In one embodiment,
- [1084] a) the VH1 comprises a Q39E mutation and the first CH1 domain comprises a S183E mutation using EU numbering;
- [1085] b) the VH2 comprises a Q39K mutation and the second CH1 domain comprises a S183K mutation using EU numbering;
- [1086] c) the VL1 comprises a Q38K mutation and the first CL domain comprises a S176K mutation using EU numbering; and
- [1087] d) the VL2 comprises a Q38E mutation and the second CL domain comprises a S176E mutation using EU numbering.
- [1088] In one embodiment,
- [1089] a) the first CH1 domain comprises G44E and S183E mutations using EU numbering;
- [1090] b) the second CH1 domain comprises G44K and S183K mutations using EU numbering;
- [1091] c) the first CL domain comprises G100K and S176K mutations using EU numbering; and
- [1092] d) the second CL domain comprises G100E and S176E mutations using EU numbering.
- [1093] In one embodiment, the hinge-CH2-CH3 polypeptide comprises an amino acid substitution selected from the group consisting of:
- [1094] (i) N297G or N297A;
- [1095] (ii) L234A and L235A; and
- [1096] (iii) R292C and V302C;
- [1097] wherein the amino acid numbering is EU numbering according to Kabat.

[1098] In one embodiment, the hinge-CH2-CH3 polypeptide comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

[1099] In one embodiment, the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human MSLN (SEQ ID NO: 2); and

[1100] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[1101] SEQ ID NOs: 58, 59, and 60, respectively;

[1102] SEQ ID NOs: 64, 65, and 66, respectively;

[1103] SEQ ID NOs: 70, 71, and 72, respectively;

[1104] SEQ ID NOs: 76, 77, and 78, respectively;

[1105] SEQ ID NOs: 82, 83, and 84, respectively;

[1106] SEQ ID NOs: 88, 89, and 90, respectively;

[1107] SEQ ID NOs: 94, 95, and 96, respectively;

[1108] SEQ ID NOs: 100, 101, and 102, respectively;

[1109] SEQ ID NOs: 106, 107, and 108, respectively;

[1110] SEQ ID NOs: 112, 113, and 114, respectively;

[1111] SEQ ID NOs: 118, 119, and 120, respectively;

[1112] SEQ ID NOs: 124, 125, and 126, respectively; and

[1113] SEQ ID NOs: 130, 131, and 132, respectively;

[1114] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[1115] SEQ ID NOs: 136, 137, and 138, respectively;

[1116] SEQ ID NOs: 142, 143, and 144, respectively;

[1117] SEQ ID NOs: 148, 149, and 150, respectively;

[1118] SEQ ID NOs: 154, 155, and 156, respectively;

[1119] SEQ ID NOs: 160, 161, and 162, respectively;

[1120] SEQ ID NOs: 166, 167, and 168, respectively;

[1121] SEQ ID NOs: 172, 173, and 174, respectively;

[1122] SEQ ID NOs: 178, 179, and 180, respectively;

[1123] SEQ ID NOs: 184, 185, and 186, respectively;

[1124] SEQ ID NOs: 190, 191, and 192, respectively;

[1125] SEQ ID NOs: 196, 197, and 198, respectively;

[1126] SEQ ID NOs: 202, 203, and 204, respectively; and

[1127] SEQ ID NOs: 208, 209, and 210, respectively;

[1128] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[1129] SEQ ID NOs: 230, 231, and 232, respectively;

- [1130] SEQ ID NOs: 236, 237, and 238, respectively;
- [1131] SEQ ID NOs: 242, 243, and 244, respectively; and
- [1132] SEQ ID NOs: 248, 249, and 250, respectively;
- [1133] and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [1134] SEQ ID NOs: 254, 255, and 256, respectively;
- [1135] SEQ ID NOs: 260, 261, and 262, respectively;
- [1136] SEQ ID NOs: 266, 267, and 268, respectively; and
- [1137] SEQ ID NOs: 272, 273, and 274, respectively.
- [1138]
- [1139] 64. The antibody construct of claim 63, wherein
- [1140] 1) the VL1 and VH1 are selected from the group consisting of:
- [1141] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;
- [1142] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;
- [1143] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;
- [1144] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;
- [1145] e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;
- [1146] f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;
- [1147] g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;
- [1148] h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

[1149] i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[1150] j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[1151] k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[1152] l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[1153] m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;

[1154] and

[1155] 2) the VL2 and VH2 are selected from the group consisting of:

[1156] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

[1157] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

[1158] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

[1159] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

[1160] In one embodiment:

[1161] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

[1162] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

[1163] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[1164] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

[1165] In one embodiment:

[1166] 1) the VL1 and VH1 are selected from the group consisting of:

[1167] a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;

[1168] b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;

[1169] c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;

[1170] d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;

[1171] e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;

[1172] f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;

[1173] g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;

[1174] h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;

[1175] i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;

[1176] j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;

[1177] k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;

[1178] l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and

[1179] m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;

[1180] and

[1181] 2) the VL2 and VH2 are selected from the group consisting of:

[1182] a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;

[1183] b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;

[1184] c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and

[1185] d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.

[1186] In one embodiment, the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1); wherein:

[1187] the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

[1188] SEQ ID NOs: 230, 231, and 232, respectively;

[1189] SEQ ID NOs: 236, 237, and 238, respectively;

[1190] SEQ ID NOs: 242, 243, and 244, respectively; and

[1191] SEQ ID NOs: 248, 249, and 250, respectively;

[1192] the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

[1193] SEQ ID NOs: 254, 255, and 256, respectively;

[1194] SEQ ID NOs: 260, 261, and 262, respectively;

[1195] SEQ ID NOs: 266, 267, and 268, respectively; and

- [1196] SEQ ID NOs: 272, 273, and 274, respectively;
- [1197] the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:
- [1198] SEQ ID NOs: 58, 59, and 60, respectively;
- [1199] SEQ ID NOs: 64, 65, and 66, respectively;
- [1200] SEQ ID NOs: 70, 71, and 72, respectively;
- [1201] SEQ ID NOs: 76, 77, and 78, respectively;
- [1202] SEQ ID NOs: 82, 83, and 84, respectively;
- [1203] SEQ ID NOs: 88, 89, and 90, respectively;
- [1204] SEQ ID NOs: 94, 95, and 96, respectively;
- [1205] SEQ ID NOs: 100, 101, and 102, respectively;
- [1206] SEQ ID NOs: 106, 107, and 108, respectively;
- [1207] SEQ ID NOs: 112, 113, and 114, respectively;
- [1208] SEQ ID NOs: 118, 119, and 120, respectively;
- [1209] SEQ ID NOs: 124, 125, and 126, respectively; and
- [1210] SEQ ID NOs: 130, 131, and 132, respectively;
- [1211] and
- [1212] the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:
- [1213] SEQ ID NOs: 136, 137, and 138, respectively;
- [1214] SEQ ID NOs: 142, 143, and 144, respectively;
- [1215] SEQ ID NOs: 148, 149, and 150, respectively;
- [1216] SEQ ID NOs: 154, 155, and 156, respectively;
- [1217] SEQ ID NOs: 160, 161, and 162, respectively;
- [1218] SEQ ID NOs: 166, 167, and 168, respectively;
- [1219] SEQ ID NOs: 172, 173, and 174, respectively;
- [1220] SEQ ID NOs: 178, 179, and 180, respectively;
- [1221] SEQ ID NOs: 184, 185, and 186, respectively;
- [1222] SEQ ID NOs: 190, 191, and 192, respectively;
- [1223] SEQ ID NOs: 196, 197, and 198, respectively;
- [1224] SEQ ID NOs: 202, 203, and 204, respectively; and
- [1225] SEQ ID NOs: 208, 209, and 210, respectively.
- [1226] In one embodiment,
- [1227] 1) the VL1 and VH1 are selected from the group consisting of:

[1228] a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

[1229] b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

[1230] c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

[1231] d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;

[1232] and

[1233] 2) the VL2 and VH2 are selected from the group consisting of:

[1234] a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

[1235] b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

[1236] c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

[1237] d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

[1238] e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

[1239] f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

[1240] g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

[1241] h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

[1242] i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

[1243] j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

[1244] k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

[1245] l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

[1246] m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

[1247] In one embodiment,

[1248] the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

[1249] the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

[1250] the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

[1251] the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

[1252] In one embodiment,

[1253] 1) the VL1 and VH1 are selected from the group consisting of:

[1254] a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;

[1255] b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;

[1256] c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and

[1257] d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;

[1258] and

[1259] 2) the VL2 and VH2 are selected from the group consisting of:

[1260] a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;

- [1261] b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;
[1262] c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;
[1263] d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;
[1264] e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;
[1265] f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;
[1266] g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;
[1267] h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;
[1268] i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;
[1269] j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;
[1270] k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;
[1271] l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and
[1272] m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.

[1273] The present invention includes one or more isolated nucleic acids encoding the multispecific antibody constructs and components thereof described herein. Nucleic acid molecules of the invention include DNA and RNA in both single-stranded and double-stranded form, as well as the corresponding complementary sequences. DNA includes, for example, cDNA, genomic DNA, chemically synthesized DNA, DNA amplified by PCR, and combinations thereof. The nucleic acid molecules of the invention include full-length genes or cDNA molecules as well as a combination of fragments thereof. In one embodiment, the nucleic acids of the invention are derived from human sources, but the invention includes those derived from non-human species, as well.

[1274] An "isolated nucleic acid," which is used interchangeably herein with "isolated polynucleotide," is a nucleic acid that has been separated from adjacent genetic sequences present in the genome of the organism from which the nucleic acid was isolated, in the case of nucleic acids isolated from naturally-occurring sources. In the case of nucleic acids synthesized enzymatically from a template or chemically, such as PCR products, cDNA molecules, or oligonucleotides for example, it is understood that the nucleic acids resulting from such processes are isolated nucleic acids. An isolated nucleic acid molecule refers to a nucleic acid molecule in the form of a separate fragment or as a component of a larger nucleic acid construct. In one embodiment, the nucleic acids are substantially free from contaminating endogenous material. The nucleic acid molecule has been derived from DNA or RNA isolated at least once in substantially pure form and in a quantity or concentration enabling identification, manipulation, and recovery of its component nucleotide sequences by standard biochemical methods (such as those outlined in Sambrook *et al.*, Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor Laboratory, Cold Spring Harbor, NY (1989)). Such sequences are provided and/or constructed in the form of an open reading frame uninterrupted by internal non-translated sequences, or introns, that are typically present in eukaryotic genes. Sequences of non-translated DNA can be present 5' or 3' from an open reading frame, where

the same do not interfere with manipulation or expression of the coding region. Unless specified otherwise, the left-hand end of any single-stranded polynucleotide sequence discussed herein is the 5' end; the left-hand direction of double-stranded polynucleotide sequences is referred to as the 5' direction. The direction of 5' to 3' production of nascent RNA transcripts is referred to as the transcription direction; sequence regions on the DNA strand having the same sequence as the RNA transcript that are 5' to the 5' end of the RNA transcript are referred to as "upstream sequences;" sequence regions on the DNA strand having the same sequence as the RNA transcript that are 3' to the 3' end of the RNA transcript are referred to as "downstream sequences."

[1275] In one aspect, the present invention is directed to a polynucleotide encoding the light chain of the antibody construct of the present invention.

[1276] In one aspect, the present invention is directed to a polynucleotide encoding the heavy chain fusion protein of the antibody construct of the present invention.

[1277] In one aspect, the present invention is directed to a vector comprising the polynucleotide encoding the light chain of the antibody construct, the polynucleotide encoding the heavy chain of the antibody construct, or both.

[1278] In one aspect, the present invention is directed to a host cell transformed or transfected with the vector or the polynucleotide encoding the light chain of the antibody construct and the polynucleotide encoding the heavy chain of the antibody construct.

[1279] In one aspect, the present invention is directed to a process for producing the antibody construct of the present invention, the process comprising culturing a host cell comprising a polynucleotide encoding the light chain and also comprising a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

[1280] Variants of the antigen binding proteins described herein can be prepared by site-specific mutagenesis of nucleotides in the DNA encoding the polypeptide, using cassette or PCR mutagenesis or other techniques well known in the art, to produce DNA encoding the variant, and thereafter expressing the recombinant DNA in cell culture as outlined herein. However, antigen binding proteins comprising variant CDRs having up to about 100-150 residues may be prepared by *in vitro* synthesis using established techniques. The variants typically exhibit the same qualitative biological activity as the naturally occurring analogue, e.g., binding to antigen. Such variants include, for example, deletions and/or insertions and/or substitutions of residues within the amino acid sequences of the antigen binding proteins. Any combination of deletion, insertion, and substitution is made to arrive at the final construct, provided that the final construct possesses the desired characteristics. The amino acid changes also may alter post-translational processes of the antigen binding protein, such as changing the number or position of glycosylation sites. In certain embodiments, antigen binding protein variants are prepared with the intent to modify those amino acid residues which are directly

involved in epitope binding. In other embodiments, modification of residues which are not directly involved in epitope binding or residues not involved in epitope binding in any way, is desirable, for purposes discussed herein. Mutagenesis within any of the CDR regions and/or framework regions is contemplated. Covariance analysis techniques can be employed by the skilled artisan to design useful modifications in the amino acid sequence of the antigen binding protein. *See, e.g.,* Choulier, *et al.*, Proteins 41:475-484, 2000; Demarest *et al.*, J. Mol. Biol. 335:41-48, 2004; Hugo *et al.*, Protein Engineering 16(5):381-86, 2003; Aurora *et al.*, US Patent Publication No. 2008/0318207 A1; Glaser *et al.*, US Patent Publication No. 2009/0048122 A1; Urech *et al.*, WO 2008/110348 A1; Borrás *et al.*, WO 2009/000099 A2. Such modifications determined by covariance analysis can improve potency, pharmacokinetic, pharmacodynamic, and/or manufacturability characteristics of an antigen binding protein.

[1281] The nucleic acid sequences of the present invention. As will be appreciated by those in the art, due to the degeneracy of the genetic code, an extremely large number of nucleic acids may be made, all of which encode the CDRs (and heavy and light chains or other components of the antigen binding proteins described herein) of the invention. Thus, having identified a particular amino acid sequence, those skilled in the art could make any number of different nucleic acids, by simply modifying the sequence of one or more codons in a way which does not change the amino acid sequence of the encoded protein.

[1282] The present invention also includes vectors comprising one or more nucleic acids encoding one or more components of the multispecific antibody constructs of the invention (e.g. variable regions, light chains, heavy chains, modified heavy chains, and Fd fragments). The term “vector” refers to any molecule or entity (e.g., nucleic acid, plasmid, bacteriophage or virus) used to transfer protein coding information into a host cell. Examples of vectors include, but are not limited to, plasmids, viral vectors, non-episomal mammalian vectors and expression vectors, for example, recombinant expression vectors. The term “expression vector” or “expression construct” as used herein refers to a recombinant DNA molecule containing a desired coding sequence and appropriate nucleic acid control sequences necessary for the expression of the operably linked coding sequence in a particular host cell. An expression vector can include, but is not limited to, sequences that affect or control transcription, translation, and, if introns are present, affect RNA splicing of a coding region operably linked thereto. Nucleic acid sequences necessary for expression in prokaryotes include a promoter, optionally an operator sequence, a ribosome binding site and possibly other sequences. Eukaryotic cells are known to utilize promoters, enhancers, and termination and polyadenylation signals. A secretory signal peptide sequence can also, optionally, be encoded by the expression vector, operably linked to the coding sequence of interest, so that the expressed polypeptide can be secreted by the recombinant host cell, for more facile isolation of the polypeptide of interest from the cell, if desired. For instance, in some embodiments, signal peptide sequences may be appended/fused to the amino terminus of any of the polypeptides sequences of the present invention. In certain

embodiments, a signal peptide having the amino acid sequence of MDMRVPAQLLGLLLLWLRGARC (SEQ ID NO: 894) is fused to the amino terminus of any of the polypeptide sequences of the present invention. In other embodiments, a signal peptide having the amino acid sequence of MAWALLLTLLTQGTGWSA (SEQ ID NO: 895) is fused to the amino terminus of any of the polypeptide sequences of the present invention. In still other embodiments, a signal peptide having the amino acid sequence of MTCSPLLLTLIHCTGWSA (SEQ ID NO: 896) is fused to the amino terminus of any of the polypeptide sequences of the present invention. Other suitable signal peptide sequences that can be fused to the amino terminus of the polypeptide sequences described herein include: MEAPAQLLFLLLLWLPDTTG (SEQ ID NO: 897), MEWTWRVFLVAAATGAHS (SEQ ID NO: 898), METPAQLLFLLLLWLPDTTG (SEQ ID NO: 899), METPAQLLFLLLLWLPDTTG (SEQ ID NO: 900), MKHLWFFLLLVAAAPRWVLS (SEQ ID NO: 901), and MEWSWVFLFSLVTTGVHS (SEQ ID NO: 902). Other signal peptides are known to those of skill in the art and may be fused to any of the polypeptide chains of the present invention, for example, to facilitate or optimize expression in particular host cells.

[1283] Typically, expression vectors used in the host cells to produce the bispecific antigen proteins of the invention will contain sequences for plasmid maintenance and for cloning and expression of exogenous nucleotide sequences encoding the components of the multispecific antibody constructs. Such sequences, collectively referred to as “flanking sequences,” in certain embodiments will typically include one or more of the following nucleotide sequences: a promoter, one or more enhancer sequences, an origin of replication, a transcriptional termination sequence, a complete intron sequence containing a donor and acceptor splice site, a sequence encoding a leader sequence for polypeptide secretion, a ribosome binding site, a polyadenylation sequence, a polylinker region for inserting the nucleic acid encoding the polypeptide to be expressed, and a selectable marker element. Each of these sequences is discussed below.

[1284] Optionally, the vector may contain a “tag”-encoding sequence, i.e., an oligonucleotide molecule located at the 5' or 3' end of the polypeptide coding sequence; the oligonucleotide tag sequence encodes polyHis (such as hexaHis (SEQ ID NO: 925)), FLAG, HA (hemagglutinin influenza virus), myc, or another “tag” molecule for which commercially available antibodies exist. This tag is typically fused to the polypeptide upon expression of the polypeptide, and can serve as a means for affinity purification or detection of the polypeptide from the host cell. Affinity purification can be accomplished, for example, by column chromatography using antibodies against the tag as an affinity matrix. Optionally, the tag can subsequently be removed from the purified polypeptide by various means such as using certain peptidases for cleavage.

[1285] Flanking sequences may be homologous (i.e., from the same species and/or strain as the host cell), heterologous (i.e., from a species other than the host cell species or strain), hybrid (i.e., a combination of flanking sequences from more than one source), synthetic or native. As such, the

source of a flanking sequence may be any prokaryotic or eukaryotic organism, any vertebrate or invertebrate organism, or any plant, provided that the flanking sequence is functional in, and can be activated by, the host cell machinery.

[1286] Flanking sequences useful in the vectors of this invention may be obtained by any of several methods well known in the art. Typically, flanking sequences useful herein will have been previously identified by mapping and/or by restriction endonuclease digestion and can thus be isolated from the proper tissue source using the appropriate restriction endonucleases. In some cases, the full nucleotide sequence of a flanking sequence may be known. Here, the flanking sequence may be synthesized using routine methods for nucleic acid synthesis or cloning.

[1287] Whether all or only a portion of the flanking sequence is known, it may be obtained using polymerase chain reaction (PCR) and/or by screening a genomic library with a suitable probe such as an oligonucleotide and/or flanking sequence fragment from the same or another species. Where the flanking sequence is not known, a fragment of DNA containing a flanking sequence may be isolated from a larger piece of DNA that may contain, for example, a coding sequence or even another gene or genes. Isolation may be accomplished by restriction endonuclease digestion to produce the proper DNA fragment followed by isolation using agarose gel purification, Qiagen® column chromatography (Chatsworth, CA), or other methods known to the skilled artisan. The selection of suitable enzymes to accomplish this purpose will be readily apparent to one of ordinary skill in the art.

[1288] An origin of replication is typically a part of those prokaryotic expression vectors purchased commercially, and the origin aids in the amplification of the vector in a host cell. If the vector of choice does not contain an origin of replication site, one may be chemically synthesized based on a known sequence, and ligated into the vector. For example, the origin of replication from the plasmid pBR322 (New England Biolabs, Beverly, MA) is suitable for most gram-negative bacteria, and various viral origins (e.g., SV40, polyoma, adenovirus, vesicular stomatitis virus (VSV), or papillomaviruses such as HPV or BPV) are useful for cloning vectors in mammalian cells. Generally, the origin of replication component is not needed for mammalian expression vectors (for example, the SV40 origin is often used only because it also contains the virus early promoter).

[1289] A transcription termination sequence is typically located 3' to the end of a polypeptide coding region and serves to terminate transcription. Usually, a transcription termination sequence in prokaryotic cells is a G-C rich fragment followed by a poly-T sequence. While the sequence is easily cloned from a library or even purchased commercially as part of a vector, it can also be readily synthesized using known methods for nucleic acid synthesis.

[1290] A selectable marker gene encodes a protein necessary for the survival and growth of a host cell grown in a selective culture medium. Typical selection marker genes encode proteins that (a) confer resistance to antibiotics or other toxins, e.g., ampicillin, tetracycline, or kanamycin for prokaryotic host cells; (b) complement auxotrophic deficiencies of the cell; or (c) supply critical

nutrients not available from complex or defined media. Specific selectable markers are the kanamycin resistance gene, the ampicillin resistance gene, and the tetracycline resistance gene. Advantageously, a neomycin resistance gene may also be used for selection in both prokaryotic and eukaryotic host cells.

[1291] Other selectable genes may be used to amplify the gene that will be expressed. Amplification is the process wherein genes that are required for production of a protein critical for growth or cell survival are reiterated in tandem within the chromosomes of successive generations of recombinant cells. Examples of suitable selectable markers for mammalian cells include dihydrofolate reductase (DHFR) and promoterless thymidine kinase genes. Mammalian cell transformants are placed under selection pressure wherein only the transformants are uniquely adapted to survive by virtue of the selectable gene present in the vector. Selection pressure is imposed by culturing the transformed cells under conditions in which the concentration of selection agent in the medium is successively increased, thereby leading to the amplification of both the selectable gene and the DNA that encodes another gene, such as one or more components of the multispecific antibody constructs described herein. As a result, increased quantities of a polypeptide are synthesized from the amplified DNA.

[1292] A ribosome-binding site is usually necessary for translation initiation of mRNA and is characterized by a Shine-Dalgarno sequence (prokaryotes) or a Kozak sequence (eukaryotes). The element is typically located 3' to the promoter and 5' to the coding sequence of the polypeptide to be expressed. In certain embodiments, one or more coding regions may be operably linked to an internal ribosome binding site (IRES), allowing translation of two open reading frames from a single RNA transcript.

[1293] In some cases, such as where glycosylation is desired in a eukaryotic host cell expression system, one may manipulate the various pre- or prosequences to improve glycosylation or yield. For example, one may alter the peptidase cleavage site of a particular signal peptide, or add prosequences, which also may affect glycosylation. The final protein product may have, in the -1 position (relative to the first amino acid of the mature protein) one or more additional amino acids incident to expression, which may not have been totally removed. For example, the final protein product may have one or two amino acid residues found in the peptidase cleavage site, attached to the amino-terminus. Alternatively, use of some enzyme cleavage sites may result in a slightly truncated form of the desired polypeptide, if the enzyme cuts at such area within the mature polypeptide.

[1294] Expression and cloning vectors of the invention will typically contain a promoter that is recognized by the host organism and operably linked to the molecule encoding the polypeptide. The term "operably linked" as used herein refers to the linkage of two or more nucleic acid sequences in such a manner that a nucleic acid molecule capable of directing the transcription of a given gene and/or the synthesis of a desired protein molecule is produced. For example, a control sequence in a vector that is "operably linked" to a protein coding sequence is ligated thereto so that expression of the protein coding sequence is achieved under conditions compatible with the transcriptional activity

of the control sequences. More specifically, a promoter and/or enhancer sequence, including any combination of cis-acting transcriptional control elements is operably linked to a coding sequence if it stimulates or modulates the transcription of the coding sequence in an appropriate host cell or other expression system.

[1295] Promoters are untranscribed sequences located upstream (i.e., 5') to the start codon of a structural gene (generally within about 100 to 1000 bp) that control transcription of the structural gene. Promoters are conventionally grouped into one of two classes: inducible promoters and constitutive promoters. Inducible promoters initiate increased levels of transcription from DNA under their control in response to some change in culture conditions, such as the presence or absence of a nutrient or a change in temperature. Constitutive promoters, on the other hand, uniformly transcribe a gene to which they are operably linked, that is, with little or no control over gene expression. A large number of promoters, recognized by a variety of potential host cells, are well known. A suitable promoter is operably linked to the DNA encoding e.g., heavy chain, light chain, modified heavy chain, or other component of the multispecific antibody constructs of the invention, by removing the promoter from the source DNA by restriction enzyme digestion and inserting the desired promoter sequence into the vector.

[1296] Suitable promoters for use with yeast hosts are also well known in the art. Yeast enhancers are advantageously used with yeast promoters. Suitable promoters for use with mammalian host cells are well known and include, but are not limited to, those obtained from the genomes of viruses such as polyoma virus, fowlpox virus, adenovirus (such as Adenovirus 2), bovine papilloma virus, avian sarcoma virus, cytomegalovirus, retroviruses, hepatitis-B virus and Simian Virus 40 (SV40). Other suitable mammalian promoters include heterologous mammalian promoters, for example, heat-shock promoters and the actin promoter.

[1297] Additional promoters which may be of interest include, but are not limited to: SV40 early promoter (Benoist and Chambon, 1981, *Nature* 290:304-310); CMV promoter (Thornsen et al., 1984, *Proc. Natl. Acad. U.S.A.* 81:659-663); the promoter contained in the 3' long terminal repeat of Rous sarcoma virus (Yamamoto et al., 1980, *Cell* 22:787-797); herpes thymidine kinase promoter (Wagner et al., 1981, *Proc. Natl. Acad. Sci. U.S.A.* 78: 1444-1445); promoter and regulatory sequences from the metallothionein gene Prinster et al., 1982, *Nature* 296:39-42); and prokaryotic promoters such as the beta-lactamase promoter (Villa-Kamaroff et al., 1978, *Proc. Natl. Acad. Sci. U.S.A.* 75:3727-3731); or the tac promoter (DeBoer et al., 1983, *Proc. Natl. Acad. Sci. U.S.A.* 80:21-25). Also of interest are the following animal transcriptional control regions, which exhibit tissue specificity and have been utilized in transgenic animals: the elastase I gene control region that is active in pancreatic acinar cells (Swift et al., 1984, *Cell* 38:639-646; Ornitz et al., 1986, *Cold Spring Harbor Symp. Quant. Biol.* 50:399-409; MacDonald, 1987, *Hepatology* 7:425-515); the insulin gene control region that is active in pancreatic beta cells (Hanahan, 1985, *Nature* 315: 115-122); the immunoglobulin gene

control region that is active in lymphoid cells (Grosschedl et al., 1984, Cell 38:647-658; Adames et al., 1985, Nature 318:533-538; Alexander et al., 1987, Mol. Cell. Biol. 7: 1436-1444); the mouse mammary tumor virus control region that is active in testicular, breast, lymphoid and mast cells (Leder et al., 1986, Cell 45:485-495); the albumin gene control region that is active in liver (Pinkert et al., 1987, Genes and Devel. 1 :268-276); the alpha-feto-protein gene control region that is active in liver (Krumlauf et al., 1985, Mol. Cell. Biol. 5: 1639-1648; Hammer et al., 1987, Science 253:53-58); the alpha 1-antitrypsin gene control region that is active in liver (Kelsey et al., 1987, Genes and Devel. 1: 161-171); the beta-globin gene control region that is active in myeloid cells (Mogrom et al., 1985, Nature 315:338-340; Kollias et al., 1986, Cell 46:89-94); the myelin basic protein gene control region that is active in oligodendrocyte cells in the brain (Readhead et al., 1987, Cell 48:703-712); the myosin light chain-2 gene control region that is active in skeletal muscle (Sani, 1985, Nature 314:283-286); and the gonadotropic releasing hormone gene control region that is active in the hypothalamus (Mason et al., 1986, Science 234: 1372-1378).

[1298] An enhancer sequence may be inserted into the vector to increase transcription of DNA encoding a component of the multispecific antibody constructs (e.g., light chain, heavy chain, modified heavy chain, Fd fragment) by higher eukaryotes. Enhancers are cis-acting elements of DNA, usually about 10-300 bp in length, that act on the promoter to increase transcription. Enhancers are relatively orientation and position independent, having been found at positions both 5' and 3' to the transcription unit. Several enhancer sequences available from mammalian genes are known (e.g., globin, elastase, albumin, alpha-feto-protein and insulin). Typically, however, an enhancer from a virus is used. The SV40 enhancer, the cytomegalovirus early promoter enhancer, the polyoma enhancer, and adenovirus enhancers known in the art are exemplary enhancing elements for the activation of eukaryotic promoters. While an enhancer may be positioned in the vector either 5' or 3' to a coding sequence, it is typically located at a site 5' from the promoter. A sequence encoding an appropriate native or heterologous signal sequence (leader sequence or signal peptide) can be incorporated into an expression vector, to promote extracellular secretion of the antibody. The choice of signal peptide or leader depends on the type of host cells in which the antibody is to be produced, and a heterologous signal sequence can replace the native signal sequence. Examples of signal peptides are described above. Other signal peptides that are functional in mammalian host cells include the signal sequence for interleukin-7 (IL-7) described in US Patent No. 4,965,195; the signal sequence for interleukin-2 receptor described in Cosman et al., 1984, Nature 312:768; the interleukin-4 receptor signal peptide described in EP Patent No. 0 367 566; the type I interleukin-1 receptor signal peptide described in U.S. Patent No. 4,968,607; the type II interleukin-1 receptor signal peptide described in EP Patent No. 0 460 846.

[1299] The expression vectors that are provided may be constructed from a starting vector such as a commercially available vector. Such vectors may or may not contain all of the desired flanking sequences. Where one or more of the flanking sequences described herein are not already present in

the vector, they may be individually obtained and ligated into the vector. Methods used for obtaining each of the flanking sequences are well known to one skilled in the art. The expression vectors can be introduced into host cells to thereby produce proteins, including fusion proteins, encoded by nucleic acids as described herein.

[1300] In certain embodiments, nucleic acids encoding the different components of the multispecific antibody constructs of the invention may be inserted into the same expression vector. For instance, the nucleic acid encoding an anti-first target antigen light chain can be cloned into the same vector as the nucleic acid encoding an anti- first target antigen heavy chain. In such embodiments, the two nucleic acids may be separated by an internal ribosome entry site (IRES) and under the control of a single promoter such that the light chain and heavy chain are expressed from the same mRNA transcript. Alternatively, the two nucleic acids may be under the control of two separate promoters such that the light chain and heavy chain are expressed from two separate mRNA transcripts. In some embodiments, nucleic acids encoding the anti- first target antigen light chain and heavy chain are cloned into one expression vector and the nucleic acids encoding the anti- second target antigen light chain and heavy chain are cloned into a second expression vector.

[1301] Similarly, for IgG-Fab multispecific antibody constructs, nucleic acids encoding each of the three components may be cloned into the same expression vector. In some embodiments, the nucleic acid encoding the light chain of the IgG-Fab molecule and the nucleic acid encoding the second polypeptide (which comprises the other half of the C-terminal Fab domain) are cloned into one expression vector, whereas the nucleic acid encoding the modified heavy chain (fusion protein comprising a heavy chain and half of a Fab domain) is cloned into a second expression vector. In certain embodiments, all components of the multispecific antibody constructs described herein are expressed from the same host cell population. For example, even if one or more components is cloned into a separate expression vector, the host cell is co-transfected with both expression vectors such that one cell produces all components of the multispecific antibody constructs.

[1302] After the vector has been constructed and the one or more nucleic acid molecules encoding the components of the multispecific antibody constructs described herein has been inserted into the proper site(s) of the vector or vectors, the completed vector(s) may be inserted into a suitable host cell for amplification and/or polypeptide expression. Thus, the present invention encompasses an isolated host cell comprising one or more expression vectors encoding the components of the multispecific antibody constructs. The term “host cell” as used herein refers to a cell that has been transformed, or is capable of being transformed, with a nucleic acid and thereby expresses a gene of interest. The term includes the progeny of the parent cell, whether or not the progeny is identical in morphology or in genetic make-up to the original parent cell, so long as the gene of interest is present. A host cell that comprises an isolated nucleic acid of the invention, in one embodiment operably linked to at least one expression control sequence (e.g. promoter or enhancer), is a “recombinant host cell.”

[1303] The transformation of an expression vector for an antigen binding protein into a selected host cell may be accomplished by well-known methods including transfection, infection, calcium phosphate co-precipitation, electroporation, microinjection, lipofection, DEAE-dextran mediated transfection, or other known techniques. The method selected will in part be a function of the type of host cell to be used. These methods and other suitable methods are well known to the skilled artisan, and are set forth, for example, in Sambrook et al., 2001, *supra*.

[1304] A host cell, when cultured under appropriate conditions, synthesizes an antigen binding protein that can subsequently be collected from the culture medium (if the host cell secretes it into the medium) or directly from the host cell producing it (if it is not secreted). The selection of an appropriate host cell will depend upon various factors, such as desired expression levels, polypeptide modifications that are desirable or necessary for activity (such as glycosylation or phosphorylation) and ease of folding into a biologically active molecule.

[1305] Exemplary host cells include prokaryote, yeast, or higher eukaryote cells. Prokaryotic host cells include eubacteria, such as Gram-negative or Gram-positive organisms, for example, *Enterobacteriaceae* such as *Escherichia*, e.g., *E. coli*, *Enterobacter*, *Erwinia*, *Klebsiella*, *Proteus*, *Salmonella*, e.g., *Salmonella typhimurium*, *Serratia*, e.g., *Serratia marcescans*, and *Shigella*, as well as *Bacillus*, such as *B. subtilis* and *B. licheniformis*, *Pseudomonas*, and *Streptomyces*. Eukaryotic microbes such as filamentous fungi or yeast are suitable cloning or expression hosts for recombinant polypeptides. *Saccharomyces cerevisiae*, or common baker's yeast, is the most commonly used among lower eukaryotic host microorganisms. However, a number of other genera, species, and strains are commonly available and useful herein, such as *Pichia*, e.g. *P. pastoris*, *Schizosaccharomyces pombe*; *Kluyveromyces*, *Yarrowia*; *Candida*; *Trichoderma reesia*; *Neurospora crassa*; *Schwanniomyces*, such as *Schwanniomyces occidentalis*; and filamentous fungi, such as, e.g., *Neurospora*, *Penicillium*, *Tolypocladium*, and *Aspergillus* hosts such as *A. nidulans* and *A. niger*.

[1306] Host cells for the expression of glycosylated antigen binding proteins can be derived from multicellular organisms. Examples of invertebrate cells include plant and insect cells. Numerous baculoviral strains and variants and corresponding permissive insect host cells from hosts such as *Spodoptera frugiperda* (caterpillar), *Aedes aegypti* (mosquito), *Aedes albopictus* (mosquito), *Drosophila melanogaster* (fruitfly), and *Bombyx mori* have been identified. A variety of viral strains for transfection of such cells are publicly available, e.g., the L-1 variant of *Autographa californica* NPV and the Bm-5 strain of *Bombyx mori* NPV.

[1307] Vertebrate host cells are also suitable hosts, and recombinant production of antigen binding proteins from such cells has become routine procedure. Mammalian cell lines available as hosts for expression are well known in the art and include, but are not limited to, immortalized cell lines available from the American Type Culture Collection (ATCC), including but not limited to Chinese hamster ovary (CHO) cells, including CHOK1 cells (ATCC CCL61), DXB-11, DG-44, and Chinese

hamster ovary cells/-DHFR (CHO, Urlaub *et al.*, Proc. Natl. Acad. Sci. USA 77: 4216, 1980); monkey kidney CV1 line transformed by SV40 (COS-7, ATCC CRL 1651); human embryonic kidney line (293 or 293 cells subcloned for growth in suspension culture, (Graham *et al.*, J. Gen Virol. 36: 59, 1977); baby hamster kidney cells (BHK, ATCC CCL 10); mouse sertoli cells (TM4, Mather, Biol. Reprod. 23: 243-251, 1980); monkey kidney cells (CV1 ATCC CCL 70); African green monkey kidney cells (VERO-76, ATCC CRL-1587); human cervical carcinoma cells (HELA, ATCC CCL 2); canine kidney cells (MDCK, ATCC CCL 34); buffalo rat liver cells (BRL 3A, ATCC CRL 1442); human lung cells (W138, ATCC CCL 75); human hepatoma cells (Hep G2, HB 8065); mouse mammary tumor (MMT 060562, ATCC CCL51); TRI cells (Mather *et al.*, Annals N.Y Acad. Sci. 383: 44-68, 1982); MRC 5 cells or FS4 cells; mammalian myeloma cells, and a number of other cell lines. In certain embodiments, cell lines may be selected through determining which cell lines have high expression levels and constitutively produce multispecific antibody constructs of the present invention. In another embodiment, a cell line from the B cell lineage that does not make its own antibody but has a capacity to make and secrete a heterologous antibody can be selected. CHO cells are host cells in some embodiments for expressing the multispecific antibody constructs of the invention.

[1308] Host cells are transformed or transfected with the above-described nucleic acids or vectors for production of multispecific antibody constructs and are cultured in conventional nutrient media modified as appropriate for inducing promoters, selecting transformants, or amplifying the genes encoding the desired sequences. In addition, novel vectors and transfected cell lines with multiple copies of transcription units separated by a selective marker are particularly useful for the expression of antigen binding proteins. Thus, the present invention also provides a method for preparing a multispecific antibody construct described herein comprising culturing a host cell comprising one or more expression vectors described herein in a culture medium under conditions permitting expression of the multispecific antibody construct encoded by the one or more expression vectors; and recovering the multispecific antibody construct from the culture medium.

[1309] The host cells used to produce the antigen binding proteins of the invention may be cultured in a variety of media. Commercially available media such as Ham's F10 (Sigma), Minimal Essential Medium ((MEM), (Sigma), RPMI-1640 (Sigma), and Dulbecco's Modified Eagle's Medium ((DMEM), Sigma) are suitable for culturing the host cells. In addition, any of the media described in Ham *et al.*, Meth. Enz. 58: 44, 1979; Barnes *et al.*, Anal. Biochem. 102: 255, 1980; U.S. Patent Nos. 4,767,704; 4,657,866; 4,927,762; 4,560,655; or 5,122,469; WO90103430; WO 87/00195; or U.S. Patent Re. No. 30,985 may be used as culture media for the host cells. Any of these media may be supplemented as necessary with hormones and/or other growth factors (such as insulin, transferrin, or epidermal growth factor), salts (such as sodium chloride, calcium, magnesium, and phosphate), buffers (such as HEPES), nucleotides (such as adenosine and thymidine), antibiotics (such as

GentamycinTM drug), trace elements (defined as inorganic compounds usually present at final concentrations in the micromolar range), and glucose or an equivalent energy source. Any other necessary supplements may also be included at appropriate concentrations that would be known to those skilled in the art. The culture conditions, such as temperature, pH, and the like, are those previously used with the host cell selected for expression, and will be apparent to the ordinarily skilled artisan.

[1310] Upon culturing the host cells, the multispecific antibody construct can be produced intracellularly, in the periplasmic space, or directly secreted into the medium. If the antigen binding protein is produced intracellularly, as a first step, the particulate debris, either host cells or lysed fragments, is removed, for example, by centrifugation or ultrafiltration. The bispecific antigen binding protein can be purified using, for example, hydroxyapatite chromatography, cation or anion exchange chromatography, or affinity chromatography, using the antigen(s) of interest or protein A or protein G as an affinity ligand. Protein A can be used to purify proteins that include polypeptides that are based on human $\gamma 1$, $\gamma 2$, or $\gamma 4$ heavy chains (Lindmark *et al.*, J. Immunol. Meth. 62: 1-13, 1983). Protein G is recommended for all mouse isotypes and for human $\gamma 3$ (Guss *et al.*, EMBO J. 5: 1567-1575, 1986). The matrix to which the affinity ligand is attached is most often agarose, but other matrices are available. Mechanically stable matrices such as controlled pore glass or poly(styrene-divinyl)benzene allow for faster flow rates and shorter processing times than can be achieved with agarose. Where the protein comprises a CH3 domain, the Bakerbond ABXTM resin (J. T. Baker, Phillipsburg, N.J.) is useful for purification. Other techniques for protein purification such as ethanol precipitation, Reverse Phase HPLC, chromatofocusing, SDS-PAGE, and ammonium sulfate precipitation are also possible depending on the particular multispecific antibody construct to be recovered.

[1311] The multispecific antibody constructs of the invention are useful for detecting target antigen(s) in biological samples and identification of cells or tissues that express the target antigen(s). The multispecific antibody constructs described herein can be used for diagnostic purposes to detect, diagnose, or monitor diseases and/or conditions associated with the target antigen(s). Also provided are methods for the detection of the presence of the target antigen(s) in a sample using classical immunohistological methods known to those of skill in the art (e.g., Tijssen, 1993, Practice and Theory of Enzyme Immunoassays, Vol 15 (Eds R.H. Burdon and P.H. van Knippenberg, Elsevier, Amsterdam); Zola, 1987, Monoclonal Antibodies: A Manual of Techniques, pp. 147-158 (CRC Press, Inc.); Jalkanen *et al.*, 1985, J. Cell. Biol. 101:976-985; Jalkanen *et al.*, 1987, J. Cell Biol. 105:3087-3096). The detection of either target can be performed *in vivo* or *in vitro*.

[1312] One embodiment provides the multispecific antibody construct of the invention or the multispecific antibody construct produced according to the process of the invention for use in the prevention, treatment or amelioration of a tumor or cancer disease or of a metastatic cancer disease.

[1313] According to a preferred embodiment of the invention said tumor or cancer disease is a solid tumor disease.

[1314] The formulations described herein are useful as pharmaceutical compositions in the treatment, amelioration and/or prevention of the pathological medical condition as described herein in a patient in need thereof. The term "treatment" refers to both therapeutic treatment and prophylactic or preventative measures. Treatment includes the application or administration of the formulation to the body, an isolated tissue, or cell from a patient who has a disease/disorder, a symptom of a disease/disorder, or a predisposition toward a disease/disorder, with the purpose to cure, heal, alleviate, relieve, alter, remedy, ameliorate, improve, or affect the disease, the symptom of the disease, or the predisposition toward the disease.

[1315] The term "amelioration" as used herein refers to any improvement of the disease state of a patient having a tumor or cancer or a metastatic cancer as specified herein below, by the administration of an multispecific antibody construct according to the invention to a subject in need thereof. Such an improvement may also be seen as a slowing or stopping of the progression of the tumor or cancer or metastatic cancer of the patient. The term "prevention" as used herein means the avoidance of the occurrence or re-occurrence of a patient having a tumor or cancer or a metastatic cancer as specified herein below, by the administration of an multispecific antibody construct according to the invention to a subject in need thereof.

[1316] The term "disease" refers to any condition that would benefit from treatment with the multispecific antibody construct or the pharmaceutical composition described herein. This includes chronic and acute disorders or diseases including those pathological conditions that predispose the mammal to the disease in question. A "neoplasm" is an abnormal growth of tissue, usually but not always forming a mass. When also forming a mass, it is commonly referred to as a "tumor". Neoplasms or tumors can be benign, potentially malignant (pre-cancerous), or malignant. Malignant neoplasms are commonly called cancer. They usually invade and destroy the surrounding tissue and may form metastases, i.e., they spread to other parts, tissues or organs of the body. Hence, the term "metastatic cancer" encompasses metastases to other tissues or organs than the one of the original tumor. Lymphomas and leukemias are lymphoid neoplasms. For the purposes of the present invention, they are also encompassed by the terms "tumor" or "cancer".

[1317] In a preferred embodiment of the invention, the tumor or cancer disease is a solid tumor disease and the metastatic cancer disease can be derived from any of the foregoing.

[1318] Preferred tumor or cancer diseases in connection with this invention are selected from a group consisting of breast cancer, Carcinoid, cervical cancer, colorectal cancer, endometrial cancer, gastric cancer, head and neck cancer, mesothelioma, liver cancer, lung cancer, ovarian cancer, pancreatic cancer, prostate cancer, skin cancer, renal cancer and stomach cancer. More preferably, the tumor or cancer disease, which is preferably a solid tumor disease, can be selected from the group consisting

of ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer. The metastatic cancer disease can be derived from any of the foregoing.

[1319] The invention also provides a method for the treatment or amelioration of tumor or cancer disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof the multispecific antibody construct of the invention or the multispecific antibody construct produced according to the process of the invention.

[1320] The terms "subject in need" or those "in need of treatment" includes those already with the disorder, as well as those in which the disorder is to be prevented. The subject in need or "patient" includes human and other mammalian subjects that receive either prophylactic or therapeutic treatment.

[1321] The multispecific antibody construct of the invention will generally be designed for specific routes and methods of administration, for specific dosages and frequencies of administration, for specific treatments of specific diseases, with ranges of bio-availability and persistence, among other things. The materials of the composition are preferably formulated in concentrations that are acceptable for the site of administration.

[1322] Formulations and compositions thus may be designed in accordance with the invention for delivery by any suitable route of administration. In the context of the present invention, the routes of administration include, but are not limited to

[1323] • topical routes (such as epicutaneous, inhalational, nasal, ophthalmic, auricular / aural, vaginal, mucosal);

[1324] • enteral routes (such as oral, gastrointestinal, sublingual, sublabial, buccal, rectal); and

[1325] • parenteral routes (such as intravenous, intraarterial, intraosseous, intramuscular, intracerebral, intracerebroventricular, epidural, intrathecal, subcutaneous, intraperitoneal, extra-amniotic, intraarticular, intracardiac, intradermal, intralesional, intrauterine, intravesical, intravitreal, transdermal, intranasal, transmucosal, intrasynovial, intraluminal).

[1326] The pharmaceutical compositions and the multispecific antibody construct of this invention are particularly useful for parenteral administration, e.g., subcutaneous or intravenous delivery, for example by injection such as bolus injection, or by infusion such as continuous infusion.

Pharmaceutical compositions may be administered using a medical device. Examples of medical devices for administering pharmaceutical compositions are described in U.S. Patent Nos. 4,475,196; 4,439,196; 4,447,224; 4,447,233; 4,486,194; 4,487,603; 4,596,556; 4,790,824; 4,941,880; 5,064,413; 5,312,335; 5,312,335; 5,383,851 ; and 5,399,163.

[1327] In particular, the present invention provides for an uninterrupted administration of the suitable composition. As a non-limiting example, uninterrupted or substantially uninterrupted, i.e. continuous administration may be realized by a small pump system worn by the patient for metering the influx of therapeutic agent into the body of the patient. The pharmaceutical composition comprising the

multispecific antibody construct of the invention can be administered by using said pump systems. Such pump systems are generally known in the art, and commonly rely on periodic exchange of cartridges containing the therapeutic agent to be infused. When exchanging the cartridge in such a pump system, a temporary interruption of the otherwise uninterrupted flow of therapeutic agent into the body of the patient may ensue. In such a case, the phase of administration prior to cartridge replacement and the phase of administration following cartridge replacement would still be considered within the meaning of the pharmaceutical means and methods of the invention together make up one "uninterrupted administration" of such therapeutic agent.

[1328] The continuous or uninterrupted administration of the multispecific antibody constructs of the invention may be intravenous or subcutaneous by way of a fluid delivery device or small pump system including a fluid driving mechanism for driving fluid out of a reservoir and an actuating mechanism for actuating the driving mechanism. Pump systems for subcutaneous administration may include a needle or a cannula for penetrating the skin of a patient and delivering the suitable composition into the patient's body. Said pump systems may be directly fixed or attached to the skin of the patient independently of a vein, artery or blood vessel, thereby allowing a direct contact between the pump system and the skin of the patient. The pump system can be attached to the skin of the patient for 24 hours up to several days. The pump system may be of small size with a reservoir for small volumes. As a non-limiting example, the volume of the reservoir for the suitable pharmaceutical composition to be administered can be between 0.1 and 50 ml.

[1329] The continuous administration may also be transdermal by way of a patch worn on the skin and replaced at intervals. One of skill in the art is aware of patch systems for drug delivery suitable for this purpose. It is of note that transdermal administration is especially amenable to uninterrupted administration, as exchange of a first exhausted patch can advantageously be accomplished simultaneously with the placement of a new, second patch, for example on the surface of the skin immediately adjacent to the first exhausted patch and immediately prior to removal of the first exhausted patch. Issues of flow interruption or power cell failure do not arise.

[1330] If the pharmaceutical composition has been lyophilized, the lyophilized material is first reconstituted in an appropriate liquid prior to administration. The lyophilized material may be reconstituted in, e.g., bacteriostatic water for injection (BWFI), physiological saline, phosphate buffered saline (PBS), or the same formulation the protein had been in prior to lyophilization.

[1331] The compositions of the present invention can be administered to the subject at a suitable dose which can be determined e.g. by dose escalating studies by administration of increasing doses of the multispecific antibody construct of the invention exhibiting cross-species specificity described herein to non-chimpanzee primates, for instance macaques. As set forth above, the multispecific antibody construct of the invention exhibiting cross-species specificity described herein can be advantageously used in identical form in preclinical testing in non-chimpanzee primates and as drug

in humans. The dosage regimen will be determined by the attending physician and clinical factors. As is well known in the medical arts, dosages for any one patient depend upon many factors, including the patient's size, body surface area, age, the particular compound to be administered, sex, time and route of administration, general health, and other drugs being administered concurrently.

[1332] The term "effective dose" or "effective dosage" is defined as an amount sufficient to achieve or at least partially achieve the desired effect. The term "therapeutically effective dose" is defined as an amount sufficient to cure or at least partially arrest the disease and its complications in a patient already suffering from the disease. Amounts or doses effective for this use will depend on the condition to be treated (the indication), the delivered multispecific antibody construct, the therapeutic context and objectives, the severity of the disease, prior therapy, the patient's clinical history and response to the therapeutic agent, the route of administration, the size (body weight, body surface or organ size) and/or condition (the age and general health) of the patient, and the general state of the patient's own immune system. The proper dose can be adjusted according to the judgment of the attending physician such that it can be administered to the patient once or over a series of administrations, and in order to obtain the optimal therapeutic effect.

[1333] A therapeutic effective amount of an multispecific antibody construct of the invention preferably results in a decrease in severity of disease symptoms, an increase in frequency or duration of disease symptom-free periods or a prevention of impairment or disability due to the disease affliction. For treating MSLN-expressing tumors, a therapeutically effective amount of the multispecific antibody construct of the invention, e.g. an anti-MSLN /anti-CD40 multispecific antibody construct, preferably inhibits cell growth or tumor growth by at least about 20%, at least about 40%, at least about 50%, at least about 60%, at least about 70%, at least about 80%, or at least about 90% relative to untreated patients. The ability of a compound to inhibit tumor growth may be evaluated in an animal model predictive of efficacy in human tumors.

[1334] The pharmaceutical composition can be administered as a sole therapeutic or in combination with additional therapies such as anti-cancer therapies as needed, e.g. other proteinaceous and non-proteinaceous drugs. These drugs may be administered simultaneously with the composition comprising the multispecific antibody construct of the invention as defined herein or separately before or after administration of said multispecific antibody construct in timely defined intervals and doses.

[1335] The term "effective and non-toxic dose" as used herein refers to a tolerable dose of an inventive multispecific antibody construct which is high enough to cause depletion of pathologic cells, tumor elimination, tumor shrinkage or stabilization of disease without or essentially without major toxic effects. Such effective and non-toxic doses may be determined e.g. by dose escalation studies described in the art and should be below the dose inducing severe adverse side events (dose limiting toxicity, DLT).

[1336] The term "toxicity" as used herein refers to the toxic effects of a drug manifested in adverse events or severe adverse events. These side events might refer to a lack of tolerability of the drug in general and/or a lack of local tolerance after administration. Toxicity could also include teratogenic or carcinogenic effects caused by the drug.

[1337] The term "safety", "in vivo safety" or "tolerability" as used herein defines the administration of a drug without inducing severe adverse events directly after administration (local tolerance) and during a longer period of application of the drug. "Safety", "in vivo safety" or "tolerability" can be evaluated e.g. at regular intervals during the treatment and follow-up period. Measurements include clinical evaluation, e.g. organ manifestations, and screening of laboratory abnormalities. Clinical evaluation may be carried out and deviations to normal findings recorded/coded according to NCI-CTC and/or MedDRA standards. Organ manifestations may include criteria such as allergy/immunology, blood/bone marrow, cardiac arrhythmia, coagulation and the like, as set forth e.g. in the Common Terminology Criteria for adverse events v3.0 (CTCAE). Laboratory parameters which may be tested include for instance hematology, clinical chemistry, coagulation profile and urine analysis and examination of other body fluids such as serum, plasma, lymphoid or spinal fluid, liquor and the like. Safety can thus be assessed e.g. by physical examination, imaging techniques (i.e. ultrasound, x-ray, CT scans, Magnetic Resonance Imaging (MRI), other measures with technical devices (i.e. electrocardiogram), vital signs, by measuring laboratory parameters and recording adverse events. For example, adverse events in non-chimpanzee primates in the uses and methods according to the invention may be examined by histopathological and/or histochemical methods. The above terms are also referred to e.g. in the Preclinical safety evaluation of biotechnology- derived pharmaceuticals S6; ICH Harmonised Tripartite Guideline; ICH Steering Committee meeting on July 16, 1997.

[1338] In a further embodiment, the invention provides a kit comprising an multispecific antibody construct of the invention, an multispecific antibody construct produced according to the process of the invention, a polynucleotide of the invention, a vector of the invention, and/or a host cell of the invention.

[1339] In the context of the present invention, the term "kit" means two or more components - one of which corresponding to the multispecific antibody construct, the pharmaceutical composition, the vector or the host cell of the invention - packaged together in a container, recipient or otherwise. A kit can hence be described as a set of products and/or utensils that are sufficient to achieve a certain goal, which can be marketed as a single unit.

[1340] The kit may comprise one or more recipients (such as vials, ampoules, containers, syringes, bottles, bags) of any appropriate shape, size and material (preferably waterproof, e.g. plastic or glass) containing the multispecific antibody construct or the pharmaceutical composition of the present invention in an appropriate dosage for administration (see above). The kit may additionally contain

directions for use (e.g. in the form of a leaflet or instruction manual), means for administering the multispecific antibody construct of the present invention such as a syringe, pump, infuser or the like, means for reconstituting the multispecific antibody construct of the invention and/or means for diluting the multispecific antibody construct of the invention.

[1341] The invention also provides kits for a single-dose administration unit. The kit of the invention may also contain a first recipient comprising a dried / lyophilized multispecific antibody construct and a second recipient comprising an aqueous formulation. In certain embodiments of this invention, kits containing single-chambered and multi-chambered pre-filled syringes (e.g., liquid syringes and lyosyringes) are provided.

[1342] Examples

[1343] **Generation and characterization of anti-mesothelin antibodies**

[1344] *Anti-mesothelin antibody generation.*

[1345] Fully human antibodies to human mesothelin were generated as previously described (US20170029502A1) or by immunizing XENOMOUSE® transgenic mice (U.S. Pat. Nos. 6,114,598; 6,162,963; 6,833,268; 7,049,426; 7,064,244, which are incorporated herein by references in their entirety; Green et al., 1994, *Nature Genetics* 7:13-21; Mendez et al., 1997, *Nature Genetics* 15:146-156; Green and Jakobovitis, 1998, *J. Ex. Med.*, 188:483-495; Kellerman and Green, *Current Opinion in Biotechnology* 13, 593-597, 2002). Animals from the XMG4-K and XMG4-KL XENOMOUSE® strains were used for these immunizations. Animals were immunized with alternating soluble human mesothelin-His and cynomolgus mesothelin-His. Animals with the highest antigen-specific serum native titers directed against human mesothelin and cynomolgus mesothelin were used for hybridoma generation (Kohler and Milstein, 1975). Pooled lymphocytes from spleen and/or draining lymph node (from each harvest) were dissociated from lymphoid tissue by grinding in a suitable medium (for example, Dulbecco's Modified Eagle Medium (DMEM); Invitrogen, Carlsbad, CA). B cells were selected and/or expanded using standard methods, and fused with a suitable fusion partner using techniques that were known in the art. Hybridoma supernatants with binding to human mesothelin and cynomolgus mesothelin were then selected for further characterization.

[1346] *Sequencing of anti-mesothelin antibodies.*

[1347] For XENOMOUSE®-derived antibodies, RNA (total or mRNA) was purified from wells containing the anti-mesothelin antibody-producing hybridoma cells using a Qiagen RNeasy mini or the Invitrogen mRNA catcher plus kit. Purified RNA was used to amplify the antibody heavy and light chain variable region (V) genes using cDNA synthesis via reverse transcription, followed by a polymerase chain reaction (RT-PCR). The fully human antibody gamma heavy chain was obtained using the Qiagen One Step Reverse Transcriptase PCR kit (Qiagen). The fully human kappa light chain was obtained using the Qiagen One Step Reverse Transcriptase PCR kit (Qiagen). Amino acid

sequences were deduced from the corresponding nucleic acid sequences bioinformatically. The derived amino acid sequences were then analyzed to determine the germline sequence origin of the antibodies and to identify deviations from the germline sequence. The amino acid sequences corresponding to complementary determining regions (CDRs) of the sequenced antibodies were aligned and these alignments were used to group the clones by similarity.

[1348] *Antibody production.*

[1349] Selected and sequenced anti-mesothelin antibody heavy and light chains were subcloned into mammalian expression vectors and individual antibodies were produced and purified. All antibodies were reformatted with human IgG1 SEFL2 (REF) heavy chain sequences.

[1350] *Binding of antibodies to human and cynomolgus monkey mesothelin.*

[1351] Binding of anti-mesothelin antibodies to human mesothelin was confirmed by flow cytometry on CHO cells engineered to express human mesothelin (huMSLN-CHO). Human MSLN-transfected CHO cells were incubated with various concentrations of purified anti-mesothelin antibodies, washed, and labeled with a fluorescent-conjugated secondary antibody specific for human IgG. Cells were then analyzed by flow cytometry. The percentage of fluorescent positive cells was plotted against antibody concentration (Figure 1) and the EC50 was determined using Prism GraphPad software (Table 1). The binding affinity of anti-human mesothelin antibodies to soluble forms of recombinant human and cynomolgus monkey (cyno) mesothelin was also measured by Octet assay (Table 1), quantifying association rate (Kon), disassociation rate (Kdis), and equilibrium binding constant (KD). These data demonstrated binding of all anti-MSLN antibodies to both human and cynomolgus monkey mesothelin.

[1352] Table 1. Binding of anti-MSLN antibodies to human and cyno Mesothelin (MSLN)

PL#	BioReg ID	Clone	On Cell Binding human MSLN EC50 (uM)	Octet Binding human MSLN			Octet Binding Cyno MSLN		
				Kon (1/Ms)	Kdis (1/s)	KD (nM)	Kon (1/Ms)	Kdis (1/s)	KD (nM)
PL-55235	4553	6F4	0.01	4.50E+05	8.77E-05	0.19	3.35E+05	1.75E-03	5.2
PL-55238	4559	7G11	0.003	5.48E+05	1.85E-04	0.34	2.94E+05	1.88E-03	6.4
PL-54455	3966	4H6	0.002	3.58E+05	5.16E-05	0.14	3.92E+05	7.64E-04	1.9
PL-54456	3967	4G12	0.001	8.42E+04	1.62E-05	0.19	1.24E+05	5.60E-04	4.5

[1353] *Generation and characterization of anti-CD40 agonist antibodies*

[1354] *Anti-CD40 antibody generation.*

[1355] Fully human antibodies to human CD40 were generated by immunizing XENOMOUSE® transgenic mice (U.S. Pat. Nos. 6,114,598; 6,162,963; 6,833,268; 7,049,426; 7,064,244, which are incorporated herein by references in their entirety; Green et al., 1994, *Nature Genetics* 7:13-21; Mendez et al., 1997, *Nature Genetics* 15:146-156; Green and Jakobovitis, 1998, *J. Ex. Med.*, 188:483-

495; Kellerman and Green, *Current Opinion in Biotechnology* 13, 593-597, 2002). Animals from the XMG4-K and XMG4-KL XENOMOUSE® strains were used for these immunizations. Multiple immunogens and routes of immunization were used to generate anti-human CD40 immune responses. For soluble recombinant protein immunizations, mice were immunized with alternating soluble human CD40-Fc and cynomolgus CD40-Fc. For cell-based immunizations, CHO-S cells were transiently transfected with either wild type human CD40 or cynomolgus CD40 as a source of immunogen. Animals were immunized with either of these transiently transfected CHO cells. Animals with the highest antigen-specific serum native titers directed against human CD40 and cynomolgus CD40 were used for hybridoma generation (Kohler and Milstein, 1975). Pooled lymphocytes from spleen and/or draining lymph node (from each harvest) were dissociated from lymphoid tissue by grinding in a suitable medium (for example, Dulbecco's Modified Eagle Medium (DMEM); Invitrogen, Carlsbad, CA). B cells were selected and/or expanded using standard methods and fused with a suitable fusion partner using techniques that were known in the art. Hybridoma supernatants with binding to human and cynomolgus monkey CD40 were identified.

[1356] *Antibody production.*

[1357] Select hybridoma supernatants showing binding to human and cynomolgus monkey CD40 were used to produce purified anti-CD40 antibodies using techniques that were known to the art (Table 2).

[1358] Table 2. Anti-CD40 antibodies

Antibody ID	HC Sequence		IgG Subclass
	VH Germline	HC CDR3	
29H10	VH1 1-02/D5 5-18 RF1/JH5	ERISMVRGVGHNWFAP (SEQ ID NO: 150)	IgG4
4G7	VH1 1-02/D3 3-16 RF3/JH4	EKITMTGIYFDY (SEQ ID NO: 198)	IgG4
33H6	VH1 1-02/D1 1-7 RF3/JH6	EKPRYFDSFYLYMDV (SEQ ID NO: 210)	IgG4
35F11	VH3 3-33/D5 5-24 RF3/JH4	DGRNYVYFDN (SEQ ID NO: 186)	IgG4
30A12	VH3 3-23/D6 6-13 RF1/JH4	GYSNSWWYFDY (SEQ ID NO: 180)	IgG4
36F3	VH6 6-01/D6 6-13 RF2/JH4	GAAPFDY (SEQ ID NO: 144)	IgG4
39C2	VH1 1-02/D1 1-1 RF1/JH4	ERCRTTNCYLDY (SEQ ID NO: 192)	IgG4
33H9	VH3 3-33/D7 7-27 RF1/JH6	GGGHWNYEGHYGMDV (SEQ ID NO: 138)	IgG4
37A6	VH3 3-11/D6 6-6 RF2/JH4	DLAAGATGGLDC (SEQ ID NO: 174)	IgG4

[1359] *Anti-CD40 antibody binding to human and cynomolgus monkey CD40.*

[1360] The binding affinity of anti-human CD40 antibodies to human and cynomolgus monkey CD40 was measured by Octet assay (Table 3), quantifying association rate (K_{on}), disassociation rate (K_{dis}), and equilibrium binding constant (K_D). Cross-reactivity of anti-CD40 antibodies to related TNF receptor superfamily (TNFRSF) members was evaluated by expressing human TNFR1, TNFR2, TNFR10, or TNFR14 on HEK293 cells by transient transfection of expression vectors encoding these

genes. Binding of indicated anti-human CD40 antibody to these transfected cells was determined by flow cytometry. Evaluating geometric mean fluorescent intensity relative to control cells demonstrated minimal binding of anti-CD40 to other TNFRSF members (Table 4). Cross-reactivity of anti-CD40 antibodies to cynomolgus monkey and mouse CD40 was evaluated by expressing cynomolgus monkey or mouse CD40 on HEK293 cells by transient transfection of expression vectors encoding these genes. Binding of indicated anti-human CD40 antibody to cells overexpressing cynomolgus monkey and mouse CD40 was determined by flow cytometry. Evaluating geometric mean fluorescent intensity relative to control cells demonstrated robust binding of anti-CD40 to cynomolgus monkey (cyno) CD40, but no binding to mouse CD40 (Table 5). Taken together these data demonstrate strong and equivalent binding of anti-CD40 antibodies to human and cynomolgus monkey CD40, but minimal binding to related TNFRSF members or mouse CD40.

Table 3. Binding affinity of anti-CD40 antibodies to human and cyno CD40

Antibody ID	Human CD40			Cyno CD40		
	kon(1/Ms)	kdis(1/s)	KD (nM)	kon(1/Ms)	kdis(1/s)	KD (nM)
29H10	3.98E+05	1.64E-04	0.41	6.12E+05	2.44E-04	0.39
4G7	1.23E+06	1.08E-03	0.88	1.13E+06	1.93E-03	1.72
33H6	5.09E+05	6.80E-04	1.34	5.56E+05	5.77E-04	1.04
35F11	5.56E+05	3.10E-04	0.55	5.06E+05	3.14E-04	0.62
30A12	5.33E+05	3.68E-04	0.69	5.44E+05	3.06E-04	0.56
36F3	6.45E+04	4.43E-04	6.87	5.22E+04	2.10E-04	4.02
39C2	5.99E+05	6.14E-04	1.02	5.18E+05	3.76E-04	0.73
33H9	4.07E+05	2.38E-04	0.58	4.02E+05	1.68E-04	4.17
37A6	6.12E+05	6.72E-04	1.10	5.31E+05	3.87E-04	0.73

Table 4. Absence of anti-CD40 antibody binding to related TNF receptor superfamily members (fold over control values)

Antibody ID	TNFR1	TNFR2	TNFR10	TNFR14
29H10	0.98	0.88	0.89	0.84
4G7	0.97	0.90	0.87	0.89
33H6	1.00	0.81	0.77	0.80
35F11	1.00	0.85	0.73	0.74
30A12	1.07	0.89	0.91	0.96
36F3	0.99	0.92	0.86	0.88
39C2	1.00	0.90	0.86	0.89
33H9	1.03	0.93	0.89	0.88
37A6	0.99	0.87	0.88	0.87

Table 5. Species cross-reactivity of anti-CD40 antibodies to cyno and mouse CD40 (fold over control values)

Antibody ID	Cyno	Mouse
29H10	71	0.8
4G7	57	0.9
33H6	74	0.9
35F11	76	0.8
30A12	71	1.3
36F3	82	0.9
39C2	87	0.9
33H9	73	0.9
37A6	92	0.9

[1361] *Cross-linking-dependent activation of CD40 by anti-CD40 antibodies.*

[1362] Stimulation of the CD40 receptor on human B cells results in activation and proliferation. Purified primary human B cells were seeded into 384-well assay plates and treated with IL-4/IL-21 as co-mitogens, varying concentrations of anti-CD40 antibodies, and the presence (Figure 2A) or absence (Figure 2B) of protein G. A single molecule of Protein G is capable of binding multiple IgG molecules and can thus cross-link antibodies in solution. Cells were incubated for 5 days and cell proliferation was examined using CellTiter-Glo. An irrelevant IgG4 isotype antibody was included as a negative control as all anti-CD40 antibodies examined were IgG4. EC50 and percent maximum activity were calculated (Table 6). All anti-CD40 antibodies induced robust proliferation of B cells in the presence of protein G but had minimal activity in the absence of protein G, indicating that anti-CD40 antibody binding to CD40 alone was insufficient to stimulate CD40 and that additional cross-linking of the antibodies was required.

Table 6. Activity of anti-CD40 antibodies with and without crosslinking in human B cell functional assay

Antibody ID	No Crosslinking EC50 (nM)	No Crosslinking Max Fold Change	With Crosslinking EC50 (nM)	With Crosslinking Max Fold Change
29H10	>16.7	1.35	0.235	18.22
4G7	>16.7	1.19	0.430	18.05
33H6	>16.7	1.21	0.237	18.24
35F11	>16.7	1.33	0.371	24.16
30A12	>16.7	1.45	0.362	23.78
36F3	>16.7	1.83	0.558	16.85
39C2	>16.7	1.24	0.529	23.01
33H9	>16.7	1.35	0.344	23.17
37A6	>16.7	1.19	0.504	27.09

[1363] *Effect of anti-CD40 antibodies on binding to human CD40 ligand to human CD40.*

[1364] To evaluate the ability of the anti-CD40 antibodies to block CD40 ligand interactions with CD40, a flow cytometry-based ligand binding assay was performed to measure the binding of fluorescent-labeled soluble human CD40L to human CD40 over-expressed on 293T cells by transient transfection with an expression vector encoding this gene. Percent ligand binding inhibition was calculated as [(CD40L binding gMFI in the absence of anti-CD40) – (CD40L binding gMFI in the presence of anti-CD40)] / (CD40L binding gMFI in the absence of anti-CD40). gMFI: geometric mean fluorescent intensity. All anti-CD40 antibodies show minimal effects on the ability of CD40L to bind to the CD40 receptor (Table 7).

Table 7. Anti-CD40 antibodies do not block CD40-CD40L interactions

Antibody ID	Ligand Binding Inhibition (%)				Ligand Blocker
	20 ug/ml	10 ug/ml	5 ug/ml	2.5 ug/ml	
29H10	-7%	1%	-8%	-11%	No
4G7	3%	14%	24%	21%	No
33H6	7%	11%	4%	-8%	No
35F11	8%	51%	-5%	-15%	No
30A12	-3%	-8%	-15%	-16%	No
36F3	1%	14%	10%	1%	No
39C2	2%	3%	-6%	7%	No
33H9	1%	3%	9%	-3%	No
37A6	12%	6%	2%	-19%	No

[1365] *Effects of anti-human CD40 antibody on CD40L-induced B cell activation.*

[1366] Some antibodies specific for TNFRSF members have been shown to potentiate activation of the receptor by its normal ligand, potentially by clustering the receptor and decreasing the threshold for ligand-induced activation. To evaluate the ability of the anti-CD40 antibodies to potentiate CD40 ligand activity, purified primary human B cells were treated with various concentrations of recombinant human CD40L in the presence or absence of 1 ug/ml of the indicated CD40 antibody. Cells were incubated for 5 days and cell proliferation was examined using CellTiter-Glo. None of the anti-CD40 antibodies induced a significant change in the EC50 of CD40 ligand to stimulate proliferation of human B cells (Table 8).

Table 8. Anti-CD40 antibodies do not potentiate activity of CD40L

Antibody ID	CD40L EC50 (ug/ml)
No Antibody	0.001085
29H10	0.000715
4G7	0.00104
33H6	0.001135
35F11	0.006655
30A12	0.00081
36F3	0.012975

39C2	0.000985
33H9	No Activity
37A6	0.00139

[1367] *Sequencing of anti-CD40 antibodies.*

[1368] RNA (total or mRNA) was purified from wells containing the anti-mesothelin antibody-producing hybridoma cells using a Qiagen RNeasy mini or the Invitrogen mRNA catcher plus kit. Purified RNA was used to amplify the antibody heavy and light chain variable region (V) genes using cDNA synthesis via reverse transcription, followed by a polymerase chain reaction (RT-PCR). The fully human antibody gamma heavy chain was obtained using the Qiagen One Step Reverse Transcriptase PCR kit (Qiagen). The fully human kappa light chain was obtained using the Qiagen One Step Reverse Transcriptase PCR kit (Qiagen). Amino acid sequences were deduced from the corresponding nucleic acid sequences bioinformatically. The derived amino acid sequences were then analyzed to determine the germline sequence origin of the antibodies and to identify deviations from the germline sequence. The amino acid sequences corresponding to complementary determining regions (CDRs) of the sequenced antibodies were aligned and these alignments were used to group the clones by similarity.

[1369] *Optimization of anti-CD40 antibody sequences.*

[1370] Engineering of select anti-CD40 antibodies was performed to remove potential sequence liabilities in the antibody heavy and light chain variable regions by standard recombinant DNA techniques. These antibodies were produced by over-expression in cell lines and purified using techniques that were known in the art. Purified antibodies were then evaluated in a human B cell proliferation assay with and without protein G cross-linking, as described above. EC₅₀ and percent maximum activity were calculated (Table 9), showing that many of the engineered antibody variants maintained similar cross-linking dependent agonist activity as their parental antibody sequence.

Table 9. Functional activity of engineered variants of anti-CD40 antibodies

Antibody Name	Antibody ID	No Crosslinking EC ₅₀ (nM)	No Crosslinking % Max Activity	With Crosslinking EC ₅₀ (nM)	With Crosslinking % Max Activity
29H10	4877-1	> 16.7	4.4	0.37	65.9
30A12	4878-1	> 16.7	8.6	0.40	89.1
30A12.001	4894-1	> 5.93	12.2	0.08	87.2
30A12.002	4879-1	> 16.7	19.0	0.36	89.2
30A12.003	4880-1	> 16.7	1.2	0.41	83.3
33H6	4883-1	> 16.7	6.5	0.26	73.4
33H6.001	4884-1	> 16.7	10.8	0.46	92.7
33H6.002	4885-1	> 16.7	11.5	0.40	81.2
33H6.003	4895-1	> 6.22	6.5	0.43	61.4

33H6.004	4886-1	> 16.7	20.9	0.44	73.5
33H9	4897-1	> 13.4	2.3	0.19	88.9
35F11	4868-1	> 16.7	4.4	0.30	119.7
35F11.001	5052-2	> 16.7	12.5	0.85	96.5
35F11.002	5053-2	> 16.7	6.4	0.77	84.2
35F11.003	5054-2	> 16.7	10.5	0.60	74.6
35F11.004	5055-2	> 16.7	10.0	0.90	83.7
35F11.005	5056-2	> 16.7	5.4	0.68	77.8
35F11.006	5057-2	> 16.7	4.8	0.49	109.0
36F3	4891-1	> 16.7	8.2	0.30	81.9
37A6	4865-1	> 16.7	0.4	0.30	80.9
37A6.001	4866-1	> 16.7	-0.2	0.32	86.7
37A6.002	4892-1	> 5.87	-1.9	0.11	87.3
37A6.003	4867-1	> 16.7	-1.1	0.21	73.0
39C2	4887-1	> 16.7	6.1	0.21	76.6
39C2.001	4888-1	> 16.7	1.2	> 16.7	2.2
39C2.002	4889-1	> 16.7	21.9	4.36	86.5
39C2.003	4890-1	> 16.7	4.6	> 16.7	1.7
4G7	4869-1	> 16.7	6.0	0.50	70.5
4G7.001	4893-1	> 5.99	2.7	0.83	80.5
4G7.002	4870-1	> 16.7	2.1	0.62	72.2
4G7.003	4871-1	> 16.7	3.9	0.77	89.1
4G7.004	4872-1	> 16.7	2.3	1.01	96.8

[1371] Generation and characterization of mesothelin-targeted CD40 agonist bivalent bi-specific antibodies

[1372] Bispecific antibody production.

[1373] In order to generate a mesothelin-dependent CD40 agonist, bivalent bispecific antibodies capable of binding to both human mesothelin and human CD40 were generated in either an IgG-Fab format (Figure 3A) or an IgG-scFv format (Figure 3B) using the variable region binding domains of the anti-mesothelin and anti-CD40 antibodies described above.

[1374] Generation and evaluation of mesothelin x CD40 bispecific agonist antibodies in the IgG-Fab format.

[1375] A panel of mesothelin x CD40 bispecific antibodies were generated in the IgG-Fab format (Table 10) and evaluated for binding to soluble forms of human and cynomolgus monkey CD40 and human and cynomolgus monkey mesothelin using the Octet assay (Table 11). Association rate (K_{on}), disassociation rate (K_{dis}), and equilibrium binding constant (K_D) were calculated. The majority of bispecific antibodies showed high affinity binding to both human and cynomolgus monkey CD40 and mesothelin. These bispecific antibodies were next evaluated for their ability to induce mesothelin-dependent activation of human B cells. CHO cells expressing human mesothelin were seeded into a 96 well plate. The next day varying concentrations of the mesothelin x CD40 IgG-Fab antibodies

were added to the wells along with purified human B cells and the plates were incubated for an additional 48 hours. Upregulation of CD86, a marker of CD40-mediated B cell activation, was quantified on B cells by flow cytometry. EC50 values for CD86 upregulation were calculated and demonstrate that the majority of mesothelin x CD40 IgG-Fab bispecific antibodies were able to B cell activation in the presence of CHO cells expressing human mesothelin (Table 12).

Table 10. Anti-CD40xMSLN IgG-Fab molecules

Antibody ID	IgG (MSLN)	Fab (CD40)	IgG (CD40)	Fab (MSLN)
13468-1	4G12	30A12		
5966-1	4H6	37A6		
5978-1	7G11	33H9		
13473-1			30A12	4G12
13474-1			35F11	4G12
13469-1	4G12	35F11		
5967-1	6F4	29H10		
6057-1	7G11	37A6		
5991-1			30A12	6F4
6003-1			35F11	6F4
13470-1	4G12	37A6		
5968-1	6F4	30A12		
6058-1	7G11	39C2		
5992-1			30A12	7G11
13475-1			36F3	4G12
13471-1	4G12	39C2		
5970-1	6F4	33H9		
5982-1			4G7	4H6
6007-1			36F3	6F4
5960-1	4H6	29H10		
5972-1	6F4	36F3		
5983-1			4G7	6F4
5995-1			33H6	6F4
5946-1			37A6	4H6
5961-1	4H6	30A12		
5973-1	6F4	37A6		
13472-1			29H10	4G12
5998-1			33H9	4H6
5947-1			37A6	7G11
5963-1	4H6	33H9		
5975-1	7G11	29H10		
5987-1			29H10	6F4
5999-1			33H9	6F4
5965-1	4H6	36F3		

5976-1	7G11	30A12		
5988-1			29H10	7G11
6000-1			33H9	7G11

Table 11. Anti-CD40xMSLN IgG-Fab molecule binding to human and cynomolgus monkey CD40 and mesothelin

Antibody ID	huCD40(1-193)			cynoCD40(1-193)			huMSLN(296-598)			cynoMSLN(296-598)		
	K_D (nM)	k_a (M ⁻¹ s ⁻¹)	k_d (s ⁻¹)	K_D (nM)	k_a (M ⁻¹ s ⁻¹)	k_d (s ⁻¹)	K_D (nM)	k_a (M ⁻¹ s ⁻¹)	k_d (s ⁻¹)	K_D (nM)	k_a (M ⁻¹ s ⁻¹)	k_d (s ⁻¹)
5959-1	0.4	5.11E+05	2.14E-04	0.7	6.21E+05	4.28E-04	<0.1	2.29E+05	>1.00E-05	6.0	1.10E+05	6.58E-04
5960-1	0.4	4.20E+05	1.82E-04	0.5	3.55E+05	1.76E-04	0.2	7.98E+05	1.79E-04	2.5	3.66E+05	9.16E-04
5961-1	0.6	6.22E+05	3.42E-04	0.8	3.39E+05	2.66E-04	<0.1	6.80E+05	>1.00E-05	2.7	3.56E+05	9.58E-04
5963-1	<0.1	8.85E+04	>1.00E-05	0.8	1.47E+05	1.15E-04	0.2	8.46E+05	1.56E-04	2.9	3.10E+05	9.03E-04
5965-1	2.5	1.19E+05	2.97E-04	2.7	9.51E+04	2.55E-04	0.2	8.03E+05	1.40E-04	2.1	4.24E+05	8.96E-04
5966-1	0.7	5.84E+05	4.03E-04	0.9	4.20E+05	3.58E-04	0.2	8.78E+05	1.76E-04	2.5	3.78E+05	9.40E-04
5967-1	0.8	3.75E+05	2.86E-04	<0.1	1.83E+05	>1.00E-05	0.3	9.12E+05	2.89E-04	6.4	3.71E+05	2.36E-03
5968-1	0.4	6.50E+05	2.56E-04	0.7	5.98E+05	4.14E-04	0.3	1.03E+06	3.53E-04	5.1	5.56E+05	2.82E-03
5970-1	1.6	2.45E+05	3.95E-04	1.3	1.63E+05	2.08E-04	0.3	7.74E+05	2.58E-04	5.8	4.31E+05	2.48E-03
5972-1	4.0	1.34E+05	5.31E-04	1.7	8.06E+04	1.37E-04	0.5	5.68E+05	2.70E-04	4.6	4.54E+05	2.10E-03
5973-1	0.6	5.41E+05	3.25E-04	0.6	6.69E+05	4.04E-04	0.5	4.51E+05	2.23E-04	5.5	3.90E+05	2.13E-03
5975-1	<0.2	4.41E+05	>1.00E-05	0.8	5.33E+05	4.02E-04	0.8	4.20E+05	3.41E-04	6.6	3.39E+05	2.24E-03
5976-1	0.6	9.01E+05	5.25E-04	0.3	7.95E+05	2.32E-04	0.8	4.06E+05	3.30E-04	7.6	3.15E+05	2.40E-03
5978-1	< 0.3	7.77E+04	>1.00E-05	1.0	1.30E+05	1.31E-04	0.8	3.90E+05	2.98E-04	8.5	2.76E+05	2.33E-03
6057-1	0.8	6.48E+05	4.97E-04	0.7	1.00E+06	7.04E-04	0.8	4.30E+05	3.65E-04	9.2	2.54E+05	2.33E-03
6058-1	0.6	9.27E+05	5.29E-04	1.0	6.01E+05	4.93E-04	0.8	3.81E+05	3.17E-04	6.8	3.42E+05	2.32E-03
5982-1	1.2	1.68E+06	1.98E-03	1.5	1.28E+06	1.89E-03	<0.1	2.17E+05	>1.00E-05	4.7	2.27E+05	1.06E-03
5983-1	1.7	1.12E+06	1.95E-03	1.5	1.43E+06	2.11E-03	1.0	2.63E+05	2.72E-04	12.3	3.42E+05	4.21E-03
5987-1	0.7	6.00E+05	4.23E-04	0.7	7.52E+05	5.26E-04	1.0	4.24E+05	4.39E-04	18.3	2.10E+05	3.83E-03
5988-1	0.3	1.66E+06	4.76E-04	0.2	2.90E+06	6.54E-04	2.5	1.44E+05	3.61E-04	29.4	7.72E+04	2.27E-03

5991-1	0.4	1.08E+06	4.67E-04	0.2	1.86E+06	4.23E-04	1.3	1.81E+05	2.42E-04	31.9	1.52E+05	4.83E-03
5992-1	0.3	1.34E+06	4.21E-04	0.3	1.98E+06	5.32E-04	1.9	1.44E+05	2.88E-04	25.5	1.05E+05	2.67E-03
5995-1	0.4	1.06E+06	3.80E-04	0.2	1.80E+06	3.77E-04	0.9	2.81E+05	2.47E-04	26.2	1.12E+05	2.93E-03
5998-1	0.3	7.12E+05	2.28E-04	0.2	1.07E+06	2.04E-04	0.6	6.20E+05	3.81E-04	15.6	6.72E+04	1.05E-03
5999-1	0.2	5.80E+05	1.11E-04	0.6	3.95E+05	2.44E-04	1.0	3.38E+05	3.22E-04	30.9	1.26E+05	3.89E-03
6000-1	0.3	6.59E+05	1.83E-04	0.5	4.10E+05	1.99E-04	1.1	3.64E+05	3.94E-04	40.0	5.37E+04	2.15E-03
6003-1	0.7	1.14E+06	8.03E-04	0.5	1.58E+06	7.73E-04	1.2	2.45E+05	2.95E-04	62.6	8.58E+04	5.37E-03
6007-1	3.3	1.41E+05	4.63E-04	2.7	2.44E+05	6.62E-04	0.8	4.92E+05	4.09E-04	40.2	1.30E+05	5.23E-03
5946-1	1.0	5.89E+05	6.01E-04	0.9	5.91E+05	5.20E-04	0.9	4.06E+05	3.61E-04	3.1	2.64E+05	8.25E-04
5947-1	1.1	5.14E+05	5.72E-04	1.1	5.77E+05	6.17E-04	1.2	2.77E+05	3.46E-04	27.3	8.60E+04	2.35E-03

Table 12. Anti-CD40xMSLN IgG-Fab molecule functional activity on human B cells in the presence of CHO cells expressing human mesothelin

Antibody ID	EC50 (nM)
13468-1	0.007
5966-1	0.008
5978-1	0.033
13473-1	0.007
13474-1	0.009
13469-1	0.011
5967-1	0.005
6057-1	0.018
5991-1	0.004
6003-1	0.007
13470-1	0.008
5968-1	0.004
6058-1	0.010
5992-1	0.007
13475-1	0.015
13471-1	0.016
5970-1	0.005
5982-1	0.005
6007-1	0.009
5960-1	0.006
5972-1	0.007
5983-1	0.004
5995-1	0.006
5946-1	0.007
5961-1	0.004
5973-1	0.006
13472-1	>0.523
5998-1	0.007
5947-1	0.003
5963-1	0.022
5975-1	0.022
5987-1	0.003
5999-1	0.005
5965-1	0.020
5976-1	0.010
5988-1	0.004
6000-1	0.002

[1377] *Generation and evaluation of mesothelin x CD40 bispecific agonist antibodies in the IgG-scFv format.*

[1378] A panel of mesothelin x CD40 bispecific antibodies were generated in the IgG-scFv format (Table 13) and evaluated for binding to soluble forms of human and cynomolgus monkey CD40 and human and cynomolgus monkey mesothelin using the Octet assay (Table 14). Association rate (Kon), disassociation rate (Kdis), and equilibrium binding constant (KD) were calculated. The majority of bispecific antibodies showed high affinity binding to both human and cynomolgus monkey CD40 and mesothelin. These bispecific antibodies were next evaluated for their ability to induce mesothelin-dependent activation of human B cells. CHO cells expressing human mesothelin were seeded into a 96 well plate. The next day varying concentrations of the mesothelin x CD40 IgG-scFv antibodies were added to the wells along with purified human B cells and the plates were incubated for an additional 48 hours. Upregulation of CD86, a marker of CD40-mediated B cell activation, was quantified on B cells by flow cytometry. EC50 values for CD86 upregulation were calculated and demonstrate that the majority of mesothelin x CD40 IgG-scFv bispecific antibodies were able to B cell activation in the presence of CHO cells expressing human mesothelin (Table 15).

Table 13. Anti-CD40xMSLN IgG-scFv molecules

Antibody ID	IgG (MSLN)	scFv (CD40)	IgG (CD40)	scFv (MSLN)
6028-2	4H6	36F3		
6030-2	4H6	39C2		
6032-2	6F4	29H10		
6043-2			30A12	4H6
6044-2			30A12	6F4
6033-2	6F4	30A12		
6034-2	6F4	33H6		
6035-2	6F4	33H9		
6056-2	6F4	39C2		
6039-2			4G7	4H6
6040-2			4G7	6F4
6041-2			29H10	4H6
6042-2			29H10	6F4
6045-2			33H6	4H6
6046-2			33H6	6F4
6047-2			33H9	4H6
6048-2			33H9	6F4
6049-2			35F11	4H6
6050-2			35F11	6F4
6051-2			36F3	4H6
6052-2			36F3	6F4
6055-2			37A6	4H6

6054-2			37A6	6F4
6053-2			39C2	4H6
6023-2			39C2	6F4

Table 14. Anti-CD40xMSLN IgG-scFv molecule binding to human and cynomolgus monkey CD40 and mesothelin

Antibody ID	huCD40(1-193)				cynoCD40(1-193)				huMSLN(296-598)				cynoMSLN(296-598)			
	K_D (nM)	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})		K_D (nM)	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})		K_D (nM)	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})		K_D (nM)	k_a ($M^{-1}s^{-1}$)	k_d (s^{-1})	
6028-2	10	7.69E+04	7.67E-04		7.1	9.64E+04	6.87E-04		0.2	7.33E+05	1.43E-04		1.8	3.71E+05	6.80E-04	
6030-2	7.9	2.14E+05	1.68E-03		6.0	3.01E+05	1.81E-03		0.2	7.08E+05	1.75E-04		1.9	3.53E+05	6.87E-04	
6032-2	2.0	2.04E+05	4.07E-04		1.7	2.38E+05	4.07E-04		0.4	6.92E+05	3.06E-04		5.8	3.45E+05	1.99E-03	
6043-2	1.2	3.21E+05	4.07E-04		0.9	4.22E+05	4.07E-04		0.1	4.44E+05	3.87E-05		4.3	2.76E+05	1.20E-03	
6044-2	1.3	3.24E+05	4.24E-04		0.8	4.85E+05	3.98E-04		1.5	3.30E+05	5.06E-04		17	3.43E+05	5.87E-03	
6033-2	4.2	3.11E+05	1.32E-03		2.9	4.88E+05	1.42E-03		0.4	4.49E+05	1.97E-04		5.9	3.31E+05	1.94E-03	
6034-2	2.1	1.66E+05	3.50E-04		2.1	1.82E+05	3.84E-04		0.3	6.00E+05	1.96E-04		5.4	2.96E+05	1.59E-03	
6035-2	<0.1	1.03E+05	<1.00E-05		<0.1	9.43E+04	<1.00E-05		0.3	6.75E+05	2.29E-04		6.3	2.71E+05	1.71E-03	
6056-2	4.4	3.63E+05	1.59E-03		5.8	2.98E+05	1.73E-03		0.3	7.19E+05	2.18E-04		5.1	3.43E+05	1.76E-03	

6039-2	1.4	1.12E+06	1.61E-03	2.1	1.04E+06	2.21E-03	0.3	5.13E+05	1.77E-04	5.4	2.81E+05	1.50E-03
6040-2	1.3	1.39E+06	1.86E-03	1.9	9.24E+05	1.73E-03	1.2	4.24E+05	5.16E-04	32	2.35E+05	7.63E-03
6041-2	0.5	7.09E+05	3.57E-04	0.6	5.39E+05	3.47E-04	0.4	5.04E+05	1.79E-04	5.7	1.58E+05	9.05E-04
6042-2	0.7	6.32E+05	4.29E-04	0.8	3.94E+05	3.03E-04	0.9	5.29E+05	4.60E-04	20	2.55E+05	5.04E-03
6045-2	0.4	8.81E+05	3.87E-04	0.6	5.80E+05	3.27E-04	0.3	5.23E+05	1.45E-04	4.6	1.87E+05	8.65E-04
6046-2	0.5	8.93E+05	4.05E-04	0.5	5.94E+05	3.14E-04	0.9	5.27E+05	4.76E-04	18	2.50E+05	4.59E-03
6047-2	0.2	5.77E+05	1.32E-04	<0.3	3.46E+04	<1.00E-05	0.4	3.75E+05	1.38E-04	6.0	1.68E+05	1.01E-03
6048-2	0.5	5.45E+05	2.59E-04	0.3	4.26E+05	1.33E-04	1.0	5.29E+05	5.53E-04	22	2.39E+05	5.35E-03
6049-2	0.9	8.34E+05	7.21E-04	1.2	5.06E+05	5.82E-04	0.3	5.36E+05	1.71E-04	5.1	1.85E+05	9.53E-04
6050-2	0.8	7.97E+05	6.49E-04	1.2	5.12E+05	6.09E-04	0.8	5.41E+05	4.51E-04	15	2.77E+05	4.27E-03
6051-2	3.5	1.24E+05	4.31E-04	4.7	1.00E+05	4.66E-04	0.2	5.99E+05	1.35E-04	5.5	1.91E+05	1.05E-03

6052-2	2.5	2.18E+05	5.56E-04	3.3	1.05E+05	3.50E-04	0.7	7.13E+05	5.33E-04	19	2.69E+05	5.19E-03
6055-2	0.5	1.46E+06	7.87E-04	0.8	7.45E+05	5.70E-04	0.4	5.52E+05	2.11E-04	6.1	1.79E+05	1.09E-03
6054-2	0.8	4.38E+05	3.33E-04	1.4	2.77E+05	4.00E-04	1.3	3.67E+05	4.62E-04	17	3.36E+05	5.78E-03
6053-2	0.9	4.53E+05	4.27E-04	0.5	2.63E+05	1.25E-04	0.4	3.41E+05	1.51E-04	4.3	2.52E+05	1.09E-03
6023-2	1.0	4.31E+05	4.40E-04	0.9	4.47E+05	4.13E-04	1.1	4.32E+05	4.60E-04	16	3.26E+05	5.35E-03

Table 15. Anti-CD40xMSLN IgG-scFv molecule functional activity on human B cells in the presence of CHO cells expressing human mesothelin

Antibody ID	EC50 (nM)
6028-2	0.012240302
6030-2	0.010141789
6032-2	0.009349861
6043-2	0.007758607
6044-2	0.005001671
6033-2	0.006030707
6034-2	0.007262489
6035-2	0.016708182
6056-2	0.004577206
6039-2	0.003317003
6040-2	0.002665906
6041-2	0.004479453
6042-2	0.003988474
6045-2	0.004663271
6046-2	0.004723116
6047-2	0.005131571
6048-2	0.004342491
6049-2	0.005853663
6050-2	0.003388309
6051-2	0.003753076
6052-2	0.003155443
6055-2	0.004340411
6054-2	0.003331971
6053-2	0.004033184
6023-2	0.002805537

[1379] *Sequence optimization of mesothelin x CD40 bispecific agonist antibodies in the IgG-scFv format*

[1380] Engineering of select mesothelin x CD40 bispecific IgG-scFv antibodies was performed to remove potential sequence liabilities in the antibody heavy and light chain variable regions by standard recombinant DNA techniques. These antibodies were produced by over-expression in cell lines and purified using techniques that were known in the art. These bispecific antibodies were next evaluated for their ability to induce mesothelin-dependent activation of human B cells. CHO cells expressing human mesothelin were seeded into a 96 well plate. The next day varying concentrations of the mesothelin x CD40 IgG-scFv sequence variant antibodies were added to the wells along with purified human B cells and the plates were incubated for an additional 48 hours. Upregulation of

CD86, a marker of CD40-mediated B cell activation, was quantified on B cells by flow cytometry. EC50 values for CD86 upregulation were calculated and demonstrate that the majority of mesothelin x CD40 IgG-scFv sequence variant bispecific antibodies maintain the ability to induce B cell activation in the presence of CHO cells expressing human mesothelin (Table 16).

Table 16. Sequence optimized anti-CD40xMSLN IgG-scFv molecule functional activity on human B cells in the presence of CHO cells expressing human mesothelin

Antibody ID	EC50 (nM)	Antibody ID	EC50 (nM)	Antibody ID	EC50 (nM)
8861-1	0.00332	8786-1	0.00322	8826-1	0.04585
8866-1	0.00306	8787-1	0.00274	8827-1	0.06588
8862-1	0.00295	8788-1	0.00371	8828-1	>0.635
8863-1	0.00557	8789-1	0.00321	8829-1	>0.635
8869-1	0.00181	8790-1	0.00480	8830-1	0.00559
8871-1	0.00259	8791-1	0.00329	8831-1	0.00523
8873-1	0.00311	8766-1	0.00355	8832-1	0.00572
8875-1	0.00241	8792-1	0.00439	8833-1	0.00788
8867-1	0.00216	8793-1	0.00405	8834-1	0.03859
8868-1	0.00363	8794-1	0.00378	8835-1	0.06977
8864-1	0.00283	8795-1	0.00511	8836-1	>0.635
8870-1	0.00267	8796-1	0.00437	8837-1	>0.635
8872-1	0.00267	8797-1	0.00384	8838-1	>0.635
8874-1	0.00262	8798-1	0.00342	8839-1	>0.635
8876-1	0.00171	8804-1	0.00323	8840-1	0.00641
8865-1	0.00329	8805-1	0.00406	8841-1	0.00572
6041-3	0.00395	8806-1	0.00326	8842-1	0.00430
8764-1	0.00434	8807-1	0.00342	8843-1	0.00376
8768-1	0.00334	8808-1	0.00331	8844-1	0.00263
8769-1	0.00434	8809-1	0.00386	8845-1	0.00362
8770-1	0.02176	8810-1	0.00451	8846-1	0.00542
8771-1	0.05341	8811-1	0.00436	8847-1	>0.635
8772-1	0.17205	8812-1	0.00415	8848-1	0.28130
8773-1	0.42410	8813-1	0.00356	8849-1	0.00315
8774-1	0.00409	8814-1	0.00453	8850-1	0.00353
8775-1	0.00309	8815-1	0.00368	8851-1	0.00648
8776-1	0.02073	8816-1	0.00438	8852-1	0.12325
8777-1	0.01153	8817-1	0.00342	8853-1	0.05720
8778-1	0.00362	8818-1	0.00518	6052-3	0.00445
8765-1	0.00305	8819-1	0.00387	8854-1	0.00303
8779-1	0.01746	8820-1	0.00366	8855-1	0.00354
8780-1	0.01673	8821-1	0.00433	8856-1	0.00363
8781-1	0.17750	8822-1	0.00404	8857-1	0.00346
8782-1	0.42500	8823-1	0.00351	8858-1	0.00357
8783-1	0.00301	8824-1	0.00331	8859-1	0.00300

8784-1	0.00311	8825-1	0.00255	8860-1	0.00357
8785-1	0.00293	8767-1	0.00209		

[1381] *Effect of varying levels of mesothelin on activity of select mesothelin x CD40 bispecific antibodies in a human B cell functional assay*

[1382] MC38 cells were engineered to express varying levels of human mesothelin ranging from low (MC38.MSLN1) to high (MC38.MSLN5). Given that MC38 cells are derived from mouse, they do not express human mesothelin (Figure 4). Parental MC38 cells and MC38 cells expressing varying levels of human mesothelin were seeded into 96 well plates and cultured overnight. The next day, mesothelin x CD40 bispecific antibodies were added to the cells at varying concentrations. Isolated human B cells were then added at to the wells and the plates were incubated for an additional 48 hours. Upregulation of CD86 on B cells was evaluated by flow cytometry and used as a measure of B cell activation. EC50 and maximum activity (Emax) values were calculated and demonstrate that the mesothelin x CD40 bispecific antibodies were able to induce B cell activation across all levels of MSLN expression (Table 17). Cell lines expressing lower levels of mesothelin induced lower maximum activity of the mesothelin x CD40 bispecific antibodies, as measured by the geometric mean fluorescence intensity of CD86 upregulation on B cells. Mesothelin x CD40 bispecific antibodies had no activity in the presence of parental MC38 cells, consistent with the mesothelin-dependent activity of these molecules for CD40 agonist activity.

Table 17. Activity of anti-CD40xMSLN bispecific molecules in human B cell and monocyte-derived dendritic cell assays in the presence of varying levels of mesothelin

		Antibody ID					
		6041	8765	8766	8767	8945	8947
Human B Cell Functional Assay (CD86 upregulation)	MC38.MSLN1 EC50 (pM)	0.2326	0.2287	0.2735	0.1472	0.2752	0.2687
	MC38.MSLN2 EC50 (pM)	0.5723	0.6028	0.6877	0.4584	0.6637	0.6626
	MC38.MSLN3 EC50 (pM)	0.801	0.6696	0.8764	0.5357	1.0162	0.6689
	MC38.MSLN4 EC50 (pM)	0.9542	0.8518	1.1365	0.7048	1.4765	0.9471
	MC38.MSLN5 EC50 (pM)	1.6005	0.8549	2.0058	1.2579	2.0122	1.0863
	MC38 parental EC50 (pM)	∞	∞	∞	∞	∞	∞
	MC38.MSLN1 Emax (MFI)	803.9	903	849	742.1	660.2	809.5
	MC38.MSLN2 Emax (MFI)	1169	1318	1223	1173	978.6	1259
	MC38.MSLN3 Emax (MFI)	1895	1861	1869	1849	1479	1733
	MC38.MSLN4 Emax (MFI)	1325	1400	1399	1341	1283	1406
Human Monocyte-Derived Dendritic Cell Functional Assay	MC38.MSLN5 Emax (MFI)	1396	1406	1474	1349	1186	1389
	MC38 parental Emax (MFI)	Bkgd	Bkgd	Bkgd	Bkgd	Bkgd	Bkgd
	MC38.MSLN2 EC50 (nM)	0.179	0.089	0.183	0.101	0.168	0.184
	MC38.MSLN5 EC50 (nM)	0.241	0.231	0.26	0.248	0.414	0.399

(IL12p40 production)	MC38 parental EC50 (nM)	∞	∞	∞	∞	∞	∞
	MC38.MSLN2 Emax (IL12p40 pg/ml)	1142	952	1205	1116	338.1	1010
	MC38.MSLN5 EC50 (nM)	2400	2148	2343	2349	1004	1869
	MC38 parental Emax (IL12p40 pg/ml)	Bkgd	Bkgd	Bkgd	Bkgd	Bkgd	Bkgd

* Poor curve fit
Bkgd= Background level of
assay

[1383] *Effect of varying levels of mesothelin on activity of select mesothelin x CD40 bispecific antibodies in a monocyte-derived dendritic cell functional assay.*

[1384] MC38 cells engineered to express varying levels of human mesothelin (Figure 4) or parental MC38 cells were seeded into 96 well plates and cultured overnight. The next day, mesothelin x CD40 bispecific antibodies were added to the cells at varying concentrations. Monocyte-derived dendritic cells differentiated from human monocytes using GM-CSF and IL4 were added to the wells and the plates were incubated for an additional 48 hours. Supernatants from each well were collected and concentrations of IL12p40 secreted from dendritic cells quantified using an ELISA kit. EC50 and maximum activity (Emax) values were calculated and demonstrate that the mesothelin x CD40 bispecific antibodies were able to induce dendritic cell activation across both high and low levels of MSLN expression (Table 17). Cell lines expressing lower levels of mesothelin induced lower maximum activity of the mesothelin x CD40 bispecific antibodies. Mesothelin x CD40 bispecific antibodies had no activity in the presence of parental MC38 cells, consistent with the mesothelin-dependent activity of these molecules for CD40 agonist activity.

[1385] *Activity of select mesothelin x CD40 bispecific antibodies in cynomolgus monkey B cell functional assay with cynomolgus monkey mesothelin-expressing CHO cells.*

[1386] CHO cells engineered to express cynomolgus monkey mesothelin were seeded into 96 well plates and cultured overnight. The next day, select mesothelin x CD40 bispecific antibodies were added to the cells at varying concentrations. Isolated cynomolgus monkey B cells were then added to the wells and the plates were incubated for an additional 48 hours. Upregulation of CD23 on the cynomolgus monkey B cells was evaluated by flow cytometry and used as a measure of CD40-induced B cell activation. EC50 were calculated and demonstrate that the mesothelin x CD40 bispecific antibodies are able to induce activation of cynomolgus monkey B cells in the presence of cynomolgus monkey mesothelin (Table 18).

Table 18. Activity of anti-CD40xMSLN bispecific molecules on cynomolgus monkey B cells stimulated with CHO cells expressing cynomolgus monkey mesothelin

Antibody ID	EC50 (nM)
6041	0.0012
8765	0.0013
8766	0.0016
8767	0.0014
8945	0.0018
8947	0.0014

[1387] *Pharmacokinetics and stability of mesothelin x CD40 bispecific antibodies in mice.*

[1388] In order to characterize the stability of mesothelin x CD40 bispecific antibodies in vivo, select mesothelin x CD40 bispecific molecules were injected into mice a 1mg/kg dose. Animals were bled at varying timepoints after injection and serum isolated. Immunoassays were used to quantify the

concentration of mesothelin x CD40 bispecific antibodies in serum, either capturing with human CD40 and detecting with anti-human IgG or capturing with human mesothelin and detecting with anti-human IgG. The combination of these two assays provide the ability to detect molecule degradation or instability. The CD40 or mesothelin capture assays yielded equivalent results, suggesting the molecules remained intact over the course of 14 days in vivo (Figure 5). All molecules with the exception of one had similar pharmacokinetic properties.

[1389] Immunostimulatory and anti-tumor activity of a tumor-targeted CD40 agonist bispecific antibody in mouse models

[1390] *Generation of a mouse surrogate human EPCAM-dependent CD40 agonist bispecific antibody and human EPCAM-expressing mouse tumor cell lines.*

[1391] The anti-mouse CD40 agonist antibody (clone FGK45) and anti-human EPCAM antibody (clone 4-7) was used to generate a surrogate CD40 bispecific antibody in the IgG-scFv format with a mouse IgG1 N297G Fc domain (anti-muCD40 x huEPCAM bispecific antibody). The bispecific antibody contains anti-CD40 at the N-terminus of the molecule and the anti-EPCAM as scFv at the C-terminus of the molecule. Human EPCAM was over-expressed in the mouse MC38 colon carcinoma cell line and the mouse B16F10 melanoma cell line by transduction with a retrovirus encoding both human EPCAM and the human truncated nerve growth factor receptor (NGFR) reporter gene (Figure 6). Human EpCAM-MC38 or human EpCAM-B16 tumor cells were inoculated in the right flank at 3e5 cells per implant and allowed to grow for 24 days or 11 days, respectively. Tumors were measured twice a week with digital calipers, demonstrating that the tumor cell lines expressing human EPCAM maintained the ability to grow and form tumors in mice. Flow cytometry on tumors harvested from mice and subjected to enzymatic disassociation demonstrated that human EPCAM expression was maintained on the surface of tumor cells during in vivo tumor formation (Figure 6).

[1392] *Human EPCAM-dependent agonist activity of the EPCAM x CD40 agonist antibody on mouse B cells.*

[1393] Human EpCAM overexpressing MC38 cells (huEpCAM-MC38) or parental MC38 cells were seeded into 96 plates and cultured overnight. The next day, the anti-muCD40 x huEPCAM bispecific antibody, the parental anti-CD40 antibody (FGK54), or an isotype control antibody was added to the cells at varying concentrations. Isolated mouse B cells were then added at to the wells and the plates were incubated for an additional 48 hours. Upregulation of CD86 on B cells was evaluated by flow cytometry and used as a measure of B cell activation. EC50 values were calculated and demonstrate that the anti-muCD40 x huEPCAM bispecific antibody was able to induce mouse B cell activation (Figure 7). The anti-muCD40 x huEPCAM bispecific antibody demonstrated robust agonist activity that was dependent on the presence of human EPCAM and showed dramatically higher potency in activating B cells compared to the FGK45 anti-CD40 monoclonal antibody.

[1394] *In vivo immunostimulatory activity of the anti-muCD40 x huEPCAM bispecific antibody in the MC38 tumor model*

[1395] To evaluate the effect of a tumor-targeted CD40 agonist bispecific antibody on immune cell activation in vivo, human EpCAM-expressing MC38 tumor cells were inoculated in the right flank of mice at 3×10^5 cells per implant and allowed to grow for 8-10 days. The mice were then randomized by tumor volume (60-100mm³) and treated with indicated antibodies administered intraperitoneally at 5 mg/kg. Tumors, tumor draining lymph nodes (dLNs), and non-draining lymph nodes (ndLNs) were harvested at 24-48 hours post-treatment and single cell suspension prepared by enzymatic digestion of tissues. Flow cytometry analysis was performed to determine cell proportion and phenotypes, staining for surface immune cell lineage markers and activation markers, and intracellular cytokines (Figure 8). As expected, CD40 agonist antibody (anti-CD40) treatment activated dendritic cells (DCs) located in tumor and in peripheral tissues including dLN and ndLN. In contrast, the anti-muCD40 x huEPCAM bispecific antibody only activated tumor infiltrating DCs in the human EPCAM-expressing MC38 tumors, but not DCs in peripheral tissues (dLN and ndLN) (Figure 8A). In addition, the anti-muCD40 x huEPCAM bispecific antibody increased the CD8 T cell to regulatory T cell ratio and interferon gamma (IFN γ)-producing CD4⁺ T cells in the tumor (Figure 8B). Total T cells were also isolated from spleens of huEpCAM-expressing MC38 tumor-bearing mice that had been treated with the indicated antibodies. Bone marrow-derived dendritic cells (BMDCs) were generated from naïve C57BL/6 bone marrow cells by incubating with recombinant GM-CSF and IL4 for 7 days. BMDCs were pulsed with the indicated tumor cells overnight and then co-cultured with isolated T cells for 18 hours. An ELISPOT assay was used to measure the production of IFN γ by antigen specific T cells, demonstrating that the anti-muCD40 x huEPCAM bispecific antibody significantly increases T cells specific for both human EPCAM-expressing MC38 and parental MC38 antigens (Figure 9). Taken together, these data demonstrate that a tumor-targeted CD40 agonist bispecific antibody can activate myeloid population in the tumor, such as DCs, in a tumor-associated antigen dependent manner, leading to enhanced anti-tumor T cell responses.

[1396] *In vivo immunostimulatory activity of the anti-muCD40 x huEPCAM bispecific antibody in the B16F10 tumor model*

[1397] To evaluate the effect of a tumor-targeted CD40 agonist bispecific antibody on immune cell activation in an additional tumor model, vector control or human EpCAM-expressing B16F10 tumor cells were inoculated in the right flank of mice at 3×10^5 cells per implant and allowed to grow for 8-10 days. The mice were then randomized by tumor volume (60-100mm³) and treated with indicated antibodies administered intraperitoneally at 5 mg/kg. Tumors, tumor dLNs, and ndLNs were harvested at 24-48 hours post-treatment and single cell suspension prepared by enzymatic digestion of tissues. Flow cytometry analysis was performed to determine cell proportion and phenotypes, staining for surface immune cell lineage markers and activation markers, and intracellular cytokines (Figure 10). As expected, CD40 agonist antibody (anti-CD40) treatment DCs located in tumor and in peripheral tissues including dLN and ndLN in parental huEpCAM negative B16F10 tumors and in

huEpCAM-expressing B16F10 tumors. In contrast, the anti-muCD40 x huEPCAM bispecific antibody only activated tumor infiltrating DCs in huEpCAM-expressing B16F10 tumors, but not in parental huEpCAM- B16F10 tumor. The anti-muCD40 x huEPCAM bispecific antibody did not activate DCs in dLN or ndLN, regardless of whether these mice had human EPCAM-expressing or non-expressing B16F10 tumors (Figure 10A). In addition, the anti-muCD40 x huEPCAM bispecific antibody increased IFN γ -producing effector CD4 $^{+}$ and CD8 $^{+}$ T cells in human EPCAM-expressing B16F10 tumors, but not in parental B16F10 tumors (Figure 10B). These data provide additional evidence that a tumor-targeted CD40 agonist bispecific antibody can activate tumor infiltrating immune cells in a tumor-associated antigen dependent manner to enhance anti-tumor T cell responses.

[1398] *The anti-muCD40 x huEPCAM bispecific antibody does not induce upregulation of systemic serum cytokines or liver damage in mice*

[1399] CD40 agonist antibodies have shown dose-limiting toxicities in clinical trials, including liver damage and elevated serum cytokine levels, presumably due to systemic immune cell activation. To compare systemic immune cell activation and liver damage between a non-targeted CD40 agonist antibody and a tumor-targeted CD40 agonist bispecific antibody, mice bearing established human EPCAM-expressing MC38 tumors (described above) were treated with a non-specific isotype control antibody, a murine CD40 agonist antibody (anti-CD40), a CD40 antibody bearing a mutation in the Fc domain that decreases Fc receptor binding rendering it a poor agonist (anti-CD40 N297G), or the anti-muCD40 x huEPCAM bispecific antibody (anti-CD40xhuEPCAM) and serum was collected 24-48 hours later. Concentrations of select cytokines was measured using mouse cytokine/chemokine multiplex assay kit. The anti-CD40 antibody increased concentrations of several pro-inflammatory cytokines in the serum of mice, consistent with systemic immune cell activation. In contrast, neither the anti-CD40 N297G nor the anti-muCD40 x huEPCAM bispecific antibody induced increased serum cytokine concentrations (Figure 11A). Similar results were observed in the B16F10 tumor model, where, in contrast to the non-targeted anti-CD40 antibody, the anti-muCD40 x huEPCAM bispecific antibody did not induce an increase in serum cytokine concentrations in either mice bearing vector control B16F10 tumors or mice bearing human EPCAM-expressing B16F10 tumors (Figure 11B). The livers of mice bearing human EPCAM-expressing MC38 tumors and treated with anti-CD40 agonist antibody or the anti-muCD40 x huEPCAM bispecific antibody were also were fixed in 10% Neutral buffered formalin, embedded in paraffin, sectioned, and stained with Hematoxylin and Eosin for histological analysis. While the anti-CD40 agonist antibody induced multifocal mononuclear cell infiltration in the liver and these livers contained single cell necrosis figures, livers from the anti-muCD40 x huEPCAM bispecific antibody-treated mice did not show any distinguishable histopathological changes (Figure 11C). Taken together, these data indicate that a tumor-targeted CD40 agonist antibody is capable of inducing local immune cell activation in the tumor, without leading to systemic cytokine production or liver damage that has been associated with anti-CD40-mediated toxicity.

[1400] *Anti-tumor effects of human EPCAM x murine CD40 bispecific antibody in mouse tumor model*

[1401] To evaluate the effect of a tumor-targeted CD40 agonist antibody on immune-mediated inhibition of tumor growth and the potential for this therapeutic approach to combine with blockade of the immune checkpoint inhibitor PD1, mice bearing human EPCAM-expressing MC38 tumors were treated with an isotype control antibody, anti-CD40 agonist antibody, anti-PDL1 (that blocks interactions between PD1 and PDL1), anti-muCD40 x huEPCAM bispecific antibody, the combination of anti-CD40 with anti-PDL1, and the combination of anti-muCD40 x huEPCAM bispecific with anti-PDL1 (Figure 12). Single agent treatment with anti-CD40 agonist antibody or anti-PDL1 blocking antibody had minimal effect on tumor growth (Figure 12A) or on improving survival (Figure 12B) relative to isotype control antibody treatment, while the combination of anti-CD40 agonist and anti-PDL1 significantly inhibited tumor growth. Strikingly, treatment with the anti-muCD40 x huEPCAM bispecific agonist antibody dramatically inhibited tumor growth and increased animal survival compared to other monotherapy treatments. Moreover, the combination of the anti-muCD40 x huEPCAM bispecific antibody with anti-PDL1 led to complete regression of tumors, with all animals surviving until the end of the study. These complete responder animals were housed for 3 months without treatment and no evidence of tumor reappearance, after which they were subject to tumor re-challenge with either human EPCAM-expressing MC38 tumor cells or B16F10 tumor cells (Figure 13). These animals completely rejected the human EPCAM-expressing MC38 tumor cells, while control B16F10 tumor grew as expected. Both tumor lines grew as expected in age-matched control animals that had not previously been exposed to human EPCAM-expressing MC38 tumors or the anti-muCD40 x huEPCAM bispecific antibody. Taken together, these data demonstrate that tumor-targeted CD40 agonist antibody therapy can enhance anti-tumor immune responses, leading to regression of established tumors that is further enhanced by combination with PD1/PDL1 blockade. In addition, the combination of tumor-targeted CD40 agonist treatment with PD1/PDL1 blockade can induce long-lasting immune memory that can protect against subsequent tumor re-challenge

[1402] All publications, patents, and patent applications discussed and cited herein are hereby incorporated by reference in their entireties. It is understood that the disclosed invention is not limited to the particular methodology, protocols and materials described as these can vary. It is also understood that the terminology used herein is for the purposes of describing particular embodiments only and is not intended to limit the scope of the appended claims.

[1403] Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

Human CD40 >NP_001241.1 tumor necrosis factor receptor superfamily member 5 isoform 1 precursor [Homo sapiens]
(SEQ ID NO: 1)
MVRPLQCVLWGCLLTAVHPEPTACREKQYLINSQCSLCPGQKLVSDCTEFTETECPCGESEFLDTWNRETHCHQHKYCDPNLGLRVQKKG
TSETDTICTCEEHWHCTSEACESCVLHRSCSPGFVKQIATGVSDTICEPCPVGFFSNVSSAFEKCHPWTSCTKDLVVQQAGTNKTDVVCGPQDRL
RALVVIPIIFGILFAILLVLFVIKKVAKKPTNKAPHKPKQEPQENFPDDLPGSNTAAPVQETLHGCQPVVTQEDGKESRISVQERQ

Human mesothelin >NP_005814.2 mesothelin isoform 1 preproprotein [Homo sapiens]
(SEQ ID NO: 2)
MALPTARPLLGSCGTPALGSLFLFLFSLGWVQPSRTLGETGQEAAPLDGVLANPPNISSLSRQLLGFPCAESVGLSTERVRELAVALAQKNVKLST
EQLRCLAHRLSEPPEDLDALPLDLLLFLNPDAFSGPQACTRFFSRITKANVDLLPRGAPERQRLPAALACWGVRSLLSEADVRLGGLACDLPGR
FVAESAELVLLPRLVSCPGPLDQDQQAARAALQGGGPPYGPSTWSVSTMDALRGLLPVLGQPIRSIPQGIWAARQSSRDPSWRQPRTLPRF
RREVEKTACPSGKKAREIDESLIFYKKWELEACVDAALLATQMDRVNAIPFTYEQLDVLKHKLDLYPQGYPESVIQHLGYLFLKMSPEDIRKWNV
TSLETLKALLEVNKGHEMSPQVATLIDRFVKGGRGQDKDLDLTLTAFYPGYLCSLSPEELSSVPSSIWAVRPQDLDTCDPRQLDVLYPKARLAFQN
MNGSEYFVKIQSFLGGAPTEDLKALSQQNVSMDLATFMKLRDTDAVLPLTVAEVQKLLGPHVEGLKAEERHRPVRDWILRQRQDDDLDTLGLGLQG
GIPNGYLVLDLSMQEALSGTPCLLGPGPVLTVLALLASTLA

Table 19: CD40 Antibody VLs and VHs

i	P	S	#	Desc	A	ri	p	e	T	y	p	e	LC	HC

2		i		P		1		-		2		S		:		5		0		5		5		6		3		3		H		9	
						(R50)				N																							
						1)		L_		A																							
						v2																											
						VL_																											
						PE_1																											
						3641																											
						057																											

2	i	1	P	1	(R14	GACATCCAGATGACCCAGTCTCCATCCTCACTGTCT	CAGTGTCAACTGGTGCAGTCTGGGGCTGAGGTGAC
-	S	2	6	G_	N	GCATCTGTAGGAGACAGAGTACCATCACTGTCTG	GAAGCTGGGGCTCAGTGAAGGTCTCTCTGCAAGG
3	:	3	vL	VH_	A	GGCAGTCAAGGACATTAGCAATAATTTAGCCTGGT	CTTCTGGATACACCTTCGCCGGCTACTATATGCACT
5	0	5	0	5		TTCAGCAGAAACCAAGGAAACCCCTAAGTCCCTG	GGGTGCGACAGGCCCTTGACAAAGGCTTGAGTGG
5	5	5	5	5		ATGTATGCTGCATCCAGTTTGACAGTGGAGTCCC	ATGGATGGATCAACCCCTCACAGTGGTGGCACAAA
5	5	5	5	5		ATCAACCTTCAGCGGCAGTGGATCTGGGACAGATT	CTATGCACAGAAGTTTCAGGACAGGGTCACCATGA
5	5	5	5	5		TCACCTTTCAACCATCAGCAGCCTGCAGCCTGAAGAT	CCAGGACACAGCTCCATCAACACAGCCTACATGGAA
8	3	3	3	3		TTTGGCAACTTATTACTGCCAACAGTATAATAGTTAC	CTGAGCAGGCTGAGATCTGACGACACGGCCGTGTA
9	H	9	9	9		CCTCTCACTTTTCGGCGGAGGACCAAGGTGGAGAT	TTACTGTGCGAGAGAACTGTTTCTATGTTCTGGG
9	H	9	9	9		CAGACGA	GAGTCGGGCACAACTGGTTGCGCCCTGGGGCCAG
9	H	9	9	9			GGAACCTGGTCAACCGTCTCCTCA
9	H	9	9	9		(SEQ ID NO: 11)	(SEQ ID NO: 12)
1	1	1	1	1		DIQMTQSPSSLSASVGDRVTITCRASQDINNNLAWFQ	QVQLVQSGAEVTKPGASVKVSCKASGYTFAGYYMH
1	1	1	1	1		QKPGKPKSLMYAASLHSGVPSFTSGSGDTFTFTI	WVRQAPGQGLEWMGWNPSPHSGGTNYAQKFQDRVT
1	1	1	1	1		SSLQPEDFATYYCQQYNISYPLTFGGGKVEIRR	MTRDTSINTAYMELSLRSDDTAVYYCARERISMVR
1	1	1	1	1			GVGHNWFAPWGQGTLVTVSS
1	1	1	1	1		(SEQ ID NO: 13)	(SEQ ID NO: 14)
2	i	1	P	1	(R16	GACATCCAGATGACCCAGTCTCCATCCTCCCTGTCT	CAGGTGCAGCTGGTGCAGTCCGGGACTGAAATGAA
-	S	2	6	G_	N	GCATCTGTAGGAGACAGAGTACCATCACTTGCCG	GAAGCTGGGGCTCAAATGAAGGTCTCTCTGCCAGA
3	:	3	vL	VH_	A	GGCAAGTCAAGGCATTAGAAATGATTTAGGTGGT	CTTCTGGATACACCTTCATCGCCTACTATATACACT
5	0	5	0	5		ATCAGCAGAAACCAAGGAAAGCCCTAAGCCCT	GGGTGCGACAGGCCCTTGACAAAGGACTTGAGTGG
5	5	5	5	5		GATCTCTGCTGATTCAGTTTGCAAGTGGGTCCC	ATGGATGGTCAACCTTGACAGTGGTGGCACAAA
5	5	5	5	5		ATCAAGGTTCAAGCTCAGTGGATCTGGGACAGAAT	CTTTGCCCGAGGTTTCAGGACAGAGTCAACCATGA
5	5	5	5	5		TCACCTCACAATCAGCAGCCTGCAGCCTGAAGAT	CCAGGACACGTCCATCAACACAGCCTACATGGAA
5	5	5	5	5		TTTGGCAACTTATTACTGTCTACAGTATAATAGTTAC	CTGAGCAGCCTGAGATCTGACGACACGGCCGTGTA
5	5	5	5	5		CCGTGGACGTTCCGCCAAGGGACCAAGGTGGAAT	TTACTGTGCGAGAGAGAAAGTTTAACTACAACATG
5	5	5	5	5		CAAAACA	GTGCTTTTGATATCTGGGGCCAAAGGACAAATGGTC
5	5	5	5	5			ACCGTCTCTTCA
5	5	5	5	5		(SEQ ID NO: 15)	(SEQ ID NO: 16)
5	5	5	5	5		DIQMTQSPSSLSASVGDRVTITCRASQGIKNDLWYQ	QVQLVQSGTEMKPGASMKVSCQTSQYTFIAYYIH
5	5	5	5	5		QKPGKAPKRLISAFLSQSGVPSRFSVSGSGTEFTLTIS	VVRQAPGQGLEWMGWNPSPHSGGTNAFPRFQDRVTMT
5	5	5	5	5		SLQPEDFATYYCQYNISYPTWTFGGQTKVEIKR	RDTSITTAAYMELSSRLSDDTAVYYCAREKFNYYGA
5	5	5	5	5			FDIWWGQGTMTVTVSS
5	5	5	5	5		(SEQ ID NO: 17)	(SEQ ID NO: 18)
5	5	5	5	5			

i	P	S	:	5	5	5	8	6	0	(R26 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 591	TCCTATGAGCTGACTCAGCCACCCCTCAGTGTCCGTG	TCTATGAGCTGAGTGGAGTCTGGGGAGGCTTAGT
											TCCTAGGACAGACAGCAATCACTACCTGCTCTGG	CAAGCCTGGAAGGTCCCTGAGACTCTCCTGTGCAG
											AGAAAGTTGGAAATAAATAATTTGCTGGTATC	CCTCTGAATTCACCTTCAGTGACTACTACATGAGCT
											AGCAGAAAGCCAGGCCAGTCCCTGTTCTGGTCATC	GGATCCGCCAGGCTCCAGGGAAGGGCTGGAGTG
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	TATCAAGATTTCAGCGGCCCTCAGGATCCCTGA	GGTTTCATATATTAGTCGAAAGTGGTGATACCATAT
											GCATTTCTCTGGTCCAACTCTGGGATCACAGCCA	ACTACGACAGACTGTGTAAGGCCGATTACCCATC
											CTCTGACCATCAGCGGAGCCAGGCTATGGATGAG	TCCAGGGACACGCCAAGAACTCACTGTATCTGCA
											GCTGACTATTACTGTGAGCGTGGACAGCAGAAC	AATGAATGCCCTGCCAGCCGAAAGACACGGCCGTGT
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	TGTGGTATTCGGCGGAGGACCAAGCTGACCGTCC	ATTACTGTGCGAGAGACTTAGCAGCAGGTGCTACA
											TAGGT	GGGGCCTTGACTGTCTGGGGCCAGGGAACCCCTGGT
											(SEQ ID NO: 27)	CACCGTCTCCTCA
											(SEQ ID NO: 28)	
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	SYELTQPPSVSPGQTASITCSGERLGNKYICWYQQ	QVQLVESGGGLVQPGGSLRLSCAASEFTFSDYYMSW
											KPGQSPVLVIYQDFKRPSPGIPERFSGSNSGITATLTISG	IRQAPGKGLEWVSYISRSGLTIYADSVKGRFTISRDN
											A TQAMDEADYYCQAWDSRTVVFVGGGKLTVLG	AKNSLYLQMNGLRAEDTAVYYCARDLAAGATGGLD
											(SEQ ID NO: 29)	CWQQGTLVTSS
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	CAGTCTGCCCTGACTCAGCCTGCCTCCGTGTCTGGG	GAGGTGACAGTGTGCTGGAGTCTGGGGAGGCTTGGT
											TCCTCTGGACAGTCGATCACCATCTCCTGCACCTGGA	ACAGCCTGGGGGTCCCTGAGACTCTCCTGTGCAG
											ACCAGCAGTGATGTGGAAATATAACCTTGTCTC	CCTCTGGATTACCTTTAGTAGAAATGCCATGAGTT
											CTGGTACCAACAGCACCCAGGCAAGCCCAAAAC	GGGTCCGCCAGGCTCCAGGGAAGGGCTGGAGTG
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	TCATGATTTTGGAGTCAATCAGCGGCCCTCAGGG	GGTCTCAGTACTGTGTGTTAGTGGTATTAGCACAT
											GTTCCTAATCGCTTCTTGGCTCCAAAGTCTGGCACC	ACTACGCAGACTCCGTGGAAGGGCCGTTCAACATC
											ACGGCCTCCCTGACAATCTCTGGCTCCAGGCTGC	TCCAGAGACAAATCCAAAGAACACGCTGTATCTGCA
											GGACGAGGCTGATTTCTGCTGCTCATATACAA	AATGAACAGTCTGAGAGCCGAGGACACGGCCGTAT
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	CTAGTAGCACTTATGTGATATTCGGCGGAGGGACC	ATTACTGTGCGAGAGGTTATAGCAACAGCTGGTGG
											AAGCTGACCGTCTAGGT	TACTTTGACTACTGGGGCCAGGGAACCCCTGGTCAC
											(SEQ ID NO: 31)	CGTCTCCTCA
											(SEQ ID NO: 32)	
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	QSALTQPAASVSGSPQSITISCTGTSSDVGNYNLVSWS	EVQLLESGGGLVQPGGSLRLSCAASGFTFSRNAMSW
											YQHPGKAPKLMIFEVNRPSGVSNRFSKSGTAS	VRQAPGKGLEWVSATGGSGISTYYADSVKGRFTISRDR
											LTISGLQAADAEADYFCCSYTTSSTYVIFGGGKLTVL	NSKNTLYLQMNSLRAEDTAVYYCARGYSNSWWYFD
											(SEQ ID NO: 33)	YWGQGTILVTSS
i	P	S	:	5	5	5	8	6	0	(R16 8)G ₋ v1 ₋ vH ₋ PE ₋₁ 3623 491	(SEQ ID NO: 34)	
											(SEQ ID NO: 35)	
											(SEQ ID NO: 36)	
											(SEQ ID NO: 37)	

2 1 - 3 : 5 5 - 3 5 8 5 1 1	i P S : 5 5 - 3 5 8 5 6 2	(R23 9)G_ vL_ vH_ PE_1 3623 562	CAGTCTGCCCTGACTCAGCCTCGCTCAGTGTCCGG GTCTCTGGACAGTCAGTACCATCTCTGCACTGG AACCAGCAGTGATGTTGGTGTATATCTTTGTCTC CTGGTACCAACAACACCCAGGCAAGCCCCCAAC TCATGATTTATGATGTCAGTAAGCGCCCTCTGGG GTCCCTGATCGCTTCTCTGGCTCCAAAGTCTGTCAAC ACGGCTCCCTGACCATCTCTGGGCTCCAGGTGA GGATGAGACTGATTATTACTGCTGCTCATATGCAG GCAACTACACTTATGTCTTCGGAAC TGGACCAAG GTCACCGTCTCTAGGT	(SEQ ID NO: 35)	(SEQ ID NO: 36)	CAGGTGCAAGCTGGTGGAGTCTGGGGAGGCGTGGT CCAGCTGGAGGTCCCTGAGACTCTCCTGTGCAG CGTCTGGATTACCCCTCAGTAGTATGGCATGCACT GGTCCGCCAGGCTCCAGGCAAGGGCTGGAGTG GGTGGCAGTTATGATGATGATGAAAGTAATAAAT ACTATGACAGACTCCGTGAAGGCCGAGTCAACATC TCCAGAGACAATTCCAGAACAACACGCTGTATCTGCA AATGAATAGCCTGAGAGCCGAGGACACGGCTGTGT ATTACTGTACGAGAGATGGCCGGAAC TACGTCTAC TTTGACAACTGGGGCCAGGGAACCTGGTCAACCGT CTCCTCA
			QSALTQPRSVSGSPGQSVTISCTGTSSDVGGYIFVSWY QQHPGKAPKLMYDVSKRPSGVDPFRFSKSVNTASL TISGLQAEDETDYYCCSYAGNYTYVFGTGKVTVLG	(SEQ ID NO: 37)	(SEQ ID NO: 38)	QVQLVESGGGVVQPGRLRLSCAASGFTLSSYGMHW VRQAPGKGLEWVA VIWYDGSNKYYADSVKGRVTIS RDNSKNTLYLQMNSLRAEDTAVYYCTRDGRNYVYF DNWQGQGLTVTVSS
			CAGTCTGCCCTGACTCAGCCTGCCTCCGTGTCTGGG TCTCTGGACAGTCGATCACCATCTCTCTGCACTGGA ACCAGCAGTGATGTTGGGAATTATAACCTTGTCTC CTGGTACCAACAACACCCAGGCAAGCCCCCAAC TCATGATTTATGAGGTCAATAGCGGCCCTCAGGG GTTTCTAATCGCTTCTCTGGCTCCAAAGTCTGGCAAC ACGGCTCCCTGACAATCTCTGGGCTCCAGGCTGA GGACGAGGCTGAATATTACTGCTGCTCATATGCAG GTAGAGACACTTTCGTGGTGTTCGGCGGAGGGACC AAGCTGACCGTCTCTAGGT	(SEQ ID NO: 39)	(SEQ ID NO: 40)	CAGGTGCAAGCTGGTGGAGTCTGGGGAGGCTGAA GAAAGCTGGGGCTCAGTGAAGGTCTCCTGCAAGG CTTCTGGATACACCTTCCCCGGCTACTATATGCACT GGGTGCACAGGCCCTTGGACAGGGCTTGAAGTG ATGGATGGATCAACCTGACAGTGGTGGCACA GTATACACAGAAGTTTCAGGGCAGGGTCACTTGA CCAGGACGCTCCGTGTCAGCACAGCCTACATTGAC CTGAACAAGGCTGAGATCTGACGACACGGCCGTATA TTACTGTGCGAGAGAGAGGTGTAGGACTACCAACT GCTATTTGGACTACTGGGGCCAAGGAAGTCTGGTC ACCGTCTCCTCA
			QSALTQPRSVSGSPGQSVTISCTGTSSDVGNYNLVS WYQHPGKAPKLMYEVNRRPSGVSNRFSGSKSGNTA SLTISGLQAEDEAEYYCCSYAGRDTFVVFVGGGTLTV LG	(SEQ ID NO: 41)	(SEQ ID NO: 42)	QVQLVQSGTEVKKPGASVKVSCKASGYTFPGYMH WVRQAPGQGLEWMGWNPDSGGTKYTKQFQGRVTL TRDASVSTAYIDLNRLRSDDTAVYYCARERCRTTNC YLDYWGQGS LVTVSS

i	P	S	:	5	5	0	6	4	3	G	7	8	1	0	336	336	N	A	GACATCCAGATGACCCAGTCTCCATCCTCCCTGTCT GCATCTGTAGGAGACATTCTCACCATCACTTGCCG GGCAAGTCAGAACATTACCACCTATTTAAATGGT ATCAGCAGAAACCAAGGAAAGCCCTAACCTCCTG ATCTCTGCTGCATCCCGTTTGCGAAGTGGGTCCCA TCAAGGTTTCAGTGGCAGTGGATCTGGGACAGATTT CAACTCTACCATCAGCAGTCTGCAACCTGTAGATTT TACAACTTTCTACTGTCAACAGACTTTCATAACCC GTGGACGTTTCGGCCAAGGGACCAAGGTGGAGATC AAACGA	(SEQ ID NO: 43)	DIQMTQSPSSLASVGDILTITCRASQNITTYLNWYQQ KPGKAPNLLISAAASRLSGVPSRFSGSGTDFTLTIS LQPVDFTTFFCYCQQTFTTPWTFGQGTKVEIKR	(SEQ ID NO: 45)	GACATCCAGATGACCCAGTCTCCATCCTCCCTGTCT GCATCTGTAGGAGACAGAGTCACCATCACTTGCCG GGCAAGTCAGGGCATTAGAAATGATTTAGGCTGGT ATCAGCAGAAACCAAGGAAAGCCCTAAGCGCT GATCTATCCTGCATCCAGTTTACAAAGTGGGTTC CATCAAGTTTCAGCGGAGTGGATCTGGGACAGAA TTCACTCTCAATCAGCAGCCTGCAGCCTGAAGA TTTTGCAACTTATTACTGTCTACAGCATATAAGTTA CCCTCTCACTTTCGGCGGAGGGACCAAGGTGGAGA TCAAGCGA	(SEQ ID NO: 47)	DIQMTQSPSSLASVGDVRVTITCRASQGIKNDLWYVQ QKPGKAPKRLIYPASSLQSGVPSRFSGSGTEFTLTIS SLQPEDFATYYCLQHSNYPLTFGGGKVEIKR	(SEQ ID NO: 49)																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																		
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<div><div>i</div><div>2</div><div>1</div><div>P</div><div>S</div><div>:</div><div>5</div><div>5</div><div>6</div><div>0</div><div>2</div><div>0</div></div>	<div><div>(R20</div><div>3)G₋</div><div>v1₋</div><div>VH₋</div><div>PE₋₁</div><div>3623</div><div>526</div><div>H</div><div>6</div></div>	N A A	GACATCCAGATGACCCAGTCTCCATCTCCCTGTCT GCATCTGTAGGAGACAGAGTCACCATCACTTGCCG GGCAGGTGAGAACATTAGCAGGCATTTAAATTGGT ATCAGCAGAATCCAGGAAAGCCCTAAGGTCTCTG ATCCATCTGCATCCAGTTTGCCAAAGTGGGTCCC GTCAAAGGTTCACTGGCAGTGGATCTGGGACAGATT TCAGTCTTACCATCAGCAGTCTGCAACCTGAAAGAT TTTGGAACCTTACTTCTGTCAACAGAGTTACAGTACC CCTCCCACCTTTCGGCGGAGGACCAAGGTGGAGCT CAAACGA (SEQ ID NO: 51)	CAGTGCAAACTGGTGCAGTCTGGGGCTGAAGTGAA GAAAGCTGGGGCTCAGTGAAGGTCTCTGCAAGG CTTCTGGATACACCTTCCCCGGCTACTATATGACT GGTTGCGACAGGCCCTGGACAAGGACTTGAGTGG ATGGATGGATCAACCTGACAGTGGTGACACAAA CTATGCACAGAAATTTACAGGCAAGGTACCATGA CCAGGACACGTCCATCAGCACAGCCTTTATGGAG CTGAGCAGGCTGAGATCAGACGACACGCGCTGTA TTACTGTGCGAGAGAGAAGCCAGATATTTGACT CCTTCTACTACTACCTTATGGACGTCTGGGGCCAAG GGACCAGGTCAACCGTCTCCTCA (SEQ ID NO: 52)
			DIQMTQSPSSLSASVGDRTVITCRAGQNIIRHLNWYQ QNP GKAPKVLHPASSLPSGVPSRFSGSGTDFSLTIS SLQPEDFGTYFCQQSYSTPPTFGGGTKVELKR (SEQ ID NO: 53)	QVQLVQSGAEVKKPGASVKVSKASGYTFPGYMY WLRQAPQGQGLEWMGWINPDSGDTNVAQKFQGRVT MTRDTSISIAFMELSLRSDDTAVYYCAREKPRYFDS FYYYLMDVWVGQGTTVTVSS (SEQ ID NO: 54)

Table 20A: CD40 Antibody VL CDRs

IPS#	Ab	Type	CDR1	CDR2	CDR3
iPS:55 6524	21- 230_33 H9	NA	ACCTTACGCAGTGGCATCAATGTTGGTT CCTCCAGGATATAT (SEQ ID NO: 55)	TACACATCAGACTCAGATAAAT TGCAGGGCTCT (SEQ ID NO: 56)	ATGATTTGGCACAGCAGCGCTG TGGTA (SEQ ID NO: 57)
		AA	TLRSGINVGSSRIY (SEQ ID NO: 58)	YTSDDSKLQGS (SEQ ID NO: 59)	MIWSSAVV (SEQ ID NO: 60)
		NA	AGGGCCAGTCAGAGTGTAGCAGCAAC TACTTAGCC (SEQ ID NO: 61)	GCTGCATCCAAACAGGGCCGCT (SEQ ID NO: 62)	CAGCAGTATGTTAGCTACCCG TCACT (SEQ ID NO: 63)
iPS:55 5854	21- 230_36 F3	AA	RASQSVSSNYLA (SEQ ID NO: 64)	AASNRAA (SEQ ID NO: 65)	QQYGSSPLT (SEQ ID NO: 66)
		AA			

iPS:55 5836	21- 230_29 H10	NA	CGGGCGAGTCAGGACATTAGCAATAAT TTAGGC	GCTGCATCCAGTTTGGCACAGT	CAACAGTATAATAGTTACCCCTC TCACT
		AA	(SEQ ID NO: 67) RASQDISNNLA (SEQ ID NO: 70)	(SEQ ID NO: 68) AASSLHS (SEQ ID NO: 71)	(SEQ ID NO: 69) QQYNSYPLT (SEQ ID NO: 72)
iPS:55 5842	21- 230_30 B9	NA	CGGGCAAGTCAGGGCATTAGAAATGAT TTAGGC	GCTGCATCCAGTTTGGCAAAGT	CTACAGTATAATAGTTACCCCGT GGACG
		AA	(SEQ ID NO: 73) RASQGIRNDLG (SEQ ID NO: 76)	(SEQ ID NO: 74) AAFSLQS (SEQ ID NO: 77)	(SEQ ID NO: 75) LQYNSYPWT (SEQ ID NO: 78)
iPS:55 5844	21- 230_31 H6	NA	CGGGCAAGTCAGGGCATTAGAAATGAG TTAGGC	GCTGCATCCAGTTTGGCACAGT	CTACAATATAGTGGGTCCCTC TCACT
		AA	(SEQ ID NO: 79) RASQGIRNELG (SEQ ID NO: 82)	(SEQ ID NO: 80) AASSLDS (SEQ ID NO: 83)	(SEQ ID NO: 81) LQYSGSPLT (SEQ ID NO: 84)
iPS:55 6022	21- 230_33 H12	NA	CGGGCAAGTCAGGACATTAGAAATGAT TTAGGC	GGTGCATCCAAATTTGGCAAAGT	CTACAACATAATATTGCTCCGC TCACC
		AA	(SEQ ID NO: 85) RASQDIRNDLG (SEQ ID NO: 88)	(SEQ ID NO: 86) GASNLQS (SEQ ID NO: 89)	(SEQ ID NO: 87) LQHNIAPLT (SEQ ID NO: 90)
iPS:55 5860	21- 230_37 A6	NA	TCTGGAGAAAGGTTGGGAAATAAATAT ATTGTC	CAAGATTTCAGCGGCCCTCA	CAGCGTGGGACAGCAGAACT GTGGTA
		AA	(SEQ ID NO: 91) SGERLGNKYIC (SEQ ID NO: 94)	(SEQ ID NO: 92) QDFKRPS (SEQ ID NO: 95)	(SEQ ID NO: 93) QAWDSRTVV (SEQ ID NO: 96)
iPS:55 5840	21- 230_30 A12	NA	ACTGGAACACAGCAGTGATGTTGGGAATT ATAACCTTGCTCC	GAGGTCAATCAGCGGCCCTCA	TGCTCATATACAACTAGTAGCA CTTATGTGATA
		AA	(SEQ ID NO: 97) TGTSSDVGNYNLVS (SEQ ID NO: 100)	(SEQ ID NO: 98) EVNQRP (SEQ ID NO: 101)	(SEQ ID NO: 99) CSYTTSSSTYVI (SEQ ID NO: 102)

iPS:55 5852	21- 230_35 F11	NA	ACTGGAACACAGCAGTGATGTTGGTGGTT ATATCTTTGTCTCC	GATGTCAGTAAGCGGGCCCTCT	TGCTCATATGCAGGCAACTACA CTTATGTC
		AA	TGTSSDVGGYIFVS (SEQ ID NO: 103)	DVSKRPS (SEQ ID NO: 104)	(SEQ ID NO: 105)
iPS:55 5862	21- 230_39 C2	NA	ACTGGAACACAGCAGTGATGTTGGGAATT ATAACCTTGCTCC	GAGGTCAATAGCGGCCCTCA (SEQ ID NO: 107)	(SEQ ID NO: 108)
		AA	TGTSSDVGNYNLVS (SEQ ID NO: 109)	EVNRRPS (SEQ ID NO: 110)	(SEQ ID NO: 111)
iPS:55 6018	21- 230_4 G7	NA	CGGGCAAGTCAGAACATTACCACCTATT TAAAT	GCTGCATCCCCGTTTGCGAAGT (SEQ ID NO: 113)	(SEQ ID NO: 114)
		AA	RASQNITTYLN (SEQ ID NO: 115)	AASRLRS (SEQ ID NO: 116)	(SEQ ID NO: 117)
iPS:55 5826	21- 230_17 H5	NA	CGGGCAAGTCAGGGCATTAGAAAATGAT TTAGGC	CCTGCATCCAGTTTACAAAGT (SEQ ID NO: 119)	(SEQ ID NO: 120)
		AA	RASQGIKNDLG (SEQ ID NO: 121)	PASSLQS (SEQ ID NO: 122)	(SEQ ID NO: 123)
iPS:55 6020	21- 230_33 H6	NA	CGGGCAGGTCAGAACATTAGCAGGCAT TTAAAT	CCTGCATCCAGTTTGCCCAAGT (SEQ ID NO: 125)	(SEQ ID NO: 126)
		AA	RAGQNISRHLN (SEQ ID NO: 127)	PASSLPS (SEQ ID NO: 128)	(SEQ ID NO: 129)
					QQSYSTPPT (SEQ ID NO: 132)

Table 20B: CD40 Antibody VH CDRs

iPS#	Ab	Type	CDR1	CDR2	CDR3
iPS:55 6524	21- 230_3 3H9	N	AGCCATGGCATG	GTTATCTGGTATGATGGAAGTAATGAATACT	GGGGGGGCCACTGGAACTACGAGGGCCA
		A	CAC	ATGGAGACTCCGTGAAGGGC	CTACTATGGTATGGACGTC
		A	(SEQ ID NO: 133)	(SEQ ID NO: 134)	(SEQ ID NO: 135)
		A	SHGMH	VIWYDGSNEYYGDSVKG	GGHWNVEGHYYGMDV
iPS:55 5854	21- 230_3 6F3	A	(SEQ ID NO: 136)	(SEQ ID NO: 137)	(SEQ ID NO: 138)
		N	AGCAGCCGTACT	AGGACATACTACAGGTCCAAAGTGGTATCATG	GGGGCTGCTCCCTTTGACTAC
		A	GCTTGGAAAC	ATTATTCAGTATCTGTGAAAAAGT	
		A	(SEQ ID NO: 139)	(SEQ ID NO: 140)	(SEQ ID NO: 141)
iPS:55 5836	21- 230_2 9H10	A	SSRTAWN	RTYYRSKWYHDYSVSVKS	GAAPFDY
		A	(SEQ ID NO: 142)	(SEQ ID NO: 143)	(SEQ ID NO: 144)
		N	GGCTACTATATG	TGGATCAACCCCTCACAGTGGTGGCACAAACT	GAACGTATTTCTATGTTCTGGGGGAGTCGGG
		A	CAC	ATGCACAGAAAGTTTCAGGAC	CACAACTGGTTCGCCCCC
iPS:55 5842	21- 230_3 0B9	A	(SEQ ID NO: 145)	(SEQ ID NO: 146)	(SEQ ID NO: 147)
		A	GYVMH	WINPHSGGTNYAQKFQD	ERISMVRGVGHNVFAP
		A	(SEQ ID NO: 148)	(SEQ ID NO: 149)	(SEQ ID NO: 150)
		N	GCCTACTATATA	TGGCTCAACCCCTGACAGTGGTGGCACAAACT	GAGAAAGTTTAACTACAACTATGGTGCTTTT
iPS:55 5844	21- 230_3 1H6	A	CAC	TTGCCCCGAGGTTTCAGGAC	GATATC
		A	(SEQ ID NO: 151)	(SEQ ID NO: 152)	(SEQ ID NO: 153)
		A	AYYIH	WLNPDSSGTNFAPRFQD	EKENYNYGAFDI
		A	(SEQ ID NO: 154)	(SEQ ID NO: 155)	(SEQ ID NO: 156)
iPS:55 5844	21- 230_3 1H6	N	GACTACTATATA	TGGATCAACCCCTAACAGTGGTGGCACAAAGCT	GAGGTGGGGCTATGGTTCGGGGACCTA
		A	CAC	ATGCACAGAAAGTTTCAGGAC	CCGCTATTACGGTATGGACGTC
		A	(SEQ ID NO: 157)	(SEQ ID NO: 158)	(SEQ ID NO: 159)
		A	DYYIH	WINPNSGDTSYAQKFQD	EVGGYSGTYRYYGMDV
		A	(SEQ ID NO: 160)	(SEQ ID NO: 161)	(SEQ ID NO: 162)

iPS:55 6022	21- 230_3 3H12	N	GACTACTATATT CAC	TGGATCAACCCTAACAGTGGCGGCACAACT ATGCACAGAAAGTTTCAGGGC	GAAGATGGGGCAGCAGTTGGTCCCCTCTA CTACTACTACGGTATGGACGTC
		A	(SEQ ID NO: 163)	(SEQ ID NO: 164)	(SEQ ID NO: 165)
iPS:55 5860	21- 230_3 7A6	A	DYYIH	WINPNSGGTNYAQKFQ	EDGAAVGPLYYYGMDV
		A	(SEQ ID NO: 166)	(SEQ ID NO: 167)	(SEQ ID NO: 168)
iPS:55 5840	21- 230_3 0A12	N	GACTACTACATG AGC	TATATTAGTCGAAAGTGGTGATACCATATACT ACGCAGACTCTGTGAAGGGC	GACTTAGCAGCAGGTGCTACAGGGGGCCT TGACTGC
		A	(SEQ ID NO: 169)	(SEQ ID NO: 170)	(SEQ ID NO: 171)
iPS:55 5852	21- 230_3 5F11	A	DYYMS	YISRSGDTIYYADSVKG	DLAAGATGGLDC
		A	(SEQ ID NO: 172)	(SEQ ID NO: 173)	(SEQ ID NO: 174)
iPS:55 5862	21- 230_3 9C2	N	AGAAATGCCATG AGT	GCTACTGGTGGTAGTGGTATTAGCACATACT ACGCAGACTCCGTGAAGGGC	GGTTATAGCAACAGCTGGTGGTACTTTGAC TAC
		A	(SEQ ID NO: 175)	(SEQ ID NO: 176)	(SEQ ID NO: 177)
iPS:55 6018	21- 230_4 G7	A	RNAMS	ATGSGISTYYADSVKG	GYSNSWWYFDY
		A	(SEQ ID NO: 178)	(SEQ ID NO: 179)	(SEQ ID NO: 180)
iPS:55 6022	21- 230_3 3H12	N	AGCTATGGCATG CAC	GTTATATGGTATGATGGAAAGTAATAAATACT ATGCAGACTCCGTGAAGGGC	GATGGCCGGAACACTACGTCTACTTTGACAAAC
		A	(SEQ ID NO: 181)	(SEQ ID NO: 182)	(SEQ ID NO: 183)
iPS:55 6018	21- 230_3 9C2	A	SYGMH	VIWYDGSNKYYADSVKG	DGRNYYVFDN
		A	(SEQ ID NO: 184)	(SEQ ID NO: 185)	(SEQ ID NO: 186)
iPS:55 6018	21- 230_3 9C2	N	GGCTACTATATG CAC	TGGATCAACCCTGACAGTGGTGGCACAAGT ATACACAGAAAGTTTCAGGGC	GAGAGGTGTAGGACTACCAACTGCTATTG GACTAC
		A	(SEQ ID NO: 187)	(SEQ ID NO: 188)	(SEQ ID NO: 189)
iPS:55 6018	21- 230_3 9C2	A	GYMH	WINPDSGGTKYTQKFQ	ERCRTNVCYLDY
		A	(SEQ ID NO: 190)	(SEQ ID NO: 191)	(SEQ ID NO: 192)
iPS:55 6018	21- 230_4 G7	N	GGCTACTATATG CAC	TGGATCAACCCTGACAGTGGGAGGCACAACT TTGCACAGCAGTTTCAGGGC	GAGAAGATAACTATGACTGGTATTACTTT GACTAT
		A	(SEQ ID NO: 193)	(SEQ ID NO: 194)	(SEQ ID NO: 195)
iPS:55 6018	21- 230_4 G7	A	GYMH	WINPDSGGTNFAQQFQ	EKITMTGLYFDY
		A	(SEQ ID NO: 196)	(SEQ ID NO: 197)	(SEQ ID NO: 198)

iPS:55 5826	21- 230_1 7H5	N	GGCTACTATATA	TGGATCAACCCTGACAGTGGTGACACAACT	GAGAGGTTAACACATTTGGTGGTGACTG
		A	CAC	ATGCACAGAA GTTTCAGGGC	CTATTCCCAATGGTTCGACCCC
		A	GYIYH	(SEQ ID NO: 199)	(SEQ ID NO: 201)
		A	GYIYH	WINPDSGDTNYAQKFQ	ERLTHCGGDCYSQWFD
iPS:55 6020	21- 230_3 3H6	A	(SEQ ID NO: 202)	(SEQ ID NO: 203)	(SEQ ID NO: 204)
		N	GGCTACTATATG	TGGATCAACCCTGACAGTGGTGACACAACT	GAGAACCCACAGATATTTGACTCCTTCTAC
		A	TAC	ATGCACAGAA GTTTCAGGGC	TACTACCTTATGACGTC
		A	(SEQ ID NO: 205)	(SEQ ID NO: 206)	(SEQ ID NO: 207)
		A	GYMY	WINPDSGDTNYAQKFQ	EKPRYFDSFYYYLMDV
		A	(SEQ ID NO: 208)	(SEQ ID NO: 209)	(SEQ ID NO: 210)

Table 21: MSLN Antibody VL and VH Sequences

iPS #	Ab	Type	LC	HC
iPS: 563 560	7G 11	NA	GACATTGTGATGACTCAGTCTCCAGACTCCCTGGC	GAGTGCAGCTGGTGCAGTCTGGAGGAGGCTTGAT
			TGTGTCCTGGGCGAGAGGGCCACCATCAACTGCA	CCAGCCTGGGGGTCCCTGAGACTCTCCTGTGCAG
			AGTCCAGCCAGAGTGTGTTTATACAGCTCCCAACAT	TCCTGGGTTCACCGTCACTAGCAAGTTCATGACC
			AAGAACTACTTAGCTTGGTACCAAGCAAGCAAGG	TGGTCCGCCAGGCTCCAGGGAAGGGGCTGGAGT
			ACAGCTCCTAAGCTGCTCATTTACTGGGCATCTA	GGGTCTCAGTTATTTATAGCGGTGGTAAGACATAC
			CCCGAGAATCCGGGTCCCTGACCGATTCAAGTGGC	TACGCAGACTCCGTGAAGGCCGATTACCCATCTC
			AGCGGTCTGGGACAGATTTCACCTCACCATCAG	CAGAGACAATTCCAGAACACGCTGTATCTTCAAA
			CAGCTGCAGGCTGAAGATGTGGCAGTTTATTACT	TGAACAGCCTGAGAGCCGAGGACACGGCCGTGTA
			GTCAGCAATATTATAGTACTCTCCGACGTTCCGGC	TTACTGTCCGAGAGATAGCGGTGGCTGGGGTACT
			CAAGGGACCAAGGTGGAGATCAAAACG	TTGACTACTGGGGCCAGGGAACCCCTGGTCAACCGTG
			(SEQ ID NO: 211)	TCCTCA
			(SEQ ID NO: 212)	(SEQ ID NO: 213)
	AA		DIVMTQSPDSLAVSLGERATINCKSSQSVLYSSNNKN	EVQLVESGGGLIQPGSLRLSCAVSGFTVSSKFMW
			YLA WYQQKPGQPKLLIYWASTRESGVDRFSGSGS	VRQAPGKGLWVSVIYSGGKTYVADSVKGRFTISR
			GTDFTLTISSLQAEDVAVYYCQYQYSTPTFGQGTKV	NSKNTLYLQMNLSRAEDTAVYYCARDSSGGWGYFDY
			EIKR	WGQGLVTVSS
			(SEQ ID NO: 213)	(SEQ ID NO: 214)
			(SEQ ID NO: 213)	(SEQ ID NO: 214)

iPS: 563 637	6F 4	NA	GACATCCAGATGACCCAGTCTCCATCTTCCGTGTCT GCTTCTGTCGAGACAGAGTCAACCATCACTTGTCTG GGCGAGTCAGGATATTAGCAGGTGGTTAGCTGGT ATCAGCAGAAACCCAGGAAAGCCCTAAGCTCCT GATTCTGCTGCATCCAGATTGCAAGTGGAGTCC CATCAAGGTTTCAGCGGCGAGTGGATCTGGGACAGAT TTCACCTCACCATCAGCAGCCTGCAGCCTGAAGA TTTTGCAATTACTATTGTCAACAGGCTAAAAGTTT TCCTCGGACGTTCCGGCCAAAGGACCAAGGTGGA ATCAAAACGA	CAGGTGCAGCTGGTGGAGTCTGGGGAGGCTTGGT CAAGCCTGGAGGTCCTGAGACTCTCCTGTGCAG CCTCTGATTACCTTCACTAGTACTACATGAGCT GGATCCGCCAGGCTCCAGGAAAGGGCTGGAGTG GATTTCATACATTAGTAGCAGTGAAGTATCATAT ATTACGTAGACTCTGTGAAGGGCCGATTCAACCATC TCCAGGGACAACGCCAAGAACTCACTGTATCTGCA AATGAACAGCCTGAGAGCCGAGGACACGGCCGTG TATTACTGTGCGAGAGATGTTGGGAGCCACTTGA CTACTGGGGCCAGGGAACCCCTGGTCAACCGTGTCT CA
			(SEQ ID NO: 215)	(SEQ ID NO: 216)
			DIQMTQSPSSVSASVGDRLVTITCRASQDISRWLAWYQ QKPGKAPKLLISAAASRLQSGVPSRFSGSGTDFTLTI SSLQPEDFAIYYCQQAQKSPRTFGQGTKVEIKR	QVQLVESGGGLVKPGGSLRLSCAASGFTFSYYMSW IRQAPGKGLEWISYSSSEIYYVDSVKGRFTISRDN KNSLYLQMNSLRAEDTAVYYCARDVGSFHDYWGQ GTLVTVSS
iPS: 344 086	4G 12	NA	(SEQ ID NO: 217)	(SEQ ID NO: 218)
			(SEQ ID NO: 219)	(SEQ ID NO: 220)
			QSVLTQPPSVSGAPGQRVTISCTGSSNIGAGYDVHW YQQVPGTAPKLLIYGNSKRPSGVPRFSGSKSGTSAS LAITGLQAEDEADYYCQSYDSSLGGWVFGGGTKLTV L	QVQLQESGPGLVKPSETLSLTCTVSGGSISSSYWYG WIRQPPGKGLEWIGSIYSGITNYPNPSLKSRTISVDI SKNQFSLKLSVTAADTAVYYCARPSNYDAFDIWGQ GTMVTVSS
iPS: 344 090	4H 6	NA	(SEQ ID NO: 221)	(SEQ ID NO: 222)
			(SEQ ID NO: 223)	(SEQ ID NO: 224)
			DIQMTQSPSSVSASVGDRLVTITCRASQGITRWLAWY QKPGKAPKLLIYAASVLQSGVPSRFSGSGTDFTL TISSLQPEDFATYYCQQSNFPRTFGQGTKVEIK	QVQLVESGGGLVKPGGSLRLSCAASGFTFSYYMTW IRQAPGKGLEWISYSSSGSTIYYADSVKGRFTISRDN AKNSLYLQMNSLRAEDTAVYYCARDNRNSHFDYWG QGTLVTVSS
iPS: 344 090	4H 6	AA	(SEQ ID NO: 225)	(SEQ ID NO: 226)

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Table 22A MSLN Antibody VL CDRs

iPS#	Ab	Ty pe	CDR1	CDR2	CDR3
iPS:563 560	7G 11	NA	AAGTCCAGCCAGAGTGTTTATACAGCTCCAACAATAA GAACTACTTAGCT	TGGCATCTACCCGA GAATCC	CAGCAATATTAGTACTCC TCCGACG
			(SEQ ID NO: 227)	(SEQ ID NO: 228)	(SEQ ID NO: 229)
		AA	KSSQSVLYSSNNKNYLA (SEQ ID NO: 230)	WASTRES (SEQ ID NO: 231)	QQYYSTPPT (SEQ ID NO: 232)
iPS:563 637	6F4	NA	CGGGCGAGTCAGGATATTAGCAGGTGGTTAGCC	GCTGCATCCAGATTG CAAAAGT	CAACAGGCTAAAAAGTTTTC TCGGACG
			(SEQ ID NO: 233)	(SEQ ID NO: 234)	(SEQ ID NO: 235)
		AA	RASQDISRWLA (SEQ ID NO: 236)	AASRLQS (SEQ ID NO: 237)	QQAASFPR (SEQ ID NO: 238)
iPS:344 086	4G 12	NA	<i>No nuc. seq. available</i> (SEQ ID NO: 239)	<i>No nuc. seq. available</i> (SEQ ID NO: 240)	<i>No nuc. seq. available</i> (SEQ ID NO: 241)
		AA	TGSSNIGAGYDVH (SEQ ID NO: 242)	GNSKRPS (SEQ ID NO: 243)	QSYDSSLGGVV (SEQ ID NO: 244)
		NA	<i>No nuc. seq. available</i> (SEQ ID NO: 245)	<i>No nuc. seq. available</i> (SEQ ID NO: 246)	<i>No nuc. seq. available</i> (SEQ ID NO: 247)
iPS:344 090	4H 6	AA	RASQGITRWLA (SEQ ID NO: 248)	AASVLQS (SEQ ID NO: 249)	QQNSFPRT (SEQ ID NO: 250)

Table 22B MSLN Antibody VH CDRs

iPS#	Ab	Ty pe	CDR1	CDR2	CDR3
iPS:563 560	7G 11	NA	AGCAAGTTCAT GACC	GTTATTTATAGCGGTGGTAAGACATACTACGCAGACTCC GTGAAGGGC	GATAGCGGTGGCTGGGGGTACT TTGACTAC
			(SEQ ID NO: 251)	(SEQ ID NO: 252)	(SEQ ID NO: 253)
		AA	SKFMT	VIYSGGKITYYADSVKG	DSGGWGYFDY

		(SEQ ID NO: 254)	(SEQ ID NO: 255)	(SEQ ID NO: 256)
		GACTACTACAT GAGC	TACATTAGTAGCAGTGAAAGTATCATATATTACGTAGAC TCTGTGAAGGGC	GATGTTGGGAGCCACTTTTGA C
iPS:563 637	6F4	NA	(SEQ ID NO: 257)	(SEQ ID NO: 259)
		AA	DYYMS YISSESIYYVDSVKG	DVGSHEFY
		NA	(SEQ ID NO: 260)	(SEQ ID NO: 262)
		AA	No nuc. seq. available	No nuc. seq. available
iPS:344 086	4G 12	NA	(SEQ ID NO: 263)	(SEQ ID NO: 265)
		AA	SSSYWYG SIYSGITNYPNLSKS	PSNYDAFDI
		NA	(SEQ ID NO: 266)	(SEQ ID NO: 268)
		AA	No nuc. seq. available	No nuc. seq. available
iPS:344 090	4H 6	NA	(SEQ ID NO: 269)	(SEQ ID NO: 271)
		AA	DYYMT YISSSGSTIYYADSVKG	DRNSHEFY
		NA	(SEQ ID NO: 272)	(SEQ ID NO: 274)

Table 23: MSLN scFv Full Sequences

iPS#	Ab	Type	SCFV
		NA	No nuc. seq. available
		NA	(SEQ ID NO: 275)
		AA	EVQLVESGGGLIQPGGSLRLSCAVSGFTVSSKFMWVRQAPGKGLEWVSVIYSGGKTTY ADSVKGRFTISR NSKNTLYLQMNLSRAEDTAVYYCARDSSGGWGYFDYWGQGTLLVTVSSGGGGGGGGSDIVMTQSP DSLAVSLGERATINCKSSQSVLYSSNNKNYLA WYQQKPGQPPKLLIYWASTRESGVPDFRSGSGSDTDFTLTI SSLQAEDVAVYYCQQYYSTPTFTGGQGTKVEIK
		NA	(SEQ ID NO: 276)
		AA	No nuc. seq. available
		NA	(SEQ ID NO: 277)

iPS:3 44086	4G12	AA	QVQLVESGGGLVKPGGSLRLSCAASGFTFSDDYYMSWIRQAPGKGLEWISYISSSEIIYYVDSVKGRFTISRDN AKNSLYLQMNSLRAEDTAVYYCARDVGSHEFYWGQGTLLVTVSSGGGGSGGGGGSGS DIQMTQSPSSVSASVGDRTVITTCRASQDISRWLA WYQQKPGKAPKLLISAAASRLQSGVPSRFSGSGSGTDFTL TISSLQPEDFAIYYCQAKSFPRFTFGQGTKVEIK
		NA	(SEQ ID NO: 278) <i>No info. seq available</i>
		AA	(SEQ ID NO: 279) QVQLQESGPGLVKPSSETLSLTCTVSGGSISSSSYYWGWIROPKPKGLEWIGSIYSGITNYYNPSLKSRVTISVD TSKNQFSLKLSSTVAADTAVYYCARPSNYDAFDIWWGQGTMTVTVSSGGGGSGGGGGSGQSVLTQPPSVS GAPGQRVTISCTGSSSNIGAGYDVHWYQQVPGTAPKLLIYGNSKRPSGVPDRFSGSKSGTASLAITGLQAE EADYYCQSYDSSSLGGWVFGGGLTLTVL
iPS:3 44090	4H6	NA	(SEQ ID NO: 280) <i>No info. seq available</i>
		AA	(SEQ ID NO: 281) QVQLVESGGGLVKPGGSLRLSCAASGFTFSDDYYMTWIRQAPGKGLEWISYISSSGSTIYYADSVKGRFTISR NAKNSLYLQMNSLRAEDTAVYYCARDNRSHFDYWGQGTLLVTVSSGGGGSGGGGGSGDIQMTQSPSSV SASVGDRTVITTCRASQGITRWLA WYQQKPGKAPKLLIYAASVLSQSGVPSRFSGSGSGTDFTLTISSLQPEDFA TYYCQQSQNSFPRTFGQGTKVEIK
		NA	(SEQ ID NO: 282)

Table 24A: CD40-MSLN IgG-scFv Full Sequences

Ab	SCFV HC	LC
ipS#	ty p e	

		<p>CAGGTGCAGTGGTGCAGTCTGGGGCTGAGGTGAAGAAG CCTGGGGCTCAGTGAAGGTGTCTCTGCAAGGCTTCTGGAT ACACCTTCGCCGGCTACTATATGCACTGGTGGACAGGC CCCTGGACAAGGCTTGAGTGGATGGATGGATCAACCT GACAGTGGAGGCACAAACTTTGCACAGAGTTTCAGGGCA GGTCAACATGACACAGGGATACGTCCATCAGCACAGCCTA CATGGAGGTGAGCAGGCTGAGATCTGACGACACGGCCGT GTTTTACTGTGCGAGAGAGAAGATCACTATGACTGGTATT TACTTTGACTATTGGGGCCAGGAACCTTGGTCAACCTGT CCTCAGCTCCACCAAGGGCCATCGGTCTTCCCCCTGGC ACCTCTCCAAAGACACCTCTGGGGCACAGGGCCCTG GGTGCCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGG TGCTGTGGAACCTCAGGCGCCCTGACCAAGCGGTGCACAC CTTCCCGGCTGTCTACAGTCTCAGGACTCTACTCCCTCA GCAGCGTGTGACCGTGCCCTCCAGCAGTTGGGCACCCA GACCTACATCTGCAACGTGAATCACAAAGCCAGCAACACC AAGGTGGACAAGAAAGTTGAGCCCAAAATCTTGTGACAAA ACTCACACATGCCACCGTGCCACGACCTGAACCTCTGG GGGACCGTCACTCTCTCTTCTTCCCCCCTCAACCCAAAGGA CACCTCATGATCTCCGGACCCCTGAGGTCAATGCGTG GTGTGGACGTGAGCCACGAAGACCTGAGGTCAAGTTCA ACTGTAACGTGGACGGCTGGAGGTGCATAATGCCAAGAC AAAGCCGTGCGAGGAGCAGTACGGCAGCAGTACCGTTG CGTACGCGTCTCACCCGTCTGCAACCAAGGACTGGCTGAAT GGCAAGGAGTACAAGTGCAAGGTGTCCAAACAAAGCCCTC CCAGCCCCATCGAGAAACCATCTCCAAAGCCAAAGGGC AGCCCCGAGAACACAGGTGTACACCTGCCCCCTATCCCG GGAGGAGATGACCAAGAACAGGTGACGCTGACCTGCCT GGTCAAAGGCTTCTATCCAGCGACATCGCCGTGGAGTGG GAGAGCAATGGCAGCCGGAGAACACTACAAGACCACG CCTCCCGTGTGACTCCGACGGCTCTTCTTCTCTCTATAG CAAGCTCACCGTGGACAAAGAGAGGTGGCAGCAGGGGAA CGTCTTCTCATGCTCCGTGATGATGAGGCTGTCACACACC ACTACACGCAGAGACCTCTCCCTGTCTCCGGGTGGTG CGGATCGGGAGGTGGCGGATCCCAAGGTGCAGCTGGTCGA</p>
		<p>N A</p>
	<p>21- 230 4 G7 Ig G 21- 233 4 H6 3</p>	<p>iP S: 5 7 7 3 8 3</p>
		<p>T</p>

	GTCTGGGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGA CTCTCTGTGCAGCTCTGGATTCACTTCAAGTGACTACTA CATGACCTGGATCAGGCAGGCTCCAGGGAAGTGCCTGGAG TGGATTTACATTAAGTAGTAGTGGTAGTACCATCTACTA CGCAGACTCTGTGAAGGGCCGATTCAACCATCTCCAGGGAC AAGGCCAAGAACTCACTGTATCTGCAAAATGAACAGCCTGA GAGCCGAGGACACGGCCGTGTATTACTGTGGGAGAGATCG GAATCCCACTTTGACTATTGGGGCCAGGGAACCCCTGGTC ACCGTGTCCTCAGGAGGGGGCGGATCTGGCGCGGAGGA AGCGGTGGGGCGGCTCCGACATTCAGATGACCCAGTCTC CATCTCCGTGTCTGCATCTGTAGGGGACAGAGTCAACCAT CACTTGTGGGGAGTCAGGTATTACAGGTGGTTAGCC TGGTATCAGCAGAAACCAAGGAAAGCCCCCTAAGCTCCTGA TCTATGCTGCATCCGTTTGCAAAAGTGGGTCCCATCAAG GTTCAAGCGCAGTGGATCTGGGACAGATTTCACCTCAAC ATCAGCAGCCTGCAGCCTGAAGATTGCAACTTACTATT GTCAACAGTCTAACAGTTCCCTCGGACGTTCCGGCTGCGG GACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 283)
		(SEQ ID NO: 284)

		<p>QVQLVQSGAEVKKPGASVKVCKASGYTFAGYYMHWVRQ APQGGLWGMWINPDSGGTNFAQQFQGRVTMTRDTSISTAY MEVSRLRSDDTAVFYCAREKITMTGIYFDYWGGQTLVTVSS ASTKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWN SGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHPKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP KPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHN AKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKKVSNK ALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFFLYSKLT VDKSRWQQGNVFSCSVMHAEALHNHYTQKSLSLSPGGGGSG GGGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWI RQAPGKCLEWISYISSGSTIYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDNRNSHFDYWGGQTLVTVSSGGG GSGGGGGGGGSDIQMTQSPSSVSASVGDRTITTCRASQGIT RWLA WYQQKPGKAPKLLIY AASVLSQSGVPSRFSGSGSGTDF TLTISSLQPEDFATYYCQQSNSFPRTFGCGTKVEIKR</p>	<p>DIQMTQSPSSLSASVGDILTTITCRASQNITTYLNWYQQKPGKA PNLLISAASRLRSGVPSRFSGSGGTDFLTITISLQPVDFTTFY CQQTFTTPWTFGGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C</p>
	A	(SEQ ID NO: 285)	(SEQ ID NO: 286)
	A		

		<p>CAGGTGCAGTGGTGCAGTCTGGGGCTGAGGTGAAGAAG CCTGGGGCTCAGTGAAGGTGTCTCTGCAAGGCTTCTGGAT ACACCTTCGCCGGCTACTATATGCACTGGTGGCAGACAGC CCCTGGACAAGGGCTTGAGTGGATGGATGGATCAACCT GACAGTGGAGGCACAAACTTTGCACAGCAAGTTTCAGGGCA GGTCAACCATGACCAGGGATACGTCCATCAGCACAGCCTA CATGGAGGTGAGCAGGCTGAGATCTGACGACACGGCCGT GTTTTACTGTGCGAGAGAGAAGATCACTATGACTGGTATT TACTTTGACTATTGGGGCCAGGAACCTTGGTCAACCTGT CCTCAGCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGC ACCTCTCCAAAGACACCTCTGGGGCACAGGGCCCTG GGTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGG TGTCTGTGAACTCAGCGCCCTGACCAAGCGGTGCACAC CTTCCCGGTGTCTACAGTCTCAGGACTCTACTCCCTCA GCAGCGTGTGACCGTGCCCTCCAGCAAGTTGGGCACCCA GACCTACATCTGCCAACGTGAATCACAAAGCCAGCAACACC AAGGTGGACAAGAAAGTTGAGCCCCAAATCTTGTGACAAA ACTCACACATGCCACCGTGCCACGACCTGAACTCCTGG GGGACCGTCAAGTCTTCTCTTCCCCCCTGCAACCCAAAGGA CACCTCATGATCTCCCGGACCCCTGAGGTCAATGCGTG GTGTGGACGTGAGCCACGAAGACCTGAGGTCAAGTTCA ACTGTAACGTGGACGGCTGGAGGTGCATAATGCCAAGAC AAAGCCGTGCGAGGAGCAAGTACGGCAGCACGTACCGTTG CGTACGCGTCTCACCGTCTCTGCCACCAAGGACTGGCTGAAT GGCAAGGAGTACAAGTGCAAGGTGTCCAAACAAGCCCTC CCAGCCCCATCGAAGAACCATCTCCAAAGCCAAAGGGC AGCCCCGAGAACACAGGTGTACACCTGCCCCCTATCCCG GGAGGAGATGACCAAGAACCAAGGTGACGCTGACCTGCCT GGTCAAAGGCTTCTATCCAGCGACATCGCCGTGGAGTGG GAGAGCAATGGCAGCCGGAGAACAACTACAAGACCACG CCTCCCGTGTGGACTCCGACGGCTCTTCTTCTCTATAG CAAGCTCACCGTGGACAAGAGAGGTGGCAGCAGGGGAA CGTCTTCTCATGCTCCGTGATGATGAGGCTCTGCACAACC ACTACACGCAGAGACCTCTCCCTGTCTCCGGGTGGTG CGGATCGGGAGGTGGCGGATCCCCAGGTGCAGCTGGTGG</p>
		<p>N A</p>
	<p>21- 230 4 G7 Ig G 21- 233 6F 4 scF v</p>	
iP S: 5 7 7 3 8 8		

	GTCTGGGGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGA CTCTCTGTGCAGCCTCTGGATTCACTTCAAGTACTACTA CATGAGCTGGATCCGCCAGGCTCCAGGGAAGTGCCTGGAG TGGATTTTCATACATTAGTAGCAGTGAAAGTATCATCTATTA CGTAGACTCTGTGAAGGGCCGATTCAACCATCTCCAGGGAC AAGCCAAAGAACTCACTGTATCTGCAAAATGAACAGCCTGA GAGCCGAGGACACGGCCGTGTATTACTGTGGGAGAGATGT TGGGAGCCACTTTGACTACTGGGGCCAGGGAACCCCTGGTC ACCGTGTCCTCAGGAGGGGGCGGATCTGGCGGCGGAGGA AGCGGTGGGGCGGC'TCCGACATCCAGATGACCCAGTCTC CATCTCCGTGTCTTCTGTCTGGAGACAGAGTCAACCATC ACTTGTCTGGGCGAGTCAGGATATTAGCAGGTGGTTAGCCT GGTATCAGCAGAAACCAAGGAAAGCCCTAAGCTCCTGAT TTCTGCTGCATCCAGATTGCAAGTGGAGTCCCATCAAGG TTCAGCGGCAGTGGATCTGGGACAGATTTCACCTCTCACC TCAGCAGCCTGCAGCCTGAAGATTTTGCAATTACTATTGT CAACAGGCTAAAAGTTTTCCTCGGACGTTCTGGCTGCGGGA CCAAAGGTGGAATCAAAACGG	(SEQ ID NO: 287)
		(SEQ ID NO: 288)

			QVQLVQSGAEVKKPGASVKVSCKASGYTFAGYYMHWVRQ APQGGLWGMWINPDSGGTNFAQQFQGRVTMTRDTSISTAY MEVSRLRSDDTAVFYCAREKITMTGIYFDYWGGQTLVTVSS ASTKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWN SGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP KPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHN AKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKKVSNK ALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFFLYSKLT VDKSRWQQGNVFSCSVMHAEALHNHYTQKSLSLSPGGGGSG GGGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWI RQAPGKCLEWISYSSSEIYYVDSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDVGSHPDYWGQGTLLVTVSSGGGG SGGGSGGGGSDIQMTQSPSSVSASVGDRTITTCRASQDISR WLAWYQQKPGKAPKLLISAAASRLQSGVPSRFSGSGGTDFTL TISSLQPEDFAIYYCQQAQKSFPRTFGCGTKVEIKR (SEQ ID NO: 289)	DIQMTQSPSSLSASVGDILTITCRASQNITTYLNWYQQKPGKA PNLLISAAASRLRSGVPSRFSGSGGTDFTLTISSLQPVDFITFY CQQTFTTPWTFGGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C (SEQ ID NO: 290)
		A		
		A		

148

	GCTGGTCAGTCTGGGGAGGCTTGGTCAAGCCTGGAGGG TCCCTGAGACTCTCCTGTGCGAGCCTCTGGATTACCTTCAG TGACTACTACATGACCTGGATCAGGCAGGCTCCAGGGAAG TGCCTGGAGTGGATTTCATACATTAGTAGTAGTGGTAGTA CCATCTACTACGACAGACTCTGTGAAGGGCCGATTCACCAT CTCCAGGACAACGCCAAGAACTCACTGTATCTGCAAAATG AACAGCCTGAGAGCCGAGGACACGGCCGTGTATTACTGTG CGAGAGATCGGAACCTCCACTTTGACTATTGGGCCAGGG AACCTGGTCAACCGTGTCTCAGGAGGGGCGGATCTGGC GGCGGAGGAAGCGGTGGGGCGGCTCCGACATTCAGATG ACCCAGTCTCCATCTCCGTGTCTGCACTGTAGGGGACAG AGTCAACCATCACTTGTCCGGCGAGTCAGGGTATTACAGG TGGTTAGCCTGGTATCAGCAGAACCAAGGAAGCCCTA AGCTCCTGATCTATGCTGCATCCGTTTGGCAAAGTGGGGTC CCATCAAGGTTTCAGCGGCAGTGGATCTGGGACAGATTTC CTCTCACCATCAGCAGCCTGCAGCCTGAAGATTTGCAAC TTACTATTGTCAACAGCTAACAGTTTCCCTCGGACGTTTCG GCTGCGGGACCAAGGTGGAATCAAAACGG	(SEQ ID NO: 291)
		(SEQ ID NO: 292)

		<p>QVQLVQSGAEVTKPGASVKVCKASGYTFAGYMHVVRQ APQGLEWMGWINPHSGGTINYAQKFQDRVTMTRDTSINTA YMELSRRLSDDTAVYYCARERISMVRGVGHNWFAPWGQGT LVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPV TVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSCDKTHCTPCPAPELLGGP SVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVD GVEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYK CKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQ VSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSF FLYSKLTVDKSRWQQGNVFSCVMHEALHNHYTQKSLSLSP GGGSGGGGSGVQLVESGGGLVKPGGSLRLSCAASGFTFSD YYMTWIRQAPGKCLEWISYISSSGSTIYYADSVKGRFTISRDN AKNSLYLQMNSLRAEDTAVYYCARDNRNSHFDYWGQGTLLVT VSSGGSGGGGGGGGSDIQMTQSPSSVSASVGDRTTTCR ASQGITRWLAWYQQKPGKAPKLLIYAASVLQSGVPSRFSGSG SGTDFTLTISLQPEDFATYYCQQSNSFPRTFGCGTKVEIKR</p>	<p>DIQMTQSPSSLSASVGDRTVTITCRASQDISNNLAWFQQKPGK PPKSLMYAASSLHSGVPTFSGSGGTDFTFTISSLQPEDFATY YCQYNSYPLTFGGGKVEIRRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK DSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC</p>
	A	(SEQ ID NO: 293)	(SEQ ID NO: 294)
	A		

ip

152

		<p>QVQLVQSGAEVTKPGASVKVCKASGYTFAGYYMHWVRQ APQGGLWGMGWINPHSGGTNYAQKFQDRVTMTRDTSINTA YMELSRRLRSDDTAVYYCARERISMVRGVGHNWFAPWGQGT LVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPV TVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSCDKTHTCPCPAPELLGGP SVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVD GVEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYK CKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQ VSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSF FLYSKLTVDKSRWQQGNVFSCVMHEALHNHYTQKSLSLSP GGGSGGGGSGVQLVESGGGLVKPGGSLRLSCAASGFTFSD YYMSWIRQAPGKCLEWISYISSSESIIYYVDSVKGRFTISRDN AKNSLYLQMNSLRAEDTAVYYCARDVGSFHDYWGQGTLLVT VSSGGGSGGGGGGGGDIQMTQSPSSVSASVGDRTITCR ASQDISRWLAWYQQKPGKAPKLLISAAASRLQSGVPSRFSGSG SGTDFTLTISLQPEDFAIYYCQQAQKSFPRTFGCGTKVEIKR</p>	<p>DIQMTQSPSSLSASVGDRTVTITCRASQDISNNLAWFQQKPGK PPKSLMYAASSLHSGVPTFSGSGGTDFTFTISSLQPEDFAFY YCQYNSYPLTFGGGKVEIRRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK DSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC</p>
	A	(SEQ ID NO: 297)	(SEQ ID NO: 298)
	A		

		<p>GAGGTGCAGCTGCTGGAGTCTGGGGGAGGCTTGGTACAGC CTGGGGGTCCCTGAGACTCTCTGTGAGCCCTCTGGATTCT ACCTTTAGTAGAAATGCCATGAGTTGGTCCGCCAGGCTC CAGGGAAGGGGCTGAGTGGGTGCAGCTACTGGTGTGA GTGGTATTAGCACATACTACGCAGACTCCGTGAAGGGCCG GTTACCATCTCCAGAGACAATTCCAAGAACACGCTGTAT CTGCAAAATGAACAGTCTGAGAGCCGAGGACACGGCCGTAT ATTACTGTGCGAGAGGTTATAGCAACAGCTGGTGGTACTT TGACTACTGGGGCCAGGGAACCTTGTCACCGTGTCTCA GGCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTC CTCCAAAGACACCTCTGGGGGCACAGGGCCCTGGGCTGC CTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGT GGAACTCAGGCGCCCTGACCAAGCGCGTGCACACCTTCCC GGCTGTCTTACAGTCTCAGGACTCTACTCCCTCAGCAGC GTGGTGACCGTGCCTCCAGCAGCTTGGGACCCAGACCT ACATCTGCAACGTGAATCACAGCCAGCAACACCAAGGT GGACAAAGAAAGTTGAGCCCAAATCTTGTGACAAAACCTAC ACATGCCACCGTGCCAGCACCTGAACTCTTGGGGGAC CGTCAAGTCTTCTCTTCCCCCCTCAAAACCCAGGACACCCCTC ATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGGTG ACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTGGTA CGTGGACGGCGTGAGGTGCATAATGCCAAGACAAAGCC GTGCGAGGAGCAGTACGGCAGCAGTACCGTTGCGTCAGC GTCTCAACCGTCTGCAACCAAGGACTGGCTGAATGGCAAGG AGTACAAGTGCAAGGTGTCCAAACAAGCCCTCCCAGCCCC CATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCG AGAACACAGGTGTACACCTTGCCCCCATCCCCGGGAGGAG ATGACCAAGAACCAGGTACGCTGACCTGCTGGTGGTCAAG GCTTCTATCCAGCGACATCGCCGTGGAGTGGGAGAGCAA TGGGCAGCCGGAGAACAACTACAAAGCCACGCCCTCCCGTG CTGGACTCCGACGGCTCTTCTTCTCTATAGCAAGCTCAC CGTGGACAAGACAGGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGCACACCACTACACG CAGAAAGAGCCTCTCCCTGTCTCCGGGTGGTGGGATCGG GAGGTGGCGGATCCCAGGTGCAGCTGGTGCAGTCTGGGGG</p>
		N A
	21- 230 _30 A1 2_1 5_gG 7_21 7_- 4_233 _4 H6 _s cFv	
iP S: 5 7 7 4 0 1		

	AGGCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCCTGT GCAGCCTCTGGATTACCTTCAGTGACTACTACATGACCTG GATCAGGCAGGCTCCAGGGAAGTGCCCTGGAGTGGATTTC TACATTAGTAGTAGTGGTAGTACCATCTACTACGCAGACT CTGTGAAGGGCCGATTACCATCTCCAGGGACAACGCCAA GAACCTCACTGTATCTGCAAAATGAACAGCCTGAGAGCCGAG GACACGGCCGTGTATTACTGTGCGAGAGATCGGAACTCCC ACTTTGACTATTGGGGCCAGGGAACCCCTGGTCAACCGTGC CTCAGGAGGGGGCGGATCTGGCGCGGAGGAAAGCGGTGG GGCGGCTCCGACATTCAGATGACCCAGTCTCCATCTTCC GTGTCTGCATCTGTAGGGGACAGAGTCAACATCACTTGT GGCGAGTCAGGTATTACAGGTGGTTAGCCTGGTATCA GCAGAAACCAGGGAAGCCCTAAGCTCCTGATCTATGCT GCATCCGTTTGC AAAGTGGGTCCCATCAAGGTTCAAGC GCAGTGGATCTGGGACAGATTTCACCTCACCATCAGCAG CCTGCAGCCTGAAGATTTTGCAACTTACTATTGTCAACAGT CTAACAGTTTCCCTCGGACGTTCCGGCTGCGGGACCAAGGT GGAAATCAAACGG	(SEQ ID NO: 299)
		(SEQ ID NO: 300)

		EVQLLESGGGLVQPGGSLRLSCAASGFTFSRNAMSWVRQAP GKGLEWVSATGGGISTYYADSVKGRFTISRDNKNTLYLQ MNSLRAEDTAVYYCARGYSNSWYFDYWGGGTLVTSSA STKGPSVFLAPSSKSTSGGTALGCLVKDYFPEPVTVSWNS GALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKHTHTCPPCPAPELGGPSVFLFPPK PKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTPPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGGGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWIR QAPGKCLEWISISSGSTIYYADSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDNRNSHFDYWGGGTLVTSSGGGG SGGGSGGGGSDIQMTQSPSSVSASVGDRTITCRASQGTR WLAWYQQKPGKAPKLLIYAASVLSQGVPSRFSGSGSGTDFT LTISSLQPEDFATYYCQQSNSFPRTFGCGTKVEIKR (SEQ ID NO: 301)	QSALTQPAVSVSGSPGQSITISCTGTSSDVGNYNLVSWYQQHP GKAPKLMIFEVNRQPSGVSNRFSGSKSGTTASLTISGLQAAD ADYFCCSYTTSSTYVIFGGGKLTVLGQPKAAPSVTLFPPSSE ELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTPPS KQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS (SEQ ID NO: 302)
	A		
	A		

		<p>GAGGTGCAGCTGCTGGAGTCTGGGGGAGGCTTGGTACAGC CTGGGGGTCCCTGAGACTCTCTGTGAGCCCTCTGGATTCT ACCTTTAGTAGAAATGCCATGAGTTGGTCCGCCAGGCTC CAGGGAAGGGGTGAGTGGGTGTGAGCTACTGGTGTGA GTGGTATTAGCACATACTACGCAGACTCCGTGAAGGGCGG GTTACCATCTCCAGAGACAAATTCGAAGAACACGCTGTAT CTGCAAAATGAACAGTCTGAGAGCCGAGGACACGGCCGTAT ATTACTGTGCGAGAGGTTATAGCAACAGCTGGTGGTACTT TGACTACTGGGGCCAGGGAACCTTGTCACCGTGTCTCA GGCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTC CTCCAAAGACACCTCTGGGGGCACAGGGCCCTGGGCTGC CTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGT GGAACTCAGGCGCCCTGACCAAGCGCGTGCACACCTTCCC GGCTGTCTTACAGTCTCAGGACTCTACTCCCTCAGCAGC GTGGTGACCGTCCCTCCAGCAGCTTGGGACCCAGACCT ACATCTGCAACGTGAATCACAGCCAGCAACACCAAGGT GGACAAAGAAAGTTGAGCCCAAATCTTGTGACAAAACCTAC ACATGCCACCCGTGCCAGCACCTGAACTCTTGGGGGAC CGTCAAGTCTTCTCTTCCCCCCTCAAAACCCAGGACACCCCTC ATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGGTG ACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTGGTA CGTGGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCC GTGCGAGGAGCAGTACGGCAGCACGTACCGTTGCGTCAGC GTCTCAACCGTCTGCACCAAGGACTGGCTGAATGGCAAGG AGTACAAGTGCAAGGTGTCCAAACAAGCCCTCCCAGCCCC CATCGAGAAAACCATCTCCAAAGCCAAAGGGCAGCCCCG AGAACCAAGGTGTACACCTTGCCCCCATCCCCGGGAGGAG ATGACCAAGAACCAGGTACGCTGACCTGCCCTGGTCAAAAG GCTTCTATCCAGCGACATCGCCGTGGAGTGGGAGAGCAA TGGGCAGCCGGAGAACAACTACAAAGCACCGCTCCCGTG CTGGACTCCGACGGCTCTTCTTCTCTATAGCAAGCTCAC CGTGGACAAGACAGGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGCACACCACTACAG CAGAAAGAGCCTCTCCCTGTCTCCGGGTGGTGGGATCGG GAGGTGGCGGATCCCAGGTGCAGCTGGTGGAGTCTGGGG</p>
		N A
21- 230 _A1 2_1 _gG _21 _ - 233 _6F _4_ scF v		
iP S: 5 7 7 4 0 5		

	<p>GAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCCTG TGCAGCCTCTGGATTCACTTTCAGTACTACTACATGAGCT GGATCCGCCAGGCTCCAGGAAAGTGCCCTGGAGTGGATTTC ATACATTAGTAGCAGTGAAAGTATCATCTATTACGTAGAC TCTGTAAAGGGCCGATTCAACCATCTCCAGGGACAACGCCA AGAACTCACTGTATCTGCAAAATGAACAGCCTGAGAGCCGA GGACACGGCCGTGTATTACTGTGCGAGAGATGTTGGGAGC CACTTTGACTACTGGGGCCAGGGAACCCCTGGTCAACCGTGT CCTCAGGAGGGGGCGGATCTGGCGCGGAGGAAAGCGGTG GGGCGGCTCCGACATCCAGATGACCCAGTCTCCATCTTC CGTGTCTGCTTCTGTCGGAGACAGAGTCACCATCACTTGTC GGCGAGTCAGGATATTAGCAGGTGGTTAGCCTGGTATCA GCAGAAACCAGGAAAGCCCTAAGCTCCTGATTCTGCT GCATCCAGATTGCAAGTGAGTGCCCATCAAGGTTCAAGCG GCAGTGGATCTGGGACAGATTTCACCTCACCATCAGCAG CCTGCAGCCTGAAGATTTTGCAATTTACTATTGTCAACAGG CTAAAAGTTTCTCGGACGTTTCGGCTGCGGGACCAAGGT GGAAATCAAACGG</p>	(SEQ ID NO: 303)
		(SEQ ID NO: 304)

		EVQLLESGGGLVQPGGSLRLSCAASGFTFSRNAMSWVRQAP GKGLEWVSATGGGISTYYADSVKGRFTISRDNKNTLYLQ MNSLRAEDTAVYYCARGYSNSWYFDYWGGGTLVTSSA STKGPSVFLAPSSKSTSGGTALGCLVKDYFPEPVTVSWNS GALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSCDKHTHTCPPCPAPELGGPSVFLFPPK PKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQYGSTYRCVSLTVLHQDWLNGKEYKCKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGGGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIR QAPGKCLEWISYSSSEIYYVDSVKGRFTISRDNKNSLYLQ MNSLRAEDTAVYYCARDVGSFHDYWGQGTLVTVSSGGGGS GGGGGGGGSDIQMTQSPSSVSASVGDRVTTTCRASQDISRW LAWYQQKPGKAPKLLISAAASRLQSGVPSRFSGSGGTDFTLTI SSLQPEDFAIYYCQQAQKSFPRTFGCGTKVEIKR (SEQ ID NO: 305)	QSALTQPAVSVSGSPGQSITISCTGTSSDVGNYNLVSQYQHP GKAPKLMIFEVNRQPSGVSNRFGSKSGTTASLTISGLQAAD ADYFCCSYTTSSTYVIFGGGKLTVLGQPKAAPSVTLFPPSSE ELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTPPS KQSNNKYAASSYLSTLTPQWKSHRYSQVTHEGSTVEKTV APTECS (SEQ ID NO: 306)
	A		
	A		

[illegible]

	TGCAGCTGGTCGAGTCTGGGGGAGGCTTGGTCAAGCCTGG AGGTCCCTGAGACTCTCTGTGCAGCCTCTGGATTACACCT TCAGTGACTACTACATGACCTGGATCAGGCAGGCTCCAGG GAAGTGCCTGGAGTGGATTTCATACATTAGTAGTAGGT AGTACCATCTACTACGACAGACTCTGTGAAGGGCCGATTCA CCATCTCCAGGGACAACGCCAAGAACTCACTGTATCTGCA AATGAACAGCCTGAGAGCCGAGGACACGGCCGTGTATTAC TGTCCGAGAGATCGGAACCTCCACTTTGACTATTGGGGCC AGGGAACCCCTGGTCACCGTGTCTCAGGAGGGGGCGGATC TGGCGCGGAGGAAAGCGGTGGGGCGGC'TCCGACATTCA GATGACCCAGTCTCCATCTTCCGTGTCTGTCATCTGTAGGGG ACAGAGTCACCATCACTTGTGGGGGAGTCAGGGTATTAC CAGGTGGTTAGCCTGGTATCAGCAGAAACCAGGGAAAGC CCCTAAGCTCCTGATCTATGCTGCATCCGTTTGGCAAAGTG GGGTCCCATCAAGGTTCA GCGGCA GTGGATCTGGGACAGA TTTCACTCTCACCATCAGCAGCCTGCAGCCTGAAGATTTG CAACTTACTATTGTCAACAGTCTAACAGTTTCCCTCGGACG TTCGGCTGCGGGACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 307)
		(SEQ ID NO: 308)

		<p>QVQLVQSGAEVKKPGASVKVCKASGYTFPGYYMYWLRQA PGGLEWMGWINDSGDTNYAQKFGQGRVTMTTRDTSISTAF MELSRLSDDTAVYYCAREKPRYFDSFYYYLMDVWQGQTT VTVSSASTKGPSVFLAPSSKSTSGGTAALGCLVKDYFPEPVT VSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPS VFLPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDG VEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKC KVSNAKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVS LTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFLL YSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLSLSPGG GGSGGGGSQLVESGGGLVKPGGSLRLSCAASGFTFSDYY MTWIRQAPGKCLEWISYISSSGSTIYYADSVKGRFTISRDNAL NSLYLQMNSLR AEDTAVYYCARDNRNSHFDYWGQGTLLVTVS SGGGSGGGGGGGGDIQMTQSPSSVSASVGDRTTTCRAS QGTRWLAWYQQKPGKAPKLLIYAASVLSQSGVPSRFSGSGG TDFTLTISSLQPEDFATYYCQQQSNFPRTFGCGTKVEIKR</p>
		<p>(SEQ ID NO: 309)</p>
		<p>DIQMTQSPSSLSASVGDRVTITCRAGQNISRHLNWKQNPQK APKVLIHPPASSLPSGVPSRFSGSGTDFSLTISSLQPEDFGTYF CQQSYSTPTTFGGGTVKVELKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C</p>
		<p>(SEQ ID NO: 310)</p>

		<div><div><div>21- 230</div><div>33</div><div>H6</div><div>Ig</div><div>G</div><div>21- 233</div><div>6F</div><div>4</div><div>scF</div><div>v</div></div><div>iP S: 5 7 7 4 1 3</div></div>	<div><div><div>CAGGTGCAACTGGTGCAGTCTGGGGCTGAAGTGAAGAAG CCTGGGGCTCAGTGAAGGTGTCTGCAAGGCTTCTGGAT ACACTTCCCCGGCTACTATATGTACTGGTTCGACAGGC CCCTGGACAAGGACTTGAGTGGATGGATGGATCAACCT GACAGTGGTGACACAACTATGCACAGAAGTTTCAGGGCA GGTCAACCATGACCAGGGACAGTCCATCAGCACAGCCTT TATGGAGCTGAGCAGGCTGAGATCAGACGACACGGCCGT GTATTACTGTGGAGAGAGAAAGCCAGATATTTGACTCC TTCTACTACTACCTTATGGACGTCTGGGGCCAGGGACCA CGGTCAACCGTGTCTCAGCCTCCACCAAGGGCCCATCGT CTTCCCCCTGGCAGCTCTCTCCAAAGAGCACCTCTGGGGG ACAGCGGCCCTGGCTGCCTGGTCAAGGACTACTTCCCG AACCAGTGACGGTGTCTGGAACTCAAGCGCCCTGACCAG CGCGTGCACACCTTCCCGGTGTCTACAGTCTCTCAGGA CTCTACTCCCTCAGCAGCGTGTGACCGTGCCCTCCAGCA GCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAA GCCAGCAACACCAAGGTGGACAAAGAAAGTTGAGCCCAA ATCTGTGACAAAACTCACACATGCCACCGTGCCCAAGCA CCTGAACCTCTGGGGGGACCGTCAAGTCTTCTCTTCCCCC AAACCCCAAGGACACCTCATGATCTCCCGGACCCCTGAG GTCACATGCGTGGTGGTGGACGTGACCAAGAACCCCTG AGGTCAGGTTCAACTGGTACGTGGACGGCGTGGAGGTGCA TAATGCCAAGACAAAGCCGTGCCAGGAGCAGTACGGCAG CAGGTACCGTTTGGCTCAGCGTCTCTCACCCTCCTGCACCA GACTGGCTGAATGGCAAGGAGTACAAGTGCAAGGTGTCC AACAAAGCCCTCCAGCCCCCATCGAGAAACCATCTCCA AAGCCAAAGGGCAGCCCCGAGAACACACAGGTGTACACCC TGCCCCCATCCCCGGGAGGAGATGACCAAGAACAGGTCA GCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGACAT CGCGTGGAGTGGGAGAGCAATGGCAGCGCGGAGAACAA CTACAAGACCACGCTCCCGTGTGGACTCCGACGGCTCC TTCTTCTCTATAGCAAGCTACCGGTGGACAAAGAGCAGGT GGCAGCAGGGGAACGCTTCTCTCATGCTCCGTGATGCATGA GGCTCTGCACAACCACTACACGCAAGAGAGCCTCTCCCTG TCTCCGGGTGGTGGCGGATCGGGAGGTGGCGGATCCCCAGG</div></div></div>	<div><div><div>GACATCCAGATGACCCAGTCTCCATCTCCTCCCTGTCTGCATC TGTAGGAGACAGAGTCAACCATCTTGCCGGGCAGGTCAAG AACATTAGCAGGCATTTAAATTGGTATCAGCAGAATCCAG GGAAAGCCCCCTAAGTCTCTGATCCATCTGCTCCAGTTT GCCAAGTGGGTCCCGTCAAGGTTCAAGTGGCAGTGGATCT GGGACAGATTTCAGTCTTACCATCAGCAGTCTGCAACCTG AAGATTTTGGAACTTACTTCTGTCAACAGAGTTACAGTAC CCCTCCCACCTTTCGGCGGAGGGACCAAGGTGGAGCTCAAA CGAACGGTGGCTGCACCATCTGTCTTCTATCTTCCCCTCATC TGATGAGCAGTTGAAATCTGGAACTGCCTCTGTGTGTGC CTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGT GGAAGGTGGATAACGCCCTCCAATCGGGTAAC'TCCCAGGA GAGTGTACAGAGCAGGACAGCAAGGACAGCACCTACAG CCTCAGCAGCACCTGACGCTGAGCAAGCAGACACTACGAG AAACACAAAGTCTAGGCTGCGGAAGTCAACCATCAGGGCC TGAGCTCGCCCCGTCAACAAAGAGCTTCAACAGGGGAGAGTG</div></div><div>T</div></div>
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	TGCAGCTGGTGGAGTCTGGGGAGGCTTGGTCAAGCCTGG AGGTCCCTGAGACTCTCTGTGCAGCCTCTGGATTACACCT TCAGTGACTACTACATGAGCTGGATCCGCCAGGCTCCAGG GAAGTGCCTGGAGTGATTTTCATACATTAGTAGCAGTGAA AGTATCATCTATTACGTAGACTCTGTGAAGGCCGATTCA CCATCTCCAGGGACAACGCCAAGAACTCACTGTATCTGCA AATGAACAGCCTGAGAGCCGAGGACACGGCCGTGTATTAC TGTCCGAGAGATGTTGGGAGCCACTTTGACTACTGGGGCC AGGGAACCCCTGGTCACCGTGTCTCAGGAGGGGGCGGATC TGGCGCGGAGGAAGCGGTGGGGCGGC'TCCGACATCCA GATGACCCAGTCTCCATCTTCCGTGTCTGTCTTCTGTGCGGAG ACAGAGTCACCATCACTTGTGGGGCGAGTCAAGGATATTAG CAGGTGGTTAGCCTGGTATCAGCAGAAACAGGGAAAGC CCCTAAGCTCCTGATTCTGTGCTGATCCAGATTGCAAAAGTG GAGTCCCATCAAGGTTCA GCGGCA GTGGATCTGGGACAGA TTTCAC'TCACCATCAGCAGCCTGCAGCCTGAAGATTGTG CAATTACTATTGTCAACAGGCTAAAAGTTTTCCTCGGACG TTCGGCTGCGGGACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 311)
		(SEQ ID NO: 312)

		<p>QVQLVQSGAEVKKPGASVKVCKASGYTFPGYYMYWLRQA PGQLEWMGWINPDSGDTNYAQKFGQGRVTMTTRDTSISTAF MELSRLSDDTA VYYCAREKPRYFDSFY YLMDVWQGQTT VTVSSASTKGPSVFLAPSSKSTSGGTAALGCLVKDYFPEPVT VSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHCTCPCPAPELLGGPS VFLPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDG VEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKC KVSNAKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVS LTCLVKGYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFLL YSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLSLSPGG GGSGGGGSQLVESGGGLVKPGGSLRLSCAASGFTFSDYY MSWIRQAPGKCLEWISYISSESIIYYVDSVKGRFTISRDNAL NSLYLQMNSLRAEDTAVYYCARDVGSFHDYWGQGTLLVTVS SGGGSGGGGGGGGDIQMTQSPSSVSASVGDRTTTCRAS QDISRWLA WYQQKPGKAPKLLISAA SRLQSGVPSRFSGSGG TDFTLTISSLQPEDFAIYYCQQA KSFPRTFGCGTKVEIKR</p>	<p>DIQMTQSPSSLSASVGDRVTITCRAGQNISRHLN WYQQNPGK APKVLIHPPASSLPSGVPSRFSGSGGTD FSLTISSLQPEDFGTYF CQQSYSTPTTFGGGTVKVELKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLN NFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C</p>
	A	(SEQ ID NO: 313)	(SEQ ID NO: 314)
	A		

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	GCAGCTGGTCGAGTCTGGGGAGGCTTGGTCAAGCCTGGA GGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTACCTT CAGTGACTACTACATGACCTGGATCAGGACGGCTCCAGGG AAGTGCCTGGAGTGGATTTCATACATTAGTAGTGGTA GTACCATCTACTACGCAGACTCTGTGAAGGCCGATTAC CATCTCAGGGACAACGCCAAGAACTCACTGTATCTGCAA ATGAACAGCCTGAGAGCCGAGGACACGGCCGTGTATTACT GTGCGAGAGATCGGAACCTCCACTTTGACTATTGGGGCCA GGGAACCCCTGGTCAACCGTGTCTCAGGAGGGGGCGGATCT GGCGCGGAGGAAAGGTGGGGCGGCTCCGACATTACAG ATGACCCAGTCTCCATCTTCCGTGTCTGCACTCTGTAGGGGA CAGAGTCACCATCACTTGTCCGGCGAGTCAGGTATTACC AGTGGTTAGCCTGGTATCAGCAGAAACCAGGGAAGCC CCTAAGCTCCTGATCTATGCTGCATCCGTTTTCGAAAAGTGG GGTCCCATCAAGGTTCAAGGCGAGTGGATCTGGGACAGAT TTCACCTCACCATCAGCAGCCTGCAGCCTGAAGATTTC AACTTACTATTGTCAACAGTCTAACAGTTTCCCTCGGACGT TCGGCTGCGGGACCAAGGTGGAATCAAAACGG	(SEQ ID NO: 315)
		(SEQ ID NO: 316)

		<p>QVQLVESGGGVVQGRSLRLSCAASGFTFSHGMHWVRQPP GKGLEWVAIVYDGSNEYYGDSVKGRFTISRDN SKNTLYLQ MNSLRVEDTAVYYCTRGGGHWNVEGHYYGMDVWVQGTT VTVSSASTKGPSVFLAPSSKSTSGTAALGCLVKDYFPEPVT VSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHCTCPAPELLGGPS VFLPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDG VEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKC KVSNAKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVS LTCLVKGYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFLL YSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGG GGSGGGGSQLVESGGGLVKPGGSLRLSCAASGFTFSYY MTWIRQAPGKCLEWISYISSSGSTIYADSVKGRFTISRDN AK NSLYLQMNSLR AEDTAVYYCARDNRNSHFDYWGQGTLLVTVS SGGGSGGGGGGGGGDIQMTQSPSSVSASVGDRTITCRAS QGTRWLAWYQQKPGKAPKLLIYAASVLSQSGVPSRFSGSGG TDFTLTISLQPEDFATYYCQQSNSFPRTFGCGTKVEIKR</p>
		<p>(SEQ ID NO: 317)</p>
		<p>(SEQ ID NO: 318)</p>

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	GCAGCTGGTGGAGTCTGGGGAGGCTTGGTCAAGCCTGGA GGTCCCTGAGACTCTCTGTGCAGCCTCTGGATTACCTT CAGTGACTACTACATGAGCTGGATCCGCCAGGCTCCAGGG AAGTGCCTGGAGTGGATTTCATACATTAGTAGCAGTGAAA GTATCATCTATTACGTAGACTCTGTGAAGGGCCGATTAC CATCTCAGGGACAACGCCAAGAACTCACTGTATCTGCAA ATGAACAGCCTGAGAGCCGAGGACACGGCCGTGTATTACT GTGCGAGAGATGTTGGAGCCACTTTGACTACTGGGGCCA GGGAACCCCTGGTACCGTGTCTCAGGAGGGGGCGGATCT GGCGCGGAGGAAAGGTGGGGCGGC'TCCGACATCCAG ATGACCCAGTCTCCATCTTCCGTGTCTTCTGTCTGGAGA CAGAGTCACCATCACTTGTGGGGCGAGTCAGGATATTAGC AGGTGGTTAGCCTGGTATCAGCAGAAACCAGGGAAAGCC CCTAAGCTCCTGATTCTGTGCTGCCATCCAGATTGCAAAGTGG AGTCCCATCAAGGTTTCAGCGGCAGTGGATCTGGGACAGAT TTCACCTCACCATCAGCAGCCTGCAGCCTGAAGATTTC AATTACTATTGTCAACAGGCTAAAAGTTTTCCTCGGACGT TCGGCTGCGGGACCAAGGTGGAATCAAAACGG	(SEQ ID NO: 319)
		(SEQ ID NO: 320)

		<p>QVQLVESGGGVVQGRSLRLSCAASGFTFSHGMHWVRQPP GKGLEWVAIVYDGSNEYYGDSVKGRFTISRDN SKNTLYLQ MNSLRVEDTAVYCTRGGGHWNYEGHYYGMDVWQGQTT VTVSSASTKGPSVFLAPSSKSTSGTAALGCLVKDYFPEPVT VSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDKTHCTCPAPELLGGPS VFLPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDG VEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKC KVSNAKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVS LTCLVKGYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFL YSKLTVDKSRWQQGNVFCSCVMHEALHNHYTQKSLSLSPGG GGSGGGGSQLVESGGGLVKPGGSLRLSCAASGFTFSDYY MSWIRQAPGKCLEWISYSSSESIYYVDSVKGRFTISRDN AK NSLYLQMNSLRAEDTAVYYCARDVGSFHDYWGQGTILVTVS SGGGSGGGGGGGGGDIQMTQSPSSVSASVGDRVTTTCRAS QDISRWLA WYQQKPGKAPKLLISAA SRLQSGVPSRFSGSGG TDFTLTISSLQPEDFAIYYCQCAKSPRTFGCGTKVEIKR</p>	<p>QAVPTQPSSLSASPGASASLTCTLRSGINVGSSRIYWYQQKPG SPQFLLRYTSDSDKLQSGGVPSRFSGSKDASANAGLLISGL QSEDEADYYCMIWHSSAVVFGGGTKLTVLGQPKAAPSVTLF PPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVE TTTTPSKQSNNKYAASSYLSLTPEQWKSRSYSCQVTHEGSTV EKTVAPTPCS</p>
	A		(SEQ ID NO: 321)
	A		(SEQ ID NO: 322)

		<div>CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGGTCCAGC CTGGAGGTCCTGAGACTCTCTGTGCAGCGTCTGGAATT CACCTCAGTAGCTATGGCATGCACTGGTCCGCCAGGCT CCAGGCAAGGGCTGGAGTGGTGGCAGTTATCTGGTATG ATGGAAGTAATAAATACTATGCAGACTCCGTGAAGGGCGG AGTCACCATCTCCAGAGACAATTCCAAGAAACACGCTGTAT CTGCAAAATGAATAGCCTGAGAGCCGAGGACACGGCTGTGT ATTACTGTACGAGAGATGGCCGGAACACTACGTCTATTGA CAACTGGGGCCAGGAACCCCTGGTCAACCGTGTCTCAGCC TCCACCAAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTCCTC CAAGAGCACCTCTGGGGGACAGGGGCCCTGGGCTGCCTG GTCAAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGTGA ACTCAGGCGCCCTGACCAGCGCGTGCACACCTTCCCGGC TGTCCTACAGTCCCTCAGGACTCTACTCCCTCAGCAGCGTGG TGACCGTGGCCTCCAGCAGCTTGGGCACCCAGACCTAGAT CTGCAACGTGAATCACAAGCCCAAGCAACACCAAGGTGGA CAAGAAAGTTGAGCCCAATCTTGTGACAAAACTCACACA TGCCACCCGTGCCAGCACCTGAACCTCTGGGGGACCGT CAGTCTTCTCTTCCCCCAAAACCCAAAGGACACCCCTCATG ATCTCCCGGACCCCTGAGGTACATGCGTGGTGGACG TGAGCCACGAAGACCTGAGGTCAAAGTTCAACTGGTACGT GGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGTG CGAGGAGCAGTACGGCAGCACGTACCGTTGCGTCAGCGTC CTCACCGTCTGCAACCAAGGACTGGCTGAATGGCAAGGAGT ACAAAGTGCAAGGTGTCCAAACAAGCCCTCCAGCCCCCAT CGAGAAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGA ACCAAGGTGTACACCTTGCCCCCATCCCGGAGGAGATG ACCAAGAACCAAGTCAAGCTGACCTGGCTGGTCAAAGGCT TCTATCCACGACATCGCCGTGGAGTGGAGAGCAATGG GCAGCCGGAGAACAACTACAAAGACCACGCTCCCGTGTCTG GACTCCGACGGCTCTTCTTCTCTATAGCAAGCTCACCGT GGACAAGAGCAGGTGGCAGCAGGGGAACGCTTCTCATG CTCCGTGATGCATGAGGCTCTGCACAACCACTACACGAG AAGAGCCTCTCCCTGTCTCCGGGTGGTGGCGGATCGGGAG GTGGCGGATCCCAAGGTGCAGCTGGTCTGAGTCTGGGGGAGG</div>
		<div>N A</div>
	<div>21- 230 35 F11 Ig G 21- 233 4 H6 s cFv</div>	

	CTTGGTCAAGCCTGGAGGTCCCTGAGACTCTCCTGTGCA GCCTCTGGATTCACTTCAGTGACTACTACATGACCTGGAT CAGGCAGGCTCCAGGGAAGTGCCCTGGAGTGGATTTTCATAC ATTAGTAGTAGTGGTAGTACCATCTACTACGCAGACTCTG TGAAAGGGCCGATTACCATCTCCAGGGACACGCCAAGAA CTCACTGTATCTGCAATGAACAGCCTGAGAGCCGAGGAC ACGGCCGTGTATTACTGTGCGAGAGATCGGAACCTCCCACT TTGACTATTGGGCCAGGGAACCCCTGGTCAACCGTGTCCTC AGGAGGGGGCGGATCTGGCGCGGAGGAAAGCGGTGGGGG CGGCTCCGACATTCAGATGACCCAGTCTCCATCTTCCGTGT CTGCATCTGTAGGGGACAGAGTCAACCATCACTTGTCTGGGC GAGTCAGGGTATTACCAGGTGGTAGCCTGGTATCAGCAG AAACCAGGGAAGCCCTAAGCTCCTGATCTATGCTGCAT CCGTTTGCAAAAGTGGGTCCCATCAAGGTTCAAGCGGCAG TGGAATCTGGGACAGATTTCACCTCAACCATCAGCAGCCTG CAGCCTGAAGATTTTGCAACTTACTATTGTCAACAGTCTAA CAGTTTCCCTCGGACGTTTCGGCTGCGGGACCAAGGTGGAA ATCAAAACGG	(SEQ ID NO: 323)
		(SEQ ID NO: 324)

		<p>QVQLVESGGGVVQGRSLRLSCAASGFTLSSYGMHWVRQAP GKGLEWVAIVYDGSNKYYADSVKGRVTISRDN SKNTLYL QMNSLRAEDTAVYYCTRDGRNYYFDNWQGGLVTVSSAS TKGPSVFLPAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSG ALTSQVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNH KPSNTKV DKKVEPKSCDKTHTCPCPAPELLGGPSVFLFPPKP KDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGSGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWIR QAPGKCLEWISYSSSGSTIYYADSVKGRFTISRDN AKNSLYL QMNSLRAEDTAVYYCARDNRNSHFDYWGQGLVTVSSGGGG SGGGSGGGGSDIQMTQSPSSVSASVGDRTITCRASQGITR WLAWYQQKPGKAPKLLIY AASVLSQGVPSRFSGSGSGTDFT LTISSLQPEDFATYYCQQSNSEFPRTFGCGTKVEIKR</p>	<p>QSALTQPRSVSGSPGQSVTISCTGTSSDVGGYIFVSWYQQHPG KAPKLMYDVSKRPSGVPDRFSGSKSVNTASLTISGLQAEDET DYCCSYAGNYTYVFGTGTKVTVLGQPKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPS KQSNNKYAASSYLSTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS</p>
	A		
	A		

		<p>CAGGTGCAGCTGGTGGAGTCTGGGGGAGGCGTGGTCCAGC CTGGGAGTCCCTGAGACTCTCTGTGACGCTCTGGAATT CACCTCAGTAGCTATGGCATGCACTGGTCCGCCAGGCT CCAGGCAAGGGCTGGAGTGGTGGCAGTTATCTGGTATG ATGGAAGTAATAAATACTATGCAGACTCCGTGAAGGGCG AGTCAACATCTCCAGAGACAATTCCAAGAAACACGCTGTAT CTGCAAAATGAATAGCTGAGAGCCGAGGACACGGCTGTGT ATTACTGTACGAGAGATGGCCGGAACACTACGTCTATTGA CAACTGGGGCCAGGAACCCCTGGTCAACCGTCTCTCAGCC TCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTCCTC CAAGAGCACCTCTGGGGGACAGGGGCCCTGGGCTGCCTG GTCAAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGGA ACTCAGGCGCCCTGACCAGCGCGTGCACACCTTCCCGGC TGTCCTACAGTCCCTCAGGACTCTACTCCCTCAGCAGCGTGG TGACCGTGGCCTCCAGCAGCTTGGGCACCCAGACTAGAT CTGCAACGTGAATCACAAGCCAGCAACACCAAGGTGGA CAAGAAAGTTGAGCCCAATCTTGTGACAAAACTCACACA TGCCACCCGTGCCAGCACCTGAACCTCTGGGGGACCGT CAGTCTTCTCTTCCCCCAAAACCCAAAGGACACCCCTCATG ATCTCCCGGACCCCTGAGGTACATGCGTGGTGGACG TGAGCCACGAAGACCTGAGGTCAAGTTCAACTGGTACGT GGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCCGTG CGAGGAGCAGTACGGCAGCACGTACCGTTGCGTCAGCGTC CTCACCGTCTGCAACAGGACTGGCTGAATGGCAAGGAGT ACAAGTGCAAGGTGTCCACAAAGCCCTCCAGCCCCCAT CGAGAAACCATCTCCAAAGCAAAAGGCGAGCCCCGAGA ACCAAGGTGTACACCTTGCCCCCATCCCGGAGGAGATG ACCAAGAACCAAGTACGCTGACCTGGCTGGTCAAAAGCT TCTATCCACGACATCGCCGTGGAGTGGAGAGCAATGG GCAGCCGGAGAACAACTACAAAGCCACGCTCCCGTCTG GACTCCGACGGCTCTTCTTCTCTATAGCAAGCTCACCGT GGACAAGAGCAGGTGGCAGCAGGGGAACGCTTCTCATG CTCCGTGATGCATGAGGCTCTGCACAACCACTACACGAG AAGAGCCTCTCCCTGTCTCCGGGTGGTGGCGGATCGGGAG GTGGCGGATCCCAAGGTGCAGCTGGTGGAGTCTGGGGGAG</p>
		<p>N A</p>
	<p>21- 230 35 F11 Ig G 21- 233 6F 4 scF v</p>	
iP S: 5 7 7 4 2 9		

	GCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCCTGTGC AGCCTCTGGATTCACTTTCAGTGACTACTACATGAGCTGG ATCCGCCAGGCTCCAGGGAAGTGCTGGAGTGGATTTCAT ACATTAGTAGCAGTGAAGTATCATCTATTACGTAGACTC TGTGAAGGGCCGATTCCACCATCTCCAGGACAAACGCCAAG AACTCACTGTATCTGCAAAATGAACAGCCTGAGAGCCGAGG ACAGGCCCTGTATTACTGTGCGAGAGATGTTGGGAGCCA CTTTGACTACTGGGGCCAGGGAACCTGGTCAACCGTGTC TCAGGAGGGGGCGGATCTGGGGCGGAGGAAGCGGTGGG GGCGCTCCGACATCCAGATGACCCAGTCTCCATCTCCG TGTCTGCTTCTGCGGAGACAGATCACCATCACTTGTGCG GCGAGTCAGGATATTAGCAGGTGGTTAGCCTGGTATCAGC AGAAACCAAGGAAAGCCCTAAGCTCCTGATTCTGCTGC ATCCAGATTGCAAGTGGAGTCCCATCAAGGTTCAAGCGGC AGTGGATCTGGACAGATTTCACTCTCACCATCAGCAGCC TGCAGCCTGAAGATTTTGCAATTTACTATTGTCAACAGGCT AAAAGTTTCTCGGACGTTGCGCTGCGGGACCAAGGTGG AAATCAAACGG	(SEQ ID NO: 327)
		(SEQ ID NO: 328)

		<p>QVQLVESGGGVVQPRSLRLSCAASGFTLSSYGMHWVRQAP GKGLEWVAIVWYDGSNKYYADSVKGRVTISRDN SKNTLYL QMNSLRAEDTAVYYCTRDGRNYYFDNWGQGLVTVSSAS TKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSG ALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNH KPSNTKV DKKVEPKSCDKTHCTCPCPAPELLGGPSVFLPPKP KDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQY GSTYRCVSVLTVLHQDWLNGKEYKCKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGSGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIR QAPGKCLEWISYSSSEIYYVDSVKGRFTISRDN AKNSLYLQ MNSLRAEDTAVYYCARDVGSFHDYWGQGLVTVSSGGGGS GGGSGGGGGDIQMTQSPSSVSASVGDRVTTITCRASQDISRW LAWYQQKPGKAPKLLISAA SRLQSGVPSRFSGSGGTDFTLTI SSLQPEDFAIYYCQQA KSFPRTFGCGTKVEIKR</p>	<p>QKALTPRSVSGSPGQSVTISCTGTSSDVGGYIFVSWYQQHPG KAPKLMYDVSKRPSGVPDRFSGSKSVNTASLTISGLQAEDET DYCCSYAGNYTYVFGTGT VTLGQPKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTPS KQSNNKYAASSYLSTPEQWKSHRSYSCQVTHEGSTVEKTV APTECS</p>
	A		
	A		

178

	GGTCAAGCCTGGAGGGTCCCTGAGACTCTCTCTGTGCAGCC TCTGGATTCACTTCACTGAGTACTACATGACCTGGATCAG GCAGGCTCCAGGGAAGTGCCTGGAGTGGATTTTCATACATT AGTAGTAGTGGTAGTACCATCTACTACGAGACTCTGTGA AGGCCGATTCAACCATCTCCAGGACAACGCCAAGAACTC ACTGTATCTGCAAAATGAACAGCCTGAGAGCCGAGGACACG GCCGTGTATTACTGTGCGAGAGATCGGAACCTCCACTTTG ACTATTGGGCCAGGGAACCTGGTCAACCGTGCTCCAGG AGGGCGGATCTGGCGGAGGAGGAGCGGTGGGGCGG CTCCGACATTCAAGATGACCCAGTCTCCATCTCCGTGCTG CATCTGAGGGGACAGAGTCAACATCACTTGTGGGCGAG TCAGGTATTACAGGTGGTAGCCTGGTATCAGCAGAAA CCAGGAAAGCCCCTAAGCTCCTGATCTATGCTGCATCCG TTTTGCAAGTGGGTCCCATCAAGGTTCAAGCGGCAAGTG ATCTGGGACAGATTTCACTCTCACCATCAGCAGCCTGCAG CCTGAAGATTTTGCAACTTACTATTGTCAACAGCTAACAG TTTCCCTCGGACGTTGGCTGCGGGACCAAGGTGGAATC AAACGG	(SEQ ID NO: 331)
		(SEQ ID NO: 332)

	QVQLQQSGPGLVKPSQTLSTCAISGDSVSSRTAWNWIQSL PSRLEWLGRTYYRSKWYHDYSVSVKSRITIDPDTSKNQFSL QLNSVTPEDTAVYYCARGAAPFDYWGGTLVTVSSASTKGP SVFPLAPSSKSTGGTAALGCLVKDYFPEPVTVSWNSGALTS GVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSN TKVDKKVEPKSCDKHTHTCPCPAPELLGGPSVFLFPPKPKDTL MISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRW QQGNVFSCSVMHEALHNHYTQKSLSLSPGGGGGGGGGSQVQ LVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWIRQAPGKC LEWISYSSSGSTIYYADSVKGRFTISRDNAKNSLYLQMNSLR AEDTAVYYCARDNRSHFDYWGGTLVTVSSGGGGGGGGGS GGGSDIQMTQSPSSVSASVGDRTITCRASQGITRWLAWYQ QKPGKAPKLLIYAASVLQSGVPSRFSGSGSGTDFTLTISSLQPE DFATYYCQQSNSFPRTFGCGTKVEIKR (SEQ ID NO: 333)	EIVLTQSPGTLSTSPGERATLSCRASQSVSSNYLAWYQQKPG QAPRALIYAASNRAAGISDRFSGSGGTDFTLTISRLEPEDFAV YFCQQYGSSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSG TASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSK DSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNR GEC (SEQ ID NO: 334)
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		<p>CAGGTACAGCTGCAACAGTCAAGTCCAGGACTGGTGAAGC CCTCGCAGACCCCTCTACACCTGTGCCATCTCCGGGAC AGTGTCTAGCAGCCGTACTGCTTGGAACTGGATCAGGC AGTCCCATCGAGAGCCCTTGAGTGGCTGGGAAGGACATA CTACAGGTCCAAGTGGTATCATGATTATTCAGATCTGTGA AAAGTGAATCACCATCGACCCAGACACATCAAGAACCA GTTCTCCCTGCAAGCTGAACCTGTGACTCCCGAGGACACG GCTGTTTATTATTGTGCAAGAGGGGCTGCTCCCTTTGACTA CTGGGGCCAGGGAACCCCTGGTCAACCGTGTCTCAGCCCTC ACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTCCCTCCA AGAGCACTCTGGGGGCACAGGGCCCTGGGCTGCCTGGT CAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGGGAAC TCAGGCGCCCTGACCAGCGCGGTGCACACCTTCCCGGCTG TCCTACAGTCCCTCAGGACTCTACTCCCTCAGCAGCGTGGTG ACCGTGGCCCTCCAGCAGCTTGGGCACCCAGACCTACATCT GCAACGTGAATCACAAGCCCCAGCAACACCAAGGTGGACA AGAAAGTTGAGCCCCAAATCTTGTGACAAAACACACATG CCCACCGTGCCAGCACCTGAACCTCTGGGGGACCGTCA GTCTTCTCTTCCCCCACAACCCAAAGGACACCCCTCATGAT CTCCCGGACCCCTGAGGTACATGCGTGGTGGACGTG AGCCACGAAGACCCCTGAGGTCAAGTTCAACTGGTACGTGG ACGGCGTGGAGGTGCATAATGCCAAGACAAGCCCGTGGC AGGAGCAGTACGGCAGCAGTACCGTTGCGTCAGCGTCCT CACCGTCTTGCACCAAGGACTGGCTGAATGGCAAGGAGTAC AAGTGCAAGGTGTCCACAAGCCCTCCAGCCCCCATCG AGAAACCATCTCCAAAGCCAAGGGCAGCCCCGAGAAC CACAGGTGTACACCTGCCCCCATCCCGGAGGAGATGAC CAAGAACCAGGTCAAGCTGACCTGGCTGGTCAAGGCTTC TATCCCAAGGACATCGCCGTGGAGTGGAGAGCAATGGGC AGCCGGAGAACAACACAAAGACCACGCTCCCGTGTGGA CTCCGAGCGCTCTTCTCTCTATAGCAAGCTCACCGTGG ACAAAGACAGGTGGCAGCAGGGGAACGTCTTCTCATGCTC CGTGATGCATGAGGCTCTGCACACCACTACACGCAAG AGCCTCTCCCTGTCTCCGGGTGGTGGCGGATCGGGAAGTG GCGGATCCCAGGTGCAGCTGGTGGAGTCTGGGGGAGGCTT</p>
		<p>N A</p>
	<p>21- 230 36 F3 IgG 7 21 - 233 6F 4 scF v</p>	
iP S: 5 7 7 4 3 7		

	GGTCAAGCCTGGAGGGTCCCTGAGACTCTCTCTGTGCAGCC TCTGGATTCACTTCACTGACTACTACATGAGCTGGATCCG CCAGGCTCCAGGGAAGTGCCTGGA GTGGATTTCATACATT AGTAGCAGTGAAAGTATCATCTATTACGTAGACTCTGTGA AGGCCGATTCACTCATCTCCAGGACAACGCCAAGAACTC ACTGTATCTGCAAAATGAACAGCCTGAGAGCCGAGGACACG GCCGTGTATTACTGTGCGAGAGATGTTGGGAGCCACTTTG ACTACTGGGGCCAGGGAACCTGGTCAACCGTGTCTCAGG AGGGGCGGATCTGGCGGAGGAGGAAAGCGGTGGGGCGG CTCCGACATCCAGATGACCCAGTCTCCATCTTCCGTGCTG CTTCTGCGGAGACAGAGTCAACATCACTTGTGCGGCGAG TCAGGATATTAGCAGGTGGTAGCTGGTATCAGCAGAAA CCAGGGAAGCCCCTAAGCTCCTGATTCTGCTGCATCCA GATTGCAAAAGTGGAGTCCCATCAAGGTTCAAGCGCAGTGG ATCTGGGACAGATTTCACTCTCACCATCAGCAGCCTGCAG CCTGAAGATTTTGCAATTTACTATTGTCAACAGGCTAAAA GTTTCTCGGACGTTCTGGCTGCGGGACCAAGGTGGAAT CAAAACGG	(SEQ ID NO: 335)
		(SEQ ID NO: 336)

		<p>QVQLQQSGPGLVKPSQTLSTCAISGDSVSSRTAWNWIQPSRGLWLGRTYYRSKWYHDYSVSVKSRITIDPDTSKNQFSLQLNSVTPEDTAVYYCARGAAPFDYWGGTLVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQYICNVNHNKPSNTKVDKKVEPKSCDKTHTCPCPAPELLGGPSVFLFPPKPKDTLMISRTPETCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTKISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGGGGGGGGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMSWIRQAPGKCLEWISYISSSEIIYYVDSVKGRTISRDNAKNSLYLQMNSLRAEDTAVYYCARDVGSFHDYWGQGLTVTVSSGGGGGGGGGGGGGGSDIQMTQSPSSVSASVGDRVTITCRASQDISRWLAWYQQKPGKAPKLLISAAASRLQSGVPSRFSGSGGTDFLTITSLQPEDFAIYYCQQAQKSFPRTFGCGTKVEIKR</p> <p>(SEQ ID NO: 337)</p>	<p>EIVLTQSPGTLSLSPGERATLSCRASQSVSSNYLAWYQQKPGQAPRALIYAASNRAAGISDRFSGSGGTDFLTISRLEPEDFAVYFCQQYGSSPLTFGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC</p>	(SEQ ID NO: 338)
A				
A				

		<p>CAGGTGCAGTTGGTGGAGTCTGGGGGAGGCTTAGTCAAGC CTGGAGGTCCCTGAGACTCTCTGTGAGCCCTCTGAATTCT ACCTTCAAGTGAATACTACATGAGCTGGATCCGCCAGGCTC CAGGGAAGGGGCTGGAGTGGTTTCATATATTAGTCGAAG TGGTGATACCATCTACTACGACAGCTCTGTGAAGGGCCGA TTCACCATCTCCAGGGACACGCCAAGAACTCACTGTATC TGCAAAATGAATGGCTGCGAGCCGAAGACACGGCCGTGTA TTACTGTGCGAGAGACTTAGCAGAGGTGTACTACAGGGGCG CTTGACTGCTGGGGCCAGGGAACCTTGGTCAACCGTGTCT CAGCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCC CTCCTCCAAAGACACCTCTGGGGGCACAGGGCCCTGGGC TGCTTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGT CGTGGAACCTCAGGCGCCCTGACCAAGCGCGTGCACACCTT CCCCGCTGTCTACAGTCTCAGGACTCTACTCCCTCAGCA GGCTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCCAGAC CTACATCTGCAACGTGAATCACAAGCCCAAGCAACACCAAG GTGGACAAGAAAGTTGAGCCCAATCTTGTGACAAAACCTC ACACATGCCACCGTGCCACGACCTGAACCTCTGGGGGG ACCGTCAAGTCTTCTCTTCCCCCAAAACCCAAAGGACACC CTCATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGG TGGACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTG GTACGTGGACGGGTGGAGGTGCATAATGCCAAGACAAA GCCGTGCGAGGAGCAGTACGGCAGCACGTACCGTTGCGTC AGCGTCTCACCGTCTCTGCACCAAGGACTGGCTGAATGGCA AGGAGTACAAGTGCAAGGTGTCCAAACAAGCCCTCCCAGC CCCCATCGAGAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAACCACAGGTGTACACCTTGCCCCCATCCCCGGGAGG AGATGACCAAGAACAGGTACGCCGTGACCTGCCTGGTCAA AGGCTTCTATCCAGCGACATCGCCGTGAGTGGGAGAGC AATGGCAGCCGGAGAACAACTACAAGACCAACGCTCCC GTGCTGGACTCCGACGGCTCTTCTTCTCTATAGCAAGCT CACCGTGGACAAGACAGGTGGCAGCAGGGGAACGCTTT CTCATGTCTCCGTGATGCATGAGGCTCTGCACAACCACTAC ACGCAGAAGAGCCCTCTCCCTGTCTCCGGGTGGTGGCGAT CGGAGGTGGCGGATCCCCAGGTGCAGTGGTGGTCTGG</p>
		<p>N A</p>
	<p>21- 230 37 A6 Ig G 21- 233 4 H6 s cFv</p>	
iP S: 5 7 7 4 4 1		

GGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCC TGTGCAGCCTCTGGATTCACTTCAGTGACTACTACATGAC CTGGATCAGGCAAGCTCCAGGGAAGTGCCCTGGAGTGGATT TCATACATTAGTAGTAGTGGTAGTACCATCTACTACGCAG ACTCTGTGAAGGGCCGATTACCATCTCCAGGGACAACGC CAAGAACTCACTGTATCTGCAAAATGAACAGCCTGAGAGCC GAGGACACGGCCGTGTATTACTGTGCGAGAGATCGGAACT CCCACCTTGACTATTGGGGCCAGGGAACCCCTGGTCACCGT GTCCTCAGGAGGGGGGGATCTGGCGGCGGAGGAAAGCGG TGGGGCGGCTCCGACATTTCAGATGACCCAGTCTCCATCT TCCGTGTCTGCATCTGTAGGGGACAGAGTCACCATCACTT GTCGGGCGAGTCAGGGTATTACCAAGTGGTTAGCCTGGTA TCAGCAGAAACCAAGGAAAGCCCCTAAGCTCCTGATCTAT GCTGCATCCGTTTTGC AAAAGTGGGTCCCATCAAGGTTCA GCGGCA GTGGATCTGGGACAGATTCACCTCACCATCAG CAGCCTGCAGCCTGAAGATTTGCAACTTACTATTGTCAAC AGTCTAACAGTTTCCCTCGGACGTTCCGGCTGCGGGACCAA GGTGGA AATCAAACGG	(SEQ ID NO: 339)	(SEQ ID NO: 340)
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		<p>QVQLVESGGGLVKPGGSLRLSCAASEFTFSDYYMSWIRQAP GKGLEWVSYISRGDTIYYADSVKGRFTISRDNAKNSLYLQM NGLRAEDTAVYYCARDLAAGATGGLDCWGQGTLLVTVSSAS TKGPSVFLPAPSSKSTSGGTAAAGCLVKDYFPEPVTVSWNSG ALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNH KPSNTKV/DKKVEPKSCDKHTHCPCPAPELLGGPSVFLFPPKP KDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTPPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGGGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWIR QAPGKCLEWISYSSSGSTIYYADSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDRNSHFDYWGQGTLLVTVSSGGGG SGGGSGGGGSDIQMTQSPSSVSASVGDRTITCRASQGITR WLAWYQQKPGKAPKLLIYAASVLSQGVPSRFSGSGSGTDFT LTISSLQPEDFATYYCQQSNSEFPRTFGCGTKVEIKR</p>	<p>SYELTQPPSVSVSPGQTASITCSGERLGNKYICWYQQKPGQSP VLVIYQDFKRPSPGIPERFSGNSGITAATLTISGTQAMDEADYY CQAWDSRTTVVFGGGTKLTVLGQPKAAPSVTLFPPSSEELQAN KATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSN NKYAASSYLSTPEQWKSHRSYSCQVTHEGSTVEKTVAPTEC</p>
	A		
	A		
			(SEQ ID NO: 341)
			(SEQ ID NO: 342)

		<p>CAGGTGCAGTTGGTGGAGTCTGGGGGAGGCTTAGTCAAGC CTGGAGGTCCCTGAGACTCTCTGTGAGCCCTCTGAATTCT ACCTTAGTGACTACTACATGAGCTGGATCCGCCAGGCTC CAGGGAAGGGGCTGGAGTGGTTTCATATATTAGTCGAAG TGGTGATACCATCTACTACGAGACTCTGTGAAGGGCCGA TTCACCATCTCCAGGGACACGCCAAGAACTCACTGTATC TGCAAAATGAATGGCTGCGAGCCGAAGACAGGCCCTGTGA TTACTGTGCGAGAGACTTAGCAGAGGTGTACTACAGGGGCG CTTGACTGCTGGGGCCAGGGAACCTTGGTCAACCTGTCTCT CAGCTCCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCC CTCCTCCAAGAGCACCTCTGGGGGCACAGGGCCCTGGGC TGCTTGCTCAAGGACTACTTCCCCGAACCGGTGACGGTGT CGTGGAACCTCAGGCGCCCTGACCAGCGGCGTGACACCTT CCCCGCTGTCTACAGTCTCAGGACTCTACTCCCTCAGCA GCGTGTGACCGTGCCCTCCAGCAGCTTGGGCACCCAGAC CTACATCTGCAACGTGAATCACAAGCCCAAGCAACACCAAG GTGGACAAGAAAGTTGAGCCCAATCTTGTGACAAAACCTC ACACATGCCACCGTGCCAGCACCTGAACCTCTGGGGGG ACCGTCAAGTCTTCTCTTCCCCCAAAACCCAAAGGACACC CTCATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGG TGGACGTGAGCCACGAAGACCCCTGAGGTCAAGTTCAACTG GTACGTGGACGGCGTGAGGTGCAATATGCCAAGACAAA GCCGTGCGAGGAGCAGTACGGCAGCACGTACCGTTGCGTC AGCGTCTCACCGTCTCTGCCACCAAGGACTGGCTGAATGGCA AGGAGTACAAGTGCAAGGTGTCCAAACAAAGCCCTCCCAGC CCCCATCGAGAAACCATCTCCAAAGCCAAAGGGCAGCCC CGAACCACAGGTGTACACCTTGCCCCCATCCCCGGGAGG AGATGACCAAGAACAGGTACGCCGTGACCTGCCTGGTCAA AGGCTTCTATCCAGCGACATCGCCGTGAGTGGGAGAGC AATGGCAGCCGGAGAACAACTACAAGACCAACGCTCCC GTGCTGGAATCCGACGGCTCTTCTTCTCTATAGCAAGCT CACCGTGGACAAGAGAGGTGGCAGCAGGGGAACGCTTT CTCATGTCTCCGTGATGCATGAGGCTCTGCACACCACTAC ACGCAAGAGAGCCCTCTCCCTGTCTCCGGGTGGTGGCGAT CGGAGGTGGCGGATCCCCAGGTGACAGTGGTGGAGTCTGG</p>
		<p>N A</p>
	<p>21- 230 37 A6 Ig G 21- 233 6F 4 scF v</p>	
iP S: 5 7 7 4 4 5		

GGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCC TGTGAGCCTCTGGATTCACTTCACTGACTACTACATGAG CTGGATCCGCCAGGCTCCAGGGAAGTGCTGGAGTGGATT TCATACATTAGTAGCAGTGAAGTATCATCTATTACGTAG ACTCTGTGAAGGGCCGATTCAACCATCTCCAGGGACAAACGC CAAGAACTCACTGTATCTGCAAAATGAACAGCCTGAGAGCC GAGGACACGGCCGTGTATTACTGTGCGAGAGATGTTGGGA GCCACTTTGACTACTGGGGCCAGGGAACCTGGTCAACCGT GTCCTCAGGAGGGGGCGGATCTGGCGGCGGAGGAAGCGG TGGGGCGGCTCCGACATCCAGATGACCCAGTCTCCATCT TCCGTGTCTGCTTCTGTCGGAGACAGATCACCATCACTTG TCGGGCGAGTCAGGATATTAGCAGGTGGTTAGCCTGGTAT CAGCAGAAACCAAGGAAAGCCCCCTAAGCTCCTGATTCTG CTGCATCCAGATTGCAAGTGGAGTCCCATCAAGGTTTCAG CGGCAGTGGATCTGGGACAGATTTCACTCTCACCATCAGC AGCCTGCAGCCTGAAGATTTTGCAATTTACTATTGTCAACA GGCTAAAAGTTTCTCTCGGACGTTGCGCTGCGGGACCAAG GTGGAAATCAAAACGG	(SEQ ID NO: 343)	(SEQ ID NO: 344)
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	QVQLVESGGGLVKPGGSLRLSCAASEFTFSDYYMSWIRQAP GKGLEWVSYISRGDTIYYADSVKGRFTISRDNAKNSLYLQM NGLRAEDTAVYYCARDLAAGATGGLDCWGQGTLLVTVSSAS TKGPSVFLPAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSG ALTSGVHTFPALVQSSGLYSLSSVTVTPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSCDKTHTCPCPAPELLGGPSVFLFPPKP KDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNA KTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKCKVSNKA LPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLT DKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGGGGSGG GGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIR QAPGKCLEWISYISSESIYYVDSVKGRFTISRDNAKNSLYLQ MNSLRAEDTAVYYCARDVGSFHDYWGQGTLLVTVSSGGGGS GGGSGGGGSDIQMTQSPSSVSASVGDRVTITCRASQDISRW LAWYQQKPKAPKLLISAA SRLQSGVPSRFSGSGSGTDFTLTI SSLQPEDFAIYYCQQA KSFPRTFGCGTKVEIKR (SEQ ID NO: 345)	SYELTQPPSVSVSPGQTASITCSGERLGNKYICWYQQKPGQSP VLVIYQDFKRPSPGIPERFSGNSGITA TLTISGTQAMDEADYY CQAWDSRTTVFVGGGKLTVLGQPKAAPSVTLFPPSSEELQAN KATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSN NKYAASSYLSTPEQWKSHRSYSCQVTHEGSTVEKTVAPTEC S (SEQ ID NO: 346)
A		
A		

		<p>CAGGTGCAGTGGTGCAGTCTGGGACTGGAGTGAGGTGAAGAAG CCTGGGGCTCAGTGAAGGTGTCTCTGCAAGGCTTCTGGAT ACACTTCCCCGGCTACTATATGCACTGGTGGGACAGGC CCCTGGACAGGGGCTTGAGTGGATGGATGGATCAACCT GACAGTGTGGCACAAAGTATACACAGAAGTTTCAGGGC AGGTACCTTGACAGGGACGCGTCCGTCAGCACAGCCT ACATTGACCTGAACAGGCTGAGATCTGACGACAGGCGCT ATATTACTGTGCGAGAGAGAGGTGTAGGACTACCAACTGC TATTTGGACTACTGGGGCCAGGGAAGTCTGGTCAACCGTG CCTCAGCTCCACCAAGGGCCATCGGTCTTCCCCCTGGC ACCTTCTCCAAAGACACCTCTGGGGCACAGGGCCCTG GGCTGCCGTGGTCAAGGACTACTTCCCCGAACCGGTGACGG TGTCTGTGAACTCAGGCGCCCTGACCAGCGGCGTGACAC CTTCCCGGTGTCTTACAGTCTCAGGACTCTACTCCCTCA GCAGCGTGTGACCGTCCCTCCAGCAGTCTGGGCACCCA GACCTACATCTGCCAACGTGAATCACAAAGCCAGCAACACC AAGGTGACAAAGAAAGTTGAGCCCCAAATCTTGTGACAAA ACTCACACATGCCACCGTGCCACGACCTGAACCTCTGG GGGACCGTCAGTCTTCTCTTCCCCCCTGAGGCAAGGA CACCCTCATGATCTCCGGACCCCTGAGGTACATGCGTG GTGTGGACGTGAGCCACGAAGACCCTGAGGTCAAGTTCA ACTGTAACGTGGACGCGTGGAGGTGCATAATGCCAAGAC AAAGCCGTGCGAGGAGCAGTACGGCAGCAGTACCGTTG CGTACGCGTCTCACCGTCTCTGCCACGAGGACTGGCTGAAT GGCAAGGAGTACAAGTGCAAGGTGTCCAACAAGCCCTC CCAGCCCCATCGAGAAACCATCTCCAAAGCCAAAGGGC AGCCCCGAGAACACAGGTGTACACCTGCCCCCTATCCCG GGAGGAGATGACCAAGAACAGGTGAGCCTGACCTGCCT GGTCAAAGGCTTCTATCCAGCGACATCGCCGTGGAGTGG GAGAGCAATGGCAGCCGGAGAACACTACAAGACCACG CCTCCCGTGTGGACTCCGACGGCTCTTCTTCTCTATAG CAAGCTCACCGTGGACAAAGAGCAGGTGGCAGCGGGGAA CGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACC ACTACACGCAAGAGCCTCTCCCTGTCTCCGGGTGGTG CGGATCGGGAGGTGGCGGATCCCCAGGTGCAGCTGTCGA</p>
		N A
21- 230 39 21 gG 21 - 233 4 H6 4 9 s cFv	iP S: 5 7 7 4 4 9	

	GTCTGGGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGA CTCTCTGTGCAGCCTCTGGATTCACTTCAAGTGAAGTACTA CATGACCTGGATCAGGCAGGCTCCAGGGAAGTGCCTGGAG TGGATTTCATACATTAGTAGTAGTGGTAGTACCATCTACTA CGCAGACTCTGTGAAGGGCCGATTCAACCATCTCCAGGGAC AAGGCCAAGAACTCACTGTATCTGCAAAATGAACAGCCTGA GAGCCGAGGACACGGCCGTGTATTACTGTGGGAGAGATCG GAATCCCACTTTGACTATTGGGGCCAGGGAACCCCTGGTC ACCGTGTCCTCAGGAGGGGGCGGATCTGGCGCGGAGGA AGCGGTGGGGCGGCTCCGACATTCAGATGACCCAGTCTC CATCTCCGTGTCTGCATCTGTAGGGGACAGAGTCAACCAT CACTTGTGGGGAGTCAGGTATTACAGGTGGTTAGCC TGGTATCAGCAGAAACCAAGGAAAGCCCCTAAGCTCCTGA TCTATGCTGCATCCGTTTGCAAAAGTGGGTCCCATCAAG GTTCAAGGCGCAGTGGATCTGGGACAGATTTCACCTCAAC ATCAGCAGCCTGCAGCCTGAAGATTGCAACTTACTATT GTCAACAGTCTAACAGTTCCCTCGGACGTTCCGGCTGCGG GACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 347)
		(SEQ ID NO: 348)

		<p>QVQLVQSGTEVKKPGASVKVCKASGYTPFGYYMHVVRQA PGGLEWMGWINDSGGTYTKYQKFGQGRVTLTRDASVSTAYI DLNRLRSDDTAVYYCARERCRTTNCYLDYW'GQGS'LVTVSS ASTKGPSVFPLAPSSKSTSGGTAAAGCLVKDYFPEPVTVSWN SGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNV NHHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP KPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWY'VDGVEVHN AKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKKVS'NK ALPAPIEKTIISKAKGQPREPQVYVTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFFLYSKLT VDKSRWQQGNVFSCSV'MHEALHNHYTQKSLSLSPGGGGSG GGGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWI RQAPGKCLEWISYISSGSTIYYADSVKGRFTISRDN'AKNSLY LQMNSLRAEDTAVYYCARDNRNSHFDYWGQGTLVTVSSGGG GSGGGGGGGGSDIQMTQSPSSVSASVGDRTVITTCRASQGIT RWLA'WYQQKPGKAPKLLIYAAASVLSQGVPSRFSGSGSGTDF TLTISSLQPEDFATYYCQQSN'SFPRTFGC'GTKVEIKR</p>	<p>QSALTQPAVS'GSPGQSITISCTGTSSDVGNYNLVS'WYQQHP GKAPKLMIEV'NRRRPSGVSNRFSGSKSGNTASLTISGLQAED EA'EYYCCSYAGRDTFV'VFGGGTKLTVLGQPKAAPSVTLFPPS SEELQANKATLVCLISDFYPGAVTV'AWKADSSPVKAGVETT TPSKQSN'NKYAAASSYLSLTPEQWKSHRSYSQVTHEGSTVEK TVAPTECS</p>
	A		(SEQ ID NO: 349)
	A		(SEQ ID NO: 350)

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	GTCTGGGGAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGA CTCTCTGTGCAGCCTCTGGATTCACTTCAAGTGACTACTA CATGAGCTGGATCCGCCAGGCTCCAGGGAAGTGCCTGGAG TGGATTTTCATACATTAGTAGCAGTGAAAGTATCATCTATTA CGTAGACTCTGTGAAGGGCCGATTACCATCTCCAGGGAC AAGCCAAAGAACTCACTGTATCTGCAAAATGAACAGCCTGA GAGCCGAGGACACGGCCGTGTATTACTGTGGGAGAGATGT TGGGAGCCACTTTGACTACTGGGGCCAGGGAACCCCTGGTC ACCGTGTCCTCAGGAGGGGGCGGATCTGGCGGCGGAGGA AGCGGTGGGGCGGC'TCCGACATCCAGATGACCCAGTCTC CATCTTCCGTGTCTTCTGTGCGGAGACAGAGTCACCATC ACTTGTGCGGGGAGTCAGGATATTAGCAGGTGGTTAGCCT GGTATCAGCAGAAACCAAGGAAAGCCCTAAGCTCCTGAT TTCTGCTGCATCCAGATTGCAAGTGGAGTCCCATCAAGG TTCAGCGGCAGTGGATCTGGGACAGATTTCAC'TCTCACCA TCAGCAGCCTGCAGCCTGAAGATTTTGCAATTACTATTGT CAACAGGCTAAAAGTTTTCCTCGGACGTTCCGGTGC'GGGA CCAAAGGTGGAATCAAAACGG	(SEQ ID NO: 351)
		(SEQ ID NO: 352)

		QVQLVQSGTEVKKPGASVKVCKASGYTPFGYYMHVVRQA PGGLEWMGWINPDSGGTKYTQKFGQGRVTLTRDASVSTAYI DLNRLRSDDTAVYYCARERCRTTNCYLDYWQGQSLVTVSS ASTKGPSVFPLAPSSKSTSGGTAAALGCLVKDYFPEPVTVSWN SGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP KPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHN AKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKCKVSNK ALPAPIEKTIISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFFLYSKLT VDKSRWQQGNVFSCSVMEALHNHYTQKSLSLSPGGGGSG GGGSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWI RQAPGKCLEWISYSSSEIYYVDSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDVGSHPDYWGQGTLVTVSSGGGG SGGGSGGGGSDIQMTQSPSSVSASVGDRTITTCRASQDISR WLAWYQQKPGKAPKLLISAAASRLQSGVPSRFSGSGSGTDFTL TISSLQPEDFAIYYCQAKSFPRTFGCGTKVEIKR (SEQ ID NO: 353)	QSALTQPASVSGSPGQSITISCTGTSSDVGNYNLVSWYQQHP GKAPKLMIEVNNRRPSGVSNRFSGSKSGNTASLTISGLQAED EAEYYCCSYAGRDTFVVFGGGTKLTVLGQPKAAPSVTLFPPS SEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETT TPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEK TVAPTECS (SEQ ID NO: 354)
	A		
	A		

Table 25

IgG-scFv v1 Full Sequences

iP S #	T y A b	SCFV_HC	
		LC	

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	CCCTGAGACTCTCTGTGGAGCCTCTGGATTACCTTCAGT GACTACTACATGAGCTGGATCCGCCAGGCTCCAGGGAAGT GCCTGGAGTGGATTTCATACATTAGTAGCAGTGAAAGTATC ATCTATTACGTAGCGCTGTGAAGGCCGATTACCATCTC CAGGGACAACGCCAAGAACTCACTGTATCTGCAAAATGAAC AGCTGAGAGCCGAGGACACGGCCGTGTAATTACTGTCCGA GAGATGTTGGGAGCCACTTTGACTACTGGGGCCAGGGAAC CCTGGTCAACCGTGCTCCAGGAGGGGGCGGATCTGGCGGC GGAGGAAGCGGTGGGGCGGCTCCGACATCCAGATGACCC AGTCTCCATCTTCCGTGCTGCTTCTGTCGGAGACAGAGTC ACCATCACTTGTGGGCGAGTCAGGATATTAGCAGGTGGTT AGCCTGGTATCAGCAGAAACCAAGGAAAGCCCTAAGCTC CTGATTTCTGCTGCATCCAGATTGCAAAAGTGAGTCCCATC AAGGTTCAAGCGGCGAGTGGATCTGGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCCTGAAAGATTTTGCAATTTACTAT TGTCACAGGCTAAAAGTTTTCCTCGGACGTTCCGGCTGCGG GACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 355)
		(SEQ ID NO: 356)

		QVQLQQSGPGLVKPSQTLSTCAISGDSVSSSRTAWNRIQSP SRGLEWLGRTYYRSKWYHDY'SVSVKSRITIDPDT'SKNQFSLQ LNSVTPEDTAVYYCARGAAPFDY'WQGTLVTVSSASTKGPSV FPLAPSSKSTSGGTAAALGCLVKDYFPEPVTVSWNSGALTSGVH TFPAVLQSSGLYSLSVVTVPSSSLGTQTYICNVNHKPSNTKV DKKVEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWY'VDGVEVHNAK'TKPCEEQY GSTYRCVSVLTVLHQDWLNGKEYCKV'SNKA'LPAPIEK'TISK AKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVE A WESNGQPENNYKTTTPVLDSGDFLYSKLTVDKSRWQQGN VFSCVMHEALHNHYTQKSLSPGGGGGGGQVQLVESG GGLVKPGGSLRLSCAASGFTFSDY'YMSWIRQAPGKCLEWISYI SSSEIYYVDAVKGRFTISRDNAKNSLYLQMNSLRAEDTAVY YCARDVGSFHDYWGQGTLVTVSSGGGGGGGGGGGSDIQ MTQSPSSVSASVGDRTVITTCRASQDISRWLA'WYQQKPGKAPK LLISAAASRLQSGVPSRFSGSGGTDFLTISSLQPEDFAIYYCQQ AKSFPRTFGCGTKVEIKR (SEQ ID NO: 357)	QNALTPASVSGSPGQSIITISCTGTSSDVGNYNLVS'WYQQHPG KAPKLMYEVNRRRPSGV'SNRFSGSKSGNTASLTISGLQAEDEA EYYCSSYAGRDTFV'VGGGTKLTVLGGPKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSK QSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTVEKTVAP TECS	(SEQ ID NO: 358)
A				
A				

199

200

	QVQLVQSGAEVTKPGASVKVCKASGYTFAGYYMHWVRQA PGQGLEWMGWINPHSGGTNYAQKFQDRVTMTTRDTSINTAYM ELSLRSDDTAVYYCARERISMVRGVGHNWFAPWGQGTLLVT VSSASTKGPSVFPLA PSSKSTSGGTAAALGCLVKDYFPEPTVS WNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLF PPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVH NAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYCKCKVSNK ALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCVMHEALHNHYTQKSLSLSPGGGGSGGG GSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQA PGKCLEWISYISSSGSTIYYADSVKGRFTISRDNAKNSLYLQM NSLRAEDTAVYYCARDNRNSHFDYWGGGTLLVTVSSGGGGSGG GGSGGGSDIQMTQSPSSVSASVGDRTITTCRASQGTRWLA WYQQKPGKAPKLLIYAASVLSQGVPSRFSGSGGTDFTLTIS LQPEDFATYYCQQNSFPRTFGCGTKVEIKR	DIQMTQSPSSLSASVGDRTITTCRASQDISNNLAWFQQKPGKP PKSLMYAASSLHSGVPSFTFSGSGGTDFTFTISLQPEDFATYY CQQYNSYPLTFGGGKVEIRRTVAAPSVFIPTPSDEQLKSGTAS VVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSST YSLSSITLTLKADYEEKHKVYACEVTHQGLSSPVTKSFNRGEC
A	(SEQ ID NO: 361)	(SEQ ID NO: 362)
A		

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	<p>GAGGCTTGGTCAAGCCTGGAGGGTCCCTGAGACTCTCCTGT GCAGCCTCTGGATTCACTTCAAGTACTACTACATGACCTG GATCAGGCAGGCTCCAGGGAAGTGCCCTGGAGTGGATTTC TACATTAGTAGTGGTAGTACCATCTACTACGCAGAATC TGTGAAGGGCCGATTCAACCATCTCCAGGGACAAGCCAAAG AACTCACTGTATCTGCAATGAACAGCCTGAGAGCCGAGG ACAGGCCCGTGTATTACTGTGCGAGAGATCGGAACCTCCAC TTTGACTATTGGGCCAGGGAACCTGGTCAACCGTGTCTC AGGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGG CGGCTCCGACATTCAAGATGACCCAGTCTCCATCTTCCGTGT CTGCATCTGTAGGGGACAGAGTCAACCATCACTTGTGGGGG AGTCAGGGTATTACCAAGTGGTAGCCTGGTATCAGCAGA AACCAGGGAAGCCCCCTAAGCTCCTGATCTATGCTGCATCC GTTTGCAAAAGTGGGTCCCATCAAGGTTCAGCGGCAGTGG ATCTGGGACAGATTTCACCTCACCATCAGCAGCCTGCAGC CTGAAGATTTTGCAACTTACTATTGTCAACAGTCTAACAGT TTCCCTCGGACGTTGGCTGGGGACCAAGGTGGAATCA AACGG</p>	(SEQ ID NO: 363)
		(SEQ ID NO: 364)

		QVQLVQSGAEVTKPGASVKVCKASGYTFAGYYMHWVRQA PGQGLEWMGWINPHSGGTNYAQKFQDRVTMTTRDTSINTAYM ELSLRSDDTAVYYCAREISMVRGVGHNWFAPWGQGTLLVT VSSASTKGPSVFPLA PSSKSTSGGT AALGCLVKDYFPEPTVS WNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLF PPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVH NAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNK ALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCVMHEALHNHYTQKSLSLSPGGGGSGGG GSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMTWIRQA PGKCLEWISYISSSGSTIYYAESVKGRFTISRDNAKNSLYLQMN SLRAEDTAVYYCARDRNSHFDYWGGGTLLVTVSSGGGGSGGG SGGGGSDIQMTQSPSSVSASVGDRTVTITCRASQGTRWLAW YQKPGKAPKLLIYAASVLQSGVPSRFSGSGSGTDFLTISLQ PEDFATYYCQQNSFPRTFGCGTKVEIKR	DIQMTQSPSSLSASVGDRTVTITCRASQDISNNLAWFQQKPGKP PKSLMYAASSLHSGVPSFTFSGSGGTDFFTTISLQPEDFATYY CQQYNSYPLTFGGGTKEIRRTVAAPSVFIPTPSDEQLKSGTAS VVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSST YSLSSITLTLKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
A		(SEQ ID NO: 365)	(SEQ ID NO: 366)
A			

		GAGGTGCAGCTGCTGGAGTCTGGGGAGGGTTGGTACAGC CTGGGGTCCCTGAGACTCTCTCTGTGCAGCCTCTGGATTCT ACCTTTAGTAAATGCCATGAGTTGGTCCGCCAGGTCC AGGGAAGGGGCTGGAGTGGGTGCAGCTACTGGTGGTAGT GGTATTAGCACATACTACGACACTCCGTGAAGGGCCGGTT CACCATCTCCAGAGACAATTCCAAGAACACAGCTGTATCTGC AAATGAACAGTCTGAGAGCCGAGGACACGGCGTATATTA CTGTGCGAGAGGTTATAGCAACAGCTGGTGGCTATTGACT ACTGGGGCCAGGGAACCTGGTCAACCGTGTCTTCAAGCCTCC ACCAAGGGCCCATCGGTCTTCCCCCTGGCACCTCTCTCCAA GAGCACCTCTGGGGGACACAGCGGCTTGGGCTGCCCTGGTC AAGGACTACTTCCCGAACCGGTGACGGTGTCTGTGGAACCTC AGGCGCCTGACCAGCGGCGTGACACCTTCCC GGCTGTCC TACAGTCTCAGGACTCTACTCCCTCAGCAGCGTGGTGACC GTGCCCTCCAGCAGCTTGGGCACCCAGACCTACATCTGCAA CGTGAATCACAAAGCCAGCAACACCAAGGTGGACAAAGAA GTTGAGCCCCAAATCTTGTGACAAAACTCACACATGCCAC GTGCCAGCACCTGAACCTCTGGGGGACCGTCAAGTCTTCC TCTTCCCCCAAAACCCAGGACACCTCTATGATCTCCCCG ACCCCTGAGGTACATCGGTGGTGGTGGACGTGAGCCACG AAGACCTGAGGTCAAGTCAACTGGTACGTGGACGGCGT GGAGGTGCATAATGCCAAGACAAAGCCGTGCGAGGAGCAG TACGGCAGCACGTACCGTTGCGTCAAGCTCTCACCGTCTCT GCACAGGACTGGCTGAATGGCAAGGAGTACAAGTGCAAG GTGTCCAAACAAAGCCCTCCAGCCCCCATCGAATAACCA TCTCCAAAGCCAAAGGGCAGCCCCGAGAACACAGGTGTA CACCTGCCCCCATCCCGGAGGAGATGACCAAGAACCAG GTCAGCCTGACCTGCCCTGGTCAAAGGCTTCTATCCCAGCGA CATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAAC AACTACAAAGACCAGCCTCCCGTGTGGACTCCGACGGCTC CTTCTCTCTATAGCAAGCTCACCGTGGACAAAGACAGGT GGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCATGAG GCTCTGCACAAACCACTACACGCAGAAAGCCCTCTCCCTGTC TCCGGGTGGTGGCGGATCGGGAGGTGGCGGATCCCAAGGTG CAGCTGGTGGAGTCTGGGGGAGGGTTGGTCAAGCCTGGAG		
		N A		
		S S T		
iP	S:	6 1 2 5 0 1	2 0 6 8 2	

	GGTCCCTGAGACTCTCCTGTGCGAGCCTCTGGATTACCTTC AGTGACTACTACATGAGCTGGATCCGCCAGGCTCCAGGGA AGTGCTGGAGTGGATTTCATACATTAGTAGCAGTGAAAGT ATCATCTATTACGTAGACGCTGTGAAGGCCGATTCAACCAT CTCCAGGGACAACGCCAAGAACTCACTGTATCTGCAAAATG AACAGCCTGAGAGCCGAGGACACGGCCGTGTATTACTGTG CGAGAGATGTTGGGAGCCACTTTGACTACTGGGGCCAGGG AACCCTGGTCAACCGTGTCTCAGGAGGGGGCGGATCTGGC GGCGAGGAAGCGGTGGGGCGGCTCCGACATCCAGATGA CCCAGTCTCCATCTTCCGTGCTGCTTCTGTCGGAGACAGA GTCACCATCACTTGTGGGGCGAGTCAGGATATTAGCAGGTG GTTAGCCTGGTATCAGCAGAAACCAAGGAAAGCCCCTAAG CTCCTGATTTCGTGTCATCCAGATTGCAAAAGTGGAGTCCC ATCAAAGGTTCAAGCGGCGAGTGGATCTGGACAGATTTCACTC TCACCATCAGCAGCCTGCAAGCTGAAGATTGTGCAATTAC TATTGTCAACAGGCTAAAAGTTTTCCTCGGACGTTTCGGCTG CGGACCAAGGTGGAATCAACGG	(SEQ ID NO: 367)
		(SEQ ID NO: 368)

		EVQLLESGGGLVQPGGSLRLSCAASGFTFSRNAMSWVRQAPG KGLEWVSATGGGISTYYADSVKGRFTISRDNKNTLYLQMN SLRAEDTAVYYCARGYSNSWWYFDYWGQGTLLVTVSSASTK GPSVFPLAPSSKSTSGGTAAALGCLVKDYFPEPVTVSWNSGALT SGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPS NTKVDKKVEPKSCDKTHCTCPAPELLGGPSVFLFPPKPKDT LMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSD IAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMHEALHNHYTQKSLSPGGGGGGGGSQVQL VESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPGKCLE WISYISSESIYYVDAVKGRFTISRDNKNSLYLQMNSLRAED TAVYYCARDVGSFHDYWGQGTLLVTVSSGGGGGGGGGGG GSDIQMTQSPSSVSASVGDRTVITCRASQDISRWLAWYQQKP GKAPKLLISAASRLQSGVPSRFSGSGGTDFTLTISSLQPEDFAI YYCQQAKSFPRTFGCGTKVEIKR	QSALETQPASVSGSPGQSIITISCTGTSSDVGNYNLVSWYQQHPG KAPKLMIFEVNRPSGVSNRFSGSKSGTTASLTISGLQAADAE/ DYFCSSYTTSTYVIFGGGKLTVLGQPKAAPSVTLFPPSSEEL QANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQ SNNKYAASSYLSTLTPQWKSRSYSCQVTHEGSTVEKTVAPT ECS
A	A	(SEQ ID NO: 369)	(SEQ ID NO: 370)

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		QVQLVQSGAEVKPGASVKVCKASGYTPGYMYWLRQA PGQGLEWMGWINDSGDTNYAQKFQGRVTMTRDTSISTAFM ELSLRSDDTAVYYCAREKPRYFDSFYLLMDVWGQGTITV VSSASTKGPSVFPLAPSSKSTSGGTAAALGCLVKDYFPEPTVS WNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLF PPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVH NAKTKPCEEQYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNK ALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLV KGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTV DKSRWQQGNVFSCVMHEALHNHYTQKSLSPGGGGGGGG GSQVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQA PGKCLEWISYSSSESHYYVDAVKGRTISRDNAKNSLYLQMN SLRAEDTAVYYCARDVGSHFDYWGQGLTVTVSSGGGGGGGG GSGGGGDIQMTQSPSSVSASVGDRTITTCRASQDISRWLA YQKPGKAPKLLISAAASRLQSGVPSRFSGSGGTDFTLTISLQ PEDFAIYYCQQAQKSFPRTFGCGTKVEIKR	DIQMTQSPSSLSASVGDRTITTCRAGQNIAHLNHWYQQNP GK APKVLIHPPASSLP SGVPSRFSGSGGTD FSLTISLQPEDFGTYF CQQSYSTPPTFGGGTKVELKRTVAAPSVFIIPPSEQLKSGTAS VVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSST YSLSSITLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC
A		(SEQ ID NO: 373)	(SEQ ID NO: 374)
A			

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	CCCTGAGACTCTCTGTGGAGCCTCTGGATTACCTTCAGT GACTACTACATGAGCTGGATCCGCCAGGCTCCAGGGAAGT GCCTGGAGTGGATTTCATACATTAGTAGCAGTGAAAGTATC ATCTATTACGTAGCGCTGTGAAGGCCGATTACCATCTC CAGGGACAACGCCAAGAACTCACTGTATCTGCAAAATGAAC AGCCTGAGAGCCGAGGACACGGCCGTGTATTACTGTGCCGA GAGATGTTGGGAGCCACTTTGACTACTGGGGCCAGGGAAC CCTGGTCACCGTGTCTCAGGAGGGGGCGGATCTGGCGGC GGAGGAAGCGGTGGGGCGGCTCCGACATCCAGATGACCC AGTCTCCATCTTCCGTGTCTGTCTGTCTGGGAGACAGAGTC ACCATCACTTGTCTGGGCGAGTCAGGATATTAGCAGGTGGTT AGCCTGGTATCAGCAGAAACCAAGGAAAGCCCTAAGCTC CTGATTTCTGCTGCATCCAGATTGC AAAAGTGGAGTCCCATC AAGGTTCAAGCGGCAGTGGATCTGGGACAGATTTCACCTCA CCATCAGCAGCCTGCAGCCTGAAAGATTTTGCAATTTACTAT TGTCACAGGCTAAAAGTTTTCCTCGGACGTTCTGGGTGCGG GACCAAGGTGGAAATCAAAACGG	(SEQ ID NO: 375)
		(SEQ ID NO: 376)

	QVQLVESGGGVVQPGRLRLSCAASGFTLSSYGMHWVRQAP GKGLEWVAIVYDGSNKYYADSVKGRVTISRDNKNTLYLQ MNSLRAEDTAVYYCTRDGRNYYVFDNWGGQTLVTVSSASTK GPSVFPLAPSSKSTSGGTAAALGCLVKDYFPEPVTVSWNSGALT SGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPS NTKVDKKVEPKSCDKTHCTCPCPAPELLGGPSVFLFPPKPKDT LMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGKEYCKVSNKALPAPIE KTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSD IAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMHEALHNHYTQKSLSLSPGGGGGGGGGSQVQL VESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPGKCLE WISYISSESIYYVDAVKGRFTISRDNKNSLYLQMNSLRAED TAVYYCARDVGSHFDYWGGQTLVTVSSGGGGGGGGGGGG GSDIQMTQSPSSVSASVGDRTVITTCRASQDISRWLAWYQQKP GKAPKLLISAASRLQSGVPSRFSGSGGTDFTLTISSLQPEDFAI YYCQQAQSFPRTFGCGTKVEIKR	(SEQ ID NO: 377)
A		(SEQ ID NO: 378)

Table 26: MSLN-CD40 IgG-scFv Full length

21- 233 4 H6 1 gG 7 7 3 0 0 G7 sc Fv	iP S: 5: 7 7 3 0 0	<p>CAGGTGACAGTGGTGCAGTCTCGGGGAGGCTTGGTCAAGC CTGAGGGTCCCTGAGACTCTCTGTGCGAGCCTCTGGATTCT ACCTTCAGTGACTACTACATGACCTGGATCAGGCAGGCTC CAGGGAAGGGGCTGGAGTGGATTTCATACATTAGTAGTAG TGGTAGTACCATCTACTACGAGACTCTGTGAAGGGCCGA TTCCACCATCTCCAGGGACACGCCAAGAACTCACTGTATCT GCAATGAACACAGGCTGAGAGCCGAGGACACGGCCCTGTAT TACTGTGCGAGAGATCGGAACCTCCCACTTGACTATTGGG GCCAGGGAACCCCTGGTCAACGCTGCTCAAGCTCCACCAA GGGCCCATCGGTCTCCCCCTGGCACCCCTCTCCCAAGAGCA CCTCTGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAAGGA CTACTTCCCCGAACCGGTGACGGTGTCTGTGGAACCTCAGGC GCCCTGACCAAGCGGCTGCACACCTTCCCCGGCTGTCTACA GTCTCAGGACTTACTCCCTCAGCAGCGTGGTGACCCGTGC CCTCCAGCAGCTTGGGCAACCCAGACCTACATCTGCAACGT GAATCAACAAGCCCAAGCAACCAAGGTGGACACAAGAAAGT TGAGCCCCAAATCTTGTGACAAAACCTCACACATGCCCAACG TGCCAGCACCTGAACCTCTCGGGGGACCGTCAGTCTTCCT CTTCCCCCAAAAACCAAGGACACCCCTCATGATCTCCCGG ACCCCTGAGGTACATGCGTGGTGGTGGACGTGAGCCACG AAGACCTGAGGTCAAGTTCAACTGTAAGTACGTGGACGGCGT GGAGTGCATAAATGCCAAGACAAAAGCCGTGCGAGGAGCA GTACGGCAGCACGTACCGTTGCGTCAAGCTCTCAACGTCC TGCAACCAAGACTGGCTGAATGGCAAGGAGTACAAGTGCAA GGTGTCACAACAAGCCCTCCCAAGCCCCATCGAAGAAACC ATCTCAAAGCCAAAGGGCAGCCCCGAGAACCCACAGGTGT ACACCTGCCCCCATCCCGGGAGGAGATGACCAAGAAACCA GGTCAGCCTGACCTGGCTGGTCAAGGCTTCTATCCCCAGCG ACATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGA ACAACTACAAGACCAACGCTCCCGTGTCTGGACTCCGACCG CTCCTTCTCTCTATAGCAAGCTCAACCGTGGACAAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTCATGCTCCGTGATGCA TGAGGCTCTGCACAACCACTACACGCAAGAAAGCCCTCC CTGTCTCCGGGTGCTGGGATCGGGAGGTGCGCGATCCC</p>	<p>GACATTACAGATGACCCAGTCTCCCATCTTCCGTGTCTGCATC TGTAGGGGACAGAGTCAACCATCTTGTGCGGGCGAGTCAAG GGTATTACCAAGGTGGTACCTGGTATCAAGCAAAACCCAG GGAAAGCCCCTAAGCTCTGATCTATGCTGCATCCGTTTGTG CAAAGTGGGTCCCATCAAGGTTCAAGCGCAGTGGATCTG GGACAGATTTCACTCTCAACCATCAGCAGCCTGCGAGCCTGA AGATTTTGCAACTTACTATTGTCAACAGTCTAACAGTTTCC CTCGGACGTTTCGGCCAAGGACCAAGGTGGAAATCAAAACG GACGGTGGCTGCACCATCTGTCTTCACTTCCCCGCCATCTG ATGAGCAGTTGAAATCTGGAACCTGCCCTCTGTTGTGTGCCGTG CTGAATAACTTCTATCCCCAGAGAGGCCAAAGTACAGTGGGA AGGTGGATAACGCCCTCCAATCGGGTAACCTCCCAGGAGAG TGTCACAGAGCAGGACAGCAAGGACAGCACTACAGCCTC AGCAGCACCTTGACGCTGAGCAAAAGCAGACTACGAGAAA CACAAAGTCTACGCTGCGAAGTCAACCATCAGGGCCTGA GCTCGCCCGTCACAAAAGAGCTTCAACAGGGGAGAGTGT</p>
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	AGGTGCAGCTGGTGCAGTCTGGGGCTGAGGTGAAGAAGCC TGGGCTCAGTGAAGGTGTCCTGCAAGGCTTCTGGATAC ACCTCGCCGGCTACTATATGCACTGGGTGCGACAGGCC CTGGACAATGCCCTTGAGTGGATGGATGGATCAACCTGA CAGTGGAGGCACAACTTTGCACAGCAGTTTCAGGGCAGG GTCACATGACCAAGGATACGTCCATCAGCACACAGCCTACA TGGAGGTGAGCAGGCTGAGATCTGACGACAGGGCCGTGTT TTACTGTGCGAGAGAGAGATCACTATGACTGGTATTTACT TTGACTATTGGGGCCAGGGAACCTGGTCACCGTGTCCCTCA GGAGGGCGGATCTGGCGCGGAGGAAGCGGTGGGGC GGTCCGACATCCAGATGACCCAGTCTCCATCCTCCCTGTC TGCACTGTAGGAGACATTTCTACCATCACTTGCCGGGCAA GTCAGAACATTACCACTATTAAATTGGTATCAGCAGAA ACCAAGGAAAGCCCTAACCTCCTGATCTCTGCTGCATCCC GTTTGCAGAGTGGGTCCCATCAAGGTTCAAGTGGCAGTGG ATCTGGGACAGATTTCACTCTCACCATCAGCAGTCTGCAAC CTGTAGATTTTACAACCTTCTACTGTCAACAGACTTTCAC ACCCCGTGGACGTTGGGCTGCGGGACCAAGGTGGAGATCA AACGA	(SEQ ID NO: 379)
		(SEQ ID NO: 380)

		QVQLVESGGGLV ^K PGGSLRLSCAASGFTFS ^D Y ^M MTWIRQAP GKLEWISY ^I SSSGSTI ^Y Y ^A DSV ^K GRFTISRDN ^A AKNSL ^Y LQMN SLRAEDTAV ^Y YCARD ^R NSHFD ^Y W ^G QGTLVTVSSAST ^K GPSV FPLAPSSK ^S TSGGTAA ^L GCLV ^K DYFPEP ^V TVSWNSGAL ^T SGV HTFPAVLQSSGL ^Y SLSSV ^V TV ^P SSSLGTQ ^T YICNVN ^H KPSNTK VDKKVEPKSCD ^K TH ^T CP ^C PAPELLG ^G PSVFLFP ^P PKPKD ^T LM SRTPEV ^T C ^V V ^V DVSHEDPE ^V KFNW ^Y V ^D GV ^E VHNA ^K TKPCEE QY ^G STYRCV ^S VL ^T VLHQD ^W LN ^G KEY ^K CKV ^S NKAL ^P APIE ^K TI SKAKGQPREPQ ^V Y ^T LPPSREEM ^T KNQ ^V SL ^T CLV ^K GFYPS ^D IA A A VEWESNGOPEN ^N YK ^T TP ^P VLDS ^D GSF ^F LY ^S SKL ^T VDKSRW ^Q Q GNV ^F SCSV ^M HEAL ^H NHY ^T Q ^K SL ^S LSPGGGGGGG ^S QV ^Q LV QSGAEV ^K KPGAS ^V K ^V SCKAS ^G Y ^T FA ^G Y ^Y MHW ^V RQAP ^G QCL EWMGW ^I NPDSGG ^T NFA ^Q QFQ ^R VT ^M TRD ^T SISTAY ^M EV ^S RL RSDDTAV ^F YCARE ^K IT ^M TG ^I YFD ^Y W ^G QGT ^L VTVSSGGG ^S G GGSGGGGSDIQ ^M TQSPSS ^L SA ^S VGDIL ^T ITCRASQ ^N ITTY ^L N WY ^Q QKPGKAP ^N LLISA ^A SLR ^L SGV ^P SRFSGSGSGTD ^F TLTISSL QPVDFTTFYCQ ^Q Q ^T FT ^T PT ^P WTFGCGTK ^V EIKR (SEQ ID NO: 381)	DIQMTQSPSSV ^S ASV ^G DRV ^T ITCRASQGITR ^W LAWY ^Q QKPGK APKLLI ^Y AASV ^L QSGV ^P SRFSGSGSGTD ^F TLTISSLQPED ^F ATY YCQ ^Q SN ^S FPRT ^F GQGT ^K VEIK ^R TVAA ^P SVFI ^F PPSDEQL ^K SGT ASV ^V CLLN ^N FYP ^R EAK ^V QW ^K VDN ^A LQSGNSQ ^E SVTEQ ^D SKD STYSL ^S STL ^T L ^S KADY ^E KE ^H KV ^Y ACEV ^T HQGLSSP ^V TKS ^F NRG EC (SEQ ID NO: 382)
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	TGGGGCCTCAGTGAAAGGTGTCTGCAAGGCTTCTGGATAC ACCTTCGCCGGCTACTATATGCACTGGGTGCGACAGGCCC CTGGACAAATGCCCTTGAGTGGATGGATGGATCAACCCCTCA CAGTGGTGGCACAAACTATGCACAGAAATTTCAGGACAGG GTCACCATGACCAAGGACACGTCCATCAACACAGCCTACA TGGAACCTGAGCAGGCTGAGATCTGACGACACGGCCGTGTA TTACTGTGCGAGAGAACGTATTTCTATGGTTCTGGGGAGTCG GGCACAACTGGTTCGCCCCCTGGGGCCAGGAAACCCCTGGT CACCGTGTCTCAGGAGGGGGCGGATCTGGCGGGAGGA AGCGGTGGGGGGCTCCGACATCCAGATGACCCAGTCTC CATCCTCACTGTCTGCATCTGTAGGAGACAGAGTCACCATC ACCTGTGGGGCAGTCAGGACATTAGCAATAATTAGCCT GGTTTCAGCAGAAACCAGGGAACCCCTAAGTCCCTGAT GTATGCTGCATCCAGTTTGACACAGTGGAGTCCCATCAACGT TCAGCGGCAGTGGATCTGGGACAGATTTCACTTTCACCATC AGCAGCCTGCAGCCTGAAGATTGTGCAACTTATTACTGCCA ACAGTATAATAGTTACCTCTCACTTTCGGCTGGGGACCA AGGTGGAGATCAGACGA	(SEQ ID NO: 383)
		(SEQ ID NO: 384)

		QVQLVESGGGLV ^K PGGSLRLSCAASGFTFS ^{DY} YMTWIRQAP GKLEWISY ^{IS} SSG ^{STI} Y ^Y ADSVKGRFTISRDN ^{AK} NSLYLQMIN SLRAEDTA ^{VY} YCARD ^{RNS} HFD ^Y WGQGT ^L VT ^V SSAST ^K GPSV FPLAPSSK ST SGGT ^A ALGCLV ^K DYFPEP ^{VT} VS ^{WN} SGAL ^T SGV HTFPAVLQSSGL ^Y SLSSV ^{VT} VPSSSLGT ^Q TYICNVN ^{HK} PSNTK VDKKVEPKSCDK TH TCPPCPAPELLGGPSVFLPPKPKD ^{TL} MI SRTP ^{EV} TCV ^V VDVSHEDPEVKFNW ^Y VDGVEVHN ^{AK} TKPCEE QYGSTYRCV ^{SV} LV ^L HQDWLNGKEY ^K CKVSNKALPAPIEK ^{TI} SKAKGQPREPQV ^Y TLPPSREEMTKNQ ^{VS} L ^T CLVKGFYPSDIA A A VEWESNGQPENNYK ^T TPPVLDSDGSFFLY ^{SK} L ^T VDKSRWQQ GNVFSCSV ^M HEALHNHY ^T QKSLSLSPGGGGGGG ^Q SV ^{QL} V QSGAEV ^{TK} PGASV ^K V ^{SK} ASGY ^{TF} AGY ^Y MHW ^{VR} QAPGQCL EWMGWINPHSGGT ^{NY} AQKFQDR ^{VT} MT RD TSINTAYMEL ^{SRL} RSDDTA ^{VY} YCARERISM ^{VR} GV ^{GH} NWFA ^{PW} GQGTL ^{VT} VSSG GGSGGGGGGGGSDIQ ^{MT} QSPSSL ^{SA} SVGDR ^{VT} ITCRASQDI SNNLAWFQ ^Q KPKPK ^{SL} MYAASSLHSGV ^{PS} TFSGSGSG ^{TD} F TFTISSLQPEDFAT ^Y YCCQ ^Y NSY ^{PL} TFGCGTK ^{VE} IRR (SEQ ID NO: 385)
		DIQMTQSPSSVSASVGDR ^{VT} ITCRASQGITR ^{WL} AWYQQK ^{PGK} APKLLI ^Y AASV ^L QSGV ^{PS} RFSGSGGT ^{DF} TLTISSLQPEDFAT ^Y YCQQSN ^{SP} RTFGQGT ^K VEIK ^{RT} V ^A APSVFI ^{FP} PSDEQL ^K SGT ASVVCLLN ^{NF} YPREAK ^{VQ} WKVD ^{NAL} QSGNSQES ^{VT} EQDS ^{KD} STYSLSSL ^{TL} SKADY ^E KEHK ^{VY} ACEV TH QGLSSP ^{VT} TKSF ^{NRG} EC (SEQ ID NO: 386)

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	TGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCA CCTTAGTAGAAATGCCATGAGTTGGTCCGCCAGGCTCC AGGAAAGTGCCTGGAGTGGGTGCAGCTACTGGTGGTAGT GGTATTAGCACATACTACGCAGACTCCGTGAAGGGCCGGT TCACCATCTCCAGAGACAATTCCAAGAACACGCTGTATCT GCAAATGAACAGTCTGAGAGCCGAGGACACGGCCGTATAT TACTGTGCGAGAGGTTATAGCAACAGCTGGTGGTACTTTG ACTACTGGGCCAGGGAACCTGGTCACCGTGTCTCTCAGG AGGGGCGGATCTGGCGGCGGAGGAAGCGGTGGGGCGG CTCCAGTCTGCCCTGACTCAGCCTGCCCTCCCTGCTGGGA GCCCTGGACAGTCGATCACCATCTCCTGCACTGGAACCCAG CAGTGATGTTGGGAATTATAACCTTGTCTCCTGGTACCAAC AGCACCCAGGCAAGCCCCCAAACTCATGATTTTGAGGT CAATCAGCGGCCCTCAGGGGTTTCTAATCGCTTCTCTGGCT CCAACTCTGGCACCAAGGCCCTCCCTGACAACTCTCTGGGCTC CAGGCTGCGGACGAGGCTGATTATTTCTGCTGCTCATATAC AACTAGTAGCACTTATGTGATCTTCGGCTGCGGGACCAAG CTGACCGTCTCTAGGT	(SEQ ID NO: 387)
		(SEQ ID NO: 388)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQAP GKLEWISYISSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGGQTLVTVSSASTKGPSV FPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHKPSNTK VDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDITLMI SRTPETCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA A A VEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMHREALHNHYTQKSLSLSPGGGGGGGGSEVQLL ESGGGLVQPGGSLRLSCAASGFTFSRNAMSWVRQAPGKCLE WVSATGGSGISTYYADSVKGRFTISRDNKNTLYLQMNSLRA EDTAVYYCARGYSNSWWYFDYWGGQTLVTVSSGGGGSGG GGSGGGGSQSALTQPASVSGSPGQSITISCTGTSSDVGNYNLV SWYQQHPGKAPKLMIFEVNRQPSGVSNRFGSKSGTTASLTIS GLQAADEADYFCCSYTTSSTYVIFGCGTKLTVLG (SEQ ID NO: 389)	DIQMTQSPSSVSASVGDRVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLAQSGVPSRFSGSGGTDFTLTISSLQPEDFATY YCQQSNSEPRFTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWVKVDNALQSGNSQESVTEQDSKD STYSLSSITLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRG EC (SEQ ID NO: 390)
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	TGGGGCCTCAGTGAAAGGTGTCCTGCAAGGCTTCTGGATAC ACCTTCCCGGCTACTATATGTACTGGTTGCGACAGGCCCC TGGACAATGCCCTTGAGTGGATGGATGGATCAACCCCTGAC AGTGGTGACACAACTATGCACAGAA GTTTCAGGGCAGGG TCACCATGACCAGGACACGTCCATCAGCACAGCCTTTAT GGAGCTGAGCAGGCTGAGATCAGACGACACGGCCGTGTAT TACTGTGCGAGAGAGAGCCAGATATTTGACTCCTTCTA CTACTACCTTATGGACGTCTGGGCCAAGGACCCACGGTC ACCGTGTCTCAGAGGGGGCGGATCTGGCGCGGAGGAA GCGGTGGGGCGGCTCCGACATCCAGATGACCCAGTCTCC ATCCTCCCTGTCTGCATCTGTAGGAGACAGAGTCACCATCA CTTGCCGGCAGGTCAAGAACATTAGCAGGCATTTAAATTG GTATCAGCAGAAATCCAGGGAAGCCCCCTAAGGTCTGATC CATCCTGCATCCAGTTTGCCAAAGTGGGGTCCCGTCAAGGTT CAGTGGCAGTGGATCTGGGACAGATTTCACTCTTACCATC AGCAGTCTGCAACCTGAAGATTTTGGAACTTACTTCTGTCA ACAGAGTTACAGTACCCCTCCCCTTTCTGGCTGCGGGACC AAGGTGGAGCTCAAACGA	(SEQ ID NO: 391)
		(SEQ ID NO: 392)

		QVQLVESGGGLV ^K PGGSLRLSCAASGFTFS ^D Y ^M MTWIRQAP GKLEWISY ^I SSSGSTI ^Y Y ^A DSV ^K GRFTISRDN ^A AKNS ^L YLQ ^M N SLRAEDTAV ^Y YCARD ^R NSHFD ^Y WGQ ^G TLVT ^V SSAST ^K GPS ^V FPLAPSSK ^S TSGGTAA ^L GCLV ^K DYFPEP ^V TVSW ^N SGAL ^T SGV HTFPAVLQSSGL ^Y SLSSV ^V TV ^P SSSLGTQ ^T YICN ^V NH ^K PS ^N TK VDKKVEPKSCDK ^T HTC ^P CPAPELLGGPSV ^F LP ^P PK ^D TL ^M I SRTP ^E VT ^C V ^V VDVSHEDPE ^V KFNW ^Y VDGVEVHN ^A KTK ^P CEE QYGSTYRCV ^S VL ^T VLHQDWLNGKEY ^K CKV ^S NKALPAPIEK ^T I SKAKGQPREPQ ^V Y ^T LPPSREEMTKNQ ^V SL ^T CLVKGFYPS ^D IA A A VEWESNGQPENNY ^K TPPVLDSDGSF ^L Y ^S KL ^T VDKSRW ^Q Q GNVFSCSV ^M HEALHNH ^Y TQKSL ^S SPGGGGGGG ^S QV ^Q LV QSGAEV ^K KPGAS ^V KV ^S CKASGY ^T FP ^G Y ^M Y ^W LRQAPG ^Q CL EWMGWINPDSG ^D TNYAQKFQGR ^V TMT ^R DT ^S ISTAFMEL ^S RL RSDDTAV ^Y YCAREKPRY ^F DSFY ^Y YLMDV ^W GQGTT ^V TVSS ^G GGSGGGGGGGGSDIQ ^M MTQSP ^S SL ^S ASVGD ^R VTITCRAG ^Q N ISRHLNW ^Y QQNP ^G KAP ^K VLIHPASS ^L PSGV ^P SRFSGSGSG ^T DF SLTISSLQPEDFG ^T YFCQQQ ^S YSTP ^T PTFGCGTK ^V EL ^K R (SEQ ID NO: 393)	DIQMTQSP ^S SV ^S ASVGD ^R VTITCRASQGIT ^R WLAW ^Y QQK ^P GK APKLLI ^Y AASV ^L QSGV ^P SRFSGSGGT ^D FTLT ^S SLQPED ^F AT ^Y YCQQSN ^S FPRTFGQ ^G TKV ^E IKRT ^V AAPSVFI ^F PPSDEQL ^K SG ^T ASVVCLLN ^N FYP ^R EAKVQ ^W KVDN ^A LQSGNSQ ^E SVTEQ ^D SK ^D STYSL ^S STLT ^L SKADY ^E E ^K HK ^V YACEV ^T HQGLSSP ^V TKS ^F NR ^G EC	(SEQ ID NO: 394)
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	TGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCA CCTTCAGTAGCCATGGCATGCACTGGGTCCGCCAACCTCCA GGCAAGTGCCTGGAGTGGGTGGCAGTTATCTGGTATGATG GAAGTAATGAATACTATGGAGACTCCGTGAAGGCCGATT CACCATCTCAGAGACAATTCCAAGAACACGCTGTATCTG CAAATGAACAGCCTGAGAGTCGAGGACACGGCTGTGTATT ACTGTACGAGAGGGGGGGCCACTGGAACTACGAGGGCC ACTACTATGGTATGGACGTCTGGGGCCAAGGACCAACGGT CACCGTGTCTCAGGAGGGGGCGGATCTGGCGGCGGAGGA AGCGTGGGGGGCTCCAGGCTGTGCCGACTCAGCCCT CTCCCTCTCTGCATCTCCTGGAGCATCAGCCAGTCTCACC TGCACCTTACGAGTGGCATCAATGTTGGTTCTCCAGGAT CTATTGGTACCAGCAGAAAGCCAGGAGTCTCCCCAGTTT CTCCTGAGGTACACATCAGACTCAGATAAAATTGCAGGGCT CTGGAGTCCCCAGCCGCTTCTCTGGATCCAAAGATGCTTCG GCCAATGCAGGACTTTTACTCATCTCTGGGCTCCAGTCTGA GGATGAGGCTGACTATTACTGTATGATTGGCACAGCAGC GCTGTGGTATTCTGGCTGCGGGACCAAACTGACCGTCTCTAG GT	(SEQ ID NO: 395)	(SEQ ID NO: 396)

		<p>QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQAP GKGLEWISYISSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGGQTLVTVSSASTKGPSV FPLAPSSKSTSGGT AALGCLVKDYFPEPTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHHKPSNTK VDKKVEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDITLMI SRTPEVTCVVVDVSHEDPEVKFNWYVVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA VEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMHREALHNHYTQKSLSLSPGGGGGGGQVQLV ESGGGVVQPGRLRLSCAASGFTFSHGMHWVRQPPGKCLE WVAWIWDGSNEYYGDSVKGRFTISRDNKNTLYLQMNSLR VEDTAVYCTRGGHWNYEGHYYGMDVWGGGTTVTVSSG GGSGGGGGGGGSAVPTQPSSLASPGASASLTCTLRSGIN VGSSRIYWYQKPGPPQLRLRYTSDSKLQSGGVPSRFGSK DASANAGLLISGLQSEDEADYYCMIWHSSAVVFGCGTKLTV LG</p>	<p>DIQMTQSPSSVSASVGDRVVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLAQSGVPSRFSGSGTDFTLTISSLQPEDFATY YCQQSNSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKD STYSLSSITLTLTKADYEEKHKVYACEVTHQGLSSPVTKSFNRG EC</p>
		(SEQ ID NO: 397)	(SEQ ID NO: 398)

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	TGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCA CCCTAGTAGCTATGGCATGGCACTGGGTCCGCCAGGCTCC AGCAAAGTGCCTGGAGTGGGTGGCAGTTATCTGGTATGAT GGAAGTAATAAATACTATGCAGACTCCGTGAAGGCCGAG TCACCATCTCCAGAGACAATTCCAGAACACGCTGTATCT GCAAATGAATAGCCTGAGAGCCGAGGACACGGCTGTGTAT TACTGTACGAGAGATGGCCGGAACACGTCTACTTTGACA ACTGGGCCAGGGAACCTGGTCACCGTGTCTCAGGAGG GGCGGATCTGGCGCGGAGGAAGCGTGGGGCGGCTC CCAGTCTGCCCTGACTCAGCCTCGCTCAGTGTCCGGAGCC CTGGACAGTCAGTCACCATCTCTGCACTGGAACCCAGCAG TGATGTTGGTGGTTATATCTTTGTCTCTGGTACCAACAAC ACCCAGGCAAGCCCCCAAACTCATGATTTATGATGTCAG TAAGCGGCCCTCTGGGTCCCTGATCGCTTCTCTGGCTCCA AGTCTGTCAACACGGCTCCCTGACCATCTCTGGGCTCCAG GCTGAGGATGAGACTGATTATTACTGCTGCTCATATGCAG GCAACTACACTTATGTCTCGGATGCGGGACCAAGGTCAC CGTCCTAGGT	(SEQ ID NO: 399)
		(SEQ ID NO: 400)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQAP GKLEWISYISSSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGQGTLTVTSSASTKGPSV FPLAPSSKSTSGGT AALGCLVKDYFPEPTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTK VDKKVEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDITLMI SRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA A A VEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMHREALHNHYTQKSLSLSPGGGGGGGGSQVQLV ESGGGVVQPGRLRLSCAASGFTLSSYGMHWVRQAPGKCLE WVAWIWYDGSNKYYADSVKGRVTISRDNKNTLYLQMNSL RAEDTAVYYCTRDRGRNYVYFDNWGGQGLTVTSGGGGSGG GGSGGGGSQSALTQPRSVSGSPGQSVTISCTGTSSDVGGYIFV SWYQQHPGKAPKLMYDVSKRPSGVPDFRFSGKSVNTASLTI SGLQAEDETDYYCCSYAGNYTYVFGCGTKVTVLG (SEQ ID NO: 401)	DIQMTQSPSSVSASVGDRVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLAQSGVPSRFSGSGGTDFTLTISSLQPEDFATY YCQQSNSEPRTFGQGTKEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKD STYSLSSITLTLTKADYEEKHKVYACEVTHQGLSSPVTKSFNRG EC (SEQ ID NO: 402)
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	CTCGAGACCCCTCTCACTCACTGCTGCCATCTCCGGGACA GTGTCTCTAGCAGCCGTACTGCTTGGAACTGGATCAGGCA GTCCCATCGAGATGCCCTTGAGTGGCTGGGAAGGACATAC TACAGGTCCAAAGTGTATCATGATTATTCAGTATCTGTGAA AAGTCGAATCACCATCGACCCAGACACATCCAAGAACCCAG TTCTCCCTGCAGCTGAACCTCTGTGACTCCCCGAGGACACGGC TGTTTATTATTGTGCAAGAGGGGTGCTCCCTTTGACTACT GGGCCAGGGAACCTGGTCACTGCTCCTCAGGAGGGGG CGGATCTGGCGGGAGGAAGCGGTGGGGCGGCTCCGA AATTGTGTGACGCAGTCTCCAGGCACCTGCTTTGTCTC CAGGGGAAAGAGCCACCCTCTCTGCAGGGCCAGTCAGAG TGTTAGCAGCAACTACTTAGCTGGTACCAACAGAAACCT GGCCAGGCTCCCAGGGCCCTTATCTATGCTGCATCCAACA GGGCCGCTGGCATCTCAGACAGGTTCAGTGGCAGTGGGTC TGGGACAGACTTCACTCAACATCAGCAGACTGGAGCCT GAAGATTTGCAGTGTATTCTGTCAAGCAGTATGGTAGCTC ACCGTCACTTTCGGCTCGGGACTAAGGTGGAGATCAAA CGA	(SEQ ID NO: 403)
		(SEQ ID NO: 404)

		<p>QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQAP GKGLEWISYISSSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGGQTLVTVSSASTKGPSV FPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHHKPSNTK VDKKVEPKSCDKTHITCPPCPAPELGGPSVFLFPPKPKDITLMI SRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI A SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA A VEWESNGQPENNNYKTPPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMHREALHNYTQKSLSPGGGGGGGGGQVQLQ QSGPGLVKPSQTLSTCAISGDSVSSRTAANNWIRQSPRCLE WLGRTYYRSKWHYHDYSVVSKRITIDPDTSKNQFSLQLNSVT PEDTAVYYCARGAAPFDYWGGQTLVTVSSGGGGGGGGGGSG GGGSEIVLTQSPGTLSPGERATLSCRASQSVSSNYLAWYQQ KPGQAPRALIYAASNRAAGISDRFSGSGGTDFTLTISRLEPED FAVYFCQQYGSSPLTFGCGTKVEIKR (SEQ ID NO: 405)</p>	<p>DIQMTQSPSSVSASVGDRVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLAQSGVPSRFSGSGGTDFTLTISSLQPEDFATY YCQQSNSEPRFTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKD STYSLSSITLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRG EC</p>
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	TGAGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGAATTCA CCTTCAGTGACTACTACATGAGCTGGATCCGCCAGGCTCCA GGGAAGTGCCCTGGAGTGGGTTTCATATATTAGTCGAAGTG GTGATACCATCTACTACGAGACTCTGTGAAGGGCCGATT CACCATCTCAGGGACAACGCCAAGAACTCACTGTATCTG CAAATGAATGGCTGCGAGCCGAAGACACGGCCGTGTATT ACTGTGCGAGAGACTTAGCAGCAGGTGCTACAGGGGGCCT TGACTGCTGGGGCCAGGGAACCTGGTCACCGTGTCCCTCA GGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGGC GGCTCCTCCTATGAGCTGACTCAGCCACCTCAGTGTCCGT GTCCCAGGACAGACAGCCAGCATCACCTGCTCTGGAGAA AGGTGGGAAATAATATATTGCTGGTATCAGCAGAAAGC CAGGCCAGTCCCTGTCTGGTCATCTATCAAGATTTC AAG CGCCCTCAGGGATCCCTGAGCGATTCTCTGGCTCCAACTC TGGGATCAGAGCCACTCTGACCATCAGCGGACCCAGGCT ATGGATGAGGCTGACTATTACTGTGAGGCGTGGGACAGCA GAACTGTGGTATTCGGCTGCGGGACCAAGCTGACCGTCT AGGT	(SEQ ID NO: 407)
		(SEQ ID NO: 408)

	QVQLVESGGGLVKPGGSLRLSCAASGFTFSDDYMTWIRQAP GKLEWISYISSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGQGTLLVTVSSASTKGPSV FPLAPSSKSTSGGTAAAGCLVKDYFPEPVTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHHKPSNTK VDKKVEPKSCDKTHITCPPCPAPPELLGGPSVFLFPPKPKDITLMI SRTPETCVVDVSHEDPEVKFNWYVVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA VEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMHEALHNHYTQKSLSPGGGGGGGGSGVQLV ESGGGLVKPGGSLRLSCAASEFTFSDDYMSWIRQAPGKCLEW VSYISRSGDTIYYADSVKGRFTISRDNAKNSLYLQMNGLRAE DTAVYYCARDLAAGATGGLDCWGQGTLLTVSSGGGGGGGG GSGGGGSSVELTQPPSVSPGQTASITCSGERLGNKYICWYQ QKPGQSPVLVIYQDFKRPSPGIPERFSGSNSGITATLTISGTQAM DEADYYCQAWDSRTVVFSGCGTKLTVLG	(SEQ ID NO: 409)
	DIQMTQSPSSVSASVGDRVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLQSGVPSRFSGSGGTDFTLTISSLQPEDFATY YCQQSNSFPRTFGQGTKEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKD STYLSSTLTLSKADYEEKHKVYACEVTHQGLSSPVTKSFNRG EC	(SEQ ID NO: 410)

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	TGGGGCCTCAGTGAAAGGTGTCCTGCAAGGCTTCTGGATAC ACCTTCCC CGGCTACTATATGCACTGGGTGCGACAGGCCCC TGGACAGTGCCTTGAGTGGATGGGATGGATCAACCCTGAC AGTGGTGGCACAAAGTATACACAGAAAGTTTCAGGGCAGGG TCACCTTGACCAGGGACGCGTCGTCAGCACAGCCTACAT TGACCTGAACACAGGCTGAGATCTGACGACACGGCCGTATAT TACTGTGCGAGAGAGAGGTGTAGGACTACCAACTGCTATT TGGACTACTGGGCCAGGGAAGTCTGGTCAACCGTGTCTC AGGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGG CGGCTCCCAGTCTGCCCTGACTCAGCCTGCCCTCCGTCTG GGAGCCCTGGACAGTCGATCACCATCTCCTGCACTGGAAC CAGCAGTGATGTTGGGAATTATAACCTTGTCTCCTGGTACC AACAGCACCCAGGCAAGCCCCCAACTCATGATTATGA GGTCAATAGCGGCCCTCAGGGGTTTCTAATCGCTTCTCTG GCTCCAAGTCTGGCAACACGGCCTCCCTGACAACTCTCTGG GCTCCAGGCTGAGGACGAGGCTGAATATTACTGCTGCTCA TATGCAGGTAGAGACACTTTCGTGGTGTTCGGCTGCGGGA CCAAGCTGACCGTCCCTAGGT	(SEQ ID NO: 411)
		(SEQ ID NO: 412)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYMTWIRQAP GKLEWISYISSGSTIYYADSVKGRFTISRDNAKNSLYLQMIN SLRAEDTAVYYCARDNRSHFDYWGGQTLVTVSSASTKGPSV FPLAPSSKSTSGGT AALGCLVKDYFPEPVTVSWNSGALTSGV HTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHHKPSNTK VDKKVEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDITLMI SRTPETCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEE QYGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI SKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIA A A VEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQ GNVFSCSVMEALHNHYTQKSLSLSPGGGGGGGSGVQLV QSGTEVKKPGASVKVSCKASGYTFPGYYMHWVRQAPGQCL EWMGWINPDSGGTKYTQKFQGRVTLTRDASVSTAYIDLNRL RSDDTAVYYCARERCRTTCYLDYWGQGLVTVSSGGGGSG GGSGGGGQSALTPASVSGSPQSITISCTGTSSDVGNYNL VSWYQQHPGKAPKLMIEVNRPRPSGVSNRPSGSKSGNTASLT ISGLQAEDEAEYYCCSYAGRDTFVVFSGCGTKLTVLG (SEQ ID NO: 413)	DIQMTQSPSSVSASVGDRVTITCRASQGITRWLAWYQQKPGK APKLLIYAASVLSQGVPSRFSGSGGTDFTLTISLSQPEDFATY YCQQSNSEPRFTGQGTKEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKD STYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRG EC (SEQ ID NO: 414)
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	TGGGGCCTCAGTGAAAGGTGTCTGCAAGGCTTCTGGATAC ACCTTCGCCGGCTACTATATGCACTGGGTGCGACAGGCCC CTGGACAAATGCCTTGAGTGGATGGATGGATCAACCCTGA CAGTGGAGGCACAAACTTTGCACAGCAGTTTCAGGGCAGG GTCACCATGACCAAGGATACGTCCATCAGCACAGCCTACA TGGAGGTGAGCAGGCTGAGATCTGACGACACGGCCGTGTT TTACTGTGCGAGAGAGAAAGATCACTATGACTGGTATTTACT TTGACTATTGGGGCCAGGGAACCTGGTCACCGTGTCTCA GGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGGC GGCTCCGACATCCAGATGACCCAGTCTCCATCCTCCCTGTC TGCATCTGTAGGAGACATTTCTACCATCACTTGCCGGGCAA GTCAGAACATTACCACTATTAAATTGGTATCAGCAGAA ACCAGGAAAGCCCCTAACCTCCTGATCTCTGCTGCATCCC GTTTGGAAAGTGGGTCCCATCAAGGTTCAAGTGGCAGTGG ATCTGGGACAGATTTCACTCTCAACCATCAGCAGTCTGCAAC CTGTAGATTTACAACCTTCTACTGTCAACAGACTTTCAC ACCCGTGGACGTTCGGCTGCGGGACCAAGGTGGAGATCA AACGA	(SEQ ID NO: 415)
		(SEQ ID NO: 416)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDY ^Y MSWIRQAPG KGLEWISYISSESI ^Y YVDSVKGRFTISRDN ^{AK} NSLYLQMN ^{SL} RAEDTAVYYCARDVGS ^H FDYWGQGT ^{LV} TVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEP ^{VT} SVWNSGALTSGVHT FPAVLQSSGLYSLSSV ^{TV} PSSSLGTQYICNVN ^{HK} PSNTKVD KKVEPKSCDK TH TCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNW ^Y VDGVEVHN ^{AK} TKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYCKVSNKALPAPIEK ^{TIS} KAKGQPREPQVY ^{TL} PPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYK ^{TT} PPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSV ^M HEALHNHYTQKSLSLSPGGGGGGGQVQLVQ SGAEVKKPGASVK ^{SK} ASGYTFA ^{GY} YMH ^W VRQAPGQCLE WMGWINPDSGGTNFAQQFQGRV TM TRDTSISTAYMEVSR ^{LR} SDDTAVFYCAREKITMTGIYFDYWGQGT ^{LV} TVSSGGGGSGG GGSGGGGSDIQMTQSPSSLSASVGDIL ^{TIT} CRASQNITTYL ^{NW} YQOKPGKAPNLLISAASRLRSGVPSRFSGSGGTDFLT ^{TI} SSLQ PVDFTTFYCQQTFTTPWTFGCGTK ^{VEIKR}	(SEQ ID NO: 417)
	A		
	A		
		DIQMTQSPSSVSASVGDRV ^{VT} TITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLT ^{TI} SSLQPEDFAI ^{YY} CQQA ^K SFPRTFGQGTK ^{VEIKR} TVAAPSVFIFPPSDEQLKSGTA SVVCLLN ^{NF} YPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C	(SEQ ID NO: 418)

[illegible]

	TGGGGCCTCAGTGAAAGGTGTCTGCAAGGCTTCTGGATAC ACCTTCGCCGGCTACTATATGCACTGGGTGCGACAGGCCC CTGGACAAATGCCCTTGAGTGGATGGATGGATCAACCTCA CAGTGGTGGCACAAACTATGCACAGAAATTTCAGGACAGG GTCACCATGACCAAGGACACGTCCATCAACACAGCCTACA TGGAACCTGAGCAGGCTGAGATCTGACGACACGGCCGTGA TTACTGTGCGAGAGAACGTATTTCTATGGTTGCGGGAGTCG GGCACAACTGGTTCGCCCCCTGGGGCCAGGAAACCTGGT CACCGTGTCTCAGGAGGGGGCGGATCTGGCGGGAGGA AGCGGTGGGGGGCTCCGACATCCAGATGACCCAGTCTC CATCCTCACTGTCTGCATCTGTAGGAGACAGAGTCACCATC ACCTGTGGGGCAGTCAGGACATTAGCAATAATTAGCCT GGTTTCAGCAGAAACCAGGAAACCCCTAAGTCCCTGAT GTATGCTGCATCCAGTTGCACAGTGGAGTCCCATCAACGT TCAGCGGCAGTGGATCTGGGACAGATTTCACTTTCACCATC AGCAGCCTGCAGCCTGAAGATTTTGCAACTTATTACTGCCA ACAGTATAATAGTTACCTCTCACTTCGGCTGCGGGACCA AGGTGGAGATCAGACGA	(SEQ ID NO: 419)
		(SEQ ID NO: 420)

	QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSESIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGLTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHITCPPCPAPELLGGPSVFLFPPKPKDTLMSIR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIIS KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSLSPGSGGGGGGQVQLVQ SGAEVTKPGASVKSCASGYTFAGYYMHWVRQAPGQCLE WMGWINPHSGGTNYAQKFQDRVTMTTRDTSINTAYMELSLRLR SDDTAVYYCARERISMVRGVGHNWFAPWGQGLTVTVSSGG GGSGGGSGGGGSDIQMTQSPSSLSASVGDRVTITCRASQDIS NNLAWFQQKPGKPPKSLMYAASSLHSGVPTSFSGSGGTDF FTISSLQPEDFATYYCQYNSYPLTFGCGTKVEIRR	DIQMTQSPSSVSASVGDRVTITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLTITSSSLQPEDFAIYY CQQAQKSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C
	(SEQ ID NO: 421)	(SEQ ID NO: 422)

[illegible]

	TGGGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGGATTCA CCTTAGTAGAAATGCCATGAGTTGGTCCGCCAGGCTCC AGGAAAGTGCCTGGAGTGGGTGCAGCTACTGGTGGTAGT GGTATTAGCACATACTACGCAGACTCCGTGAAGGGCCGGT TCACCATCTCCAGAGACAATTCCAGAACACGCTGTATCT GCAAAATGAACAGTCTGAGAGCCGAGGACACGGCCGTATAT TACTGTGCGAGAGGTTATAGCAACAGCTGGTGGTACTTTG ACTACTGGGCCAGGGAACCTGGTCACCGTGTCTCTCAGG AGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGGCGG CTCCAGTCTGCCCTGACTCAGCCTGCCCTCCCTGCTGGGA GCCCTGGACAGTCGATCACCATCTCCTGCACTGGAAACCAG CAGTGATGTTGGGAATTATAACCTTGTCTCCTGGTACCAAC AGCACCCAGGCAAGCCCCCAAACTCATGATTTTGAGGT CAATCAGCGGCCCTCAGGGGTTTCTAATCGCTTCTCTGGCT CCAAAGTCTGGCACCAAGGCCCTCCCTGACAACTCTCTGGGCTC CAGGCTGCGGACGAGGCTGATTATTTCTGCTGCTCATATAC AACTAGTAGCACTTATGTGATCTTCGGCTGCGGGACCAAG CTGACCGTCTCTAGGT	(SEQ ID NO: 423)
		(SEQ ID NO: 424)

			QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSESIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGTLVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHITCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTIS KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSPGGGGGGGGSEVQLLES GGGLVQPGGSLRLSCAASGFTFSRNAMSWVRQAPGKCLEWV SATGGGISTYYADSVKGRFTISRDNKNTLYLQMNSLRAED TAVYYCARGYSNSWWYFDYWGGGTLVTVSSGGGGGGGGGS GGGGSQSALTQPASVSGSPGQSITISCTGTSSDVGNYNLVSWY QQHPGKAPKLMIFEVNRQPSGVSNRFGSKSGTTASLTISGLQ AADEADYFCCSYTTSSTYVIFGCGTKLTVLG (SEQ ID NO: 425)	DIQMTQSPSSVSASVGDRVTITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLTITSSLQPEDFAIYY CQQAQKSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQQLSSPVTKSFNRRGE C (SEQ ID NO: 426)

250

	TGGGGCCTCAGTGAAAGGTGTCCTGCAAGGCTTCTGGATAC ACCTTCCCGGCTACTATATGTACTGGTTGCGACAGGCCCC TGGACAATGCCCTTGAGTGGATGGATGGATCAACCCCTGAC AGTGGTGACACAACTATGCACAGAA GTTTCAGGGCAGGG TCACCATGACCAGGACACGTCCATCAGCACAGCCTTTAT GGAGCTGAGCAGGCTGAGATCAGACGACACGGCCGTGTAT TACTGTGCGAGAGAGAGCCAGATATTTGACTCCTTCTA CTACTACCTTATGGACGTCTGGGGCCAAGGACCCACGGTC ACCGTGTCTCAGAGGGGGCGGATCTGGCGCGGAGGAA GCGGTGGGGCGGCTCCGACATCCAGATGACCCAGTCTCC ATCCTCCCTGTCTGCATCTGTAGGAGACAGAGTCACCATCA CTTGCCGGGCAGGTCAAGAACATTAGCAGGCATTAAATTG GTATCAGCAGAAATCCAGGGAAGCCCCCTAAGGTCTGATC CATCCTGCATCCAGTTTGCCAAAGTGGGGTCCCGTCAAGGTT CAGTGGCAGTGGATCTGGGACAGATTTCA GTCTTACCATC AGCAGTCTGCAACCTGAAGATTTTGGAACTTACTTCTGTCA ACAGAGTTACAGTACCCCTCCC ACTTTGCGCTGCGGGACC AAGGTGGAGCTCAACGA	(SEQ ID NO: 427)
		(SEQ ID NO: 428)

			QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSESIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGTLVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTIIS KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSLSPGSGGGGGGQVQLVQ SGAEVKKPGASVKVSCKASGYTFPGYYMYWLRQAPGQCLE WMGWINPDSGDTNYAQKFQGRVTMTRTDSISTAFMELSLRLR SDDTAVYYCAREKPRYFDSFYLYLMDVWGQGTITVTVSSGG GGSGGGSGGGSDIQMTQSPSSLSASVGDRTVITTCRAGQNIIS RHLNYYQQNPGKAPKVLHPASSLPAGVPSRFSFGSGGTDGSL TISSLQPEDFGTYFCQQQSYSTPPTFGCGTKVELKR (SEQ ID NO: 429)	DIQMTQSPSSVSASVGDRVITTCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSFGSGGTDFTLTITSSSLQPEDFAIYY CQQAQKSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C (SEQ ID NO: 430)

	TGGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCA CCTTCAGTAGCCATGGCATGCACTGGGTCCGCCAACCTCCA GGCAAGTGCCTGGAGTGGGTGGCAGTTATCTGGTATGATG GAAGTAATGAATACTATGGAGACTCCGTGAAGGCCGATT CACCATCTCAGAGACAATTCCAAGAACACGCTGTATCTG CAAATGAACAGCCTGAGAGTCGAGGACACGGCTGTGTATT ACTGTACGAGAGGGGGGGCCACTGGAACTACGAGGGCC ACTACTATGGTATGGACGTCTGGGGCCAAGGACCAACGGT CACCGTGTCTCAGGAGGGGGCGGATCTGGCGGGGAGGA AGCGTGGGGGGCTCCAGGCTGTGCCGACTCAGCCCT CTCCCTCTCTGCATCTCCTGGAGCATCAGCCAGTCTCACC TGCACCTTACGAGTGGCATCAATGTTGGTTCTCCAGGAT CTATTGGTACCAGCAGAAAGCCAGGAGTCTCCCCAGTTT CTCCTGAGGTACACATCAGACTCAGATAAAATTGCAGGGCT CTGGAGTCCCCAGCCGCTTCTCTGGATCCAAAGATGCTTCG GCCAATGCAGGACTTTTACTCATCTCTGGGCTCCAGTCTGA GGATGAGGCTGACTATTACTGTATGATTGGCACAGCAGC GCTGTGGTATTCTGGCTGCGGGACCAAACTGACCGTCTCTAG GT	(SEQ ID NO: 431)	(SEQ ID NO: 432)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSEIIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGLTVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHITCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTI KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSLSPGSGGGGGGQVQLVE SGGVVQPGRLRLSCAASGFTSSHGMHWVRQPPGKCLEW VAVIWDGSNEYYGDSVKGRFTISRDNKNTLYLQMNSLRV EDTAVYCTRGGHWNYEGHYYGMDVWGGGTTTVSSGG GGSGGGGGGGGQA VPTQPSSLASPGASASLTCTLRSGINV GSSRIYWYQQKPGSPQQFLRYTSDSKLQSGGVPSRFGSKD ASANAAGLLISGLQSEDEADYYCMHWSSAVVFGCGTKLTVL G	DIQMTQSPSSVSASVGDRVVTITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLTITSSLQPEDFAIYY CQQA KSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQQLSSPVTKSFNRGE C
		(SEQ ID NO: 433)	(SEQ ID NO: 434)

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	TGGAGGTCCCTGAGACTCTCCTGTGCAGCGTCTGGATTCA CCCTAGTAGCTATGGCATGGCACTGGGTCCGCCAGGCTCC AGCAAAGTGCCTGGAGTGGGTGGCAGTTATCTGGTATGAT GGAAGTAATAAATACTATGCAGACTCCGTGAAGGCCGAG TCACCATCTCCAGAGACAATTCCAGAACAACGCTGTATCT GCAAATGAATAGCCTGAGAGCCGAGGACACGGCTGTGTAT TACTGTACGAGAGATGGCCGGAACACGTCTACTTTGACA ACTGGGCCAGGGAACCTGGTCACTCCGTCTCAGGAGG GGCGGATCTGGCGCGGAGGAAGCGTGGGGCGGCTC CCAGTCTGCCCTGACTCAGCCTCGCTCAGTGTCCGGAGCC CTGGACAGTCAGTCACCATCTCTGCACTGGAACCAAGCAG TGATGTTGGTGGTTATATCTTTGTCTCTGGTACCAACAAC ACCCAGGCAAGCCCCCAAACTCATGATTTATGATGTCAG TAAGCGGCCCTCTGGGTCCCTGATCGCTTCTCTGGCTCCA AGTCTGTCAACACGGCTCCCTGACCATCTCTGGGCTCCAG GCTGAGGATGAGACTGATTATTACTGCTGCTCATATGCAG GCAACTACACTTATGTCTCGGATGCGGGACCAAGGTCAC CGTCCTAGGT	(SEQ ID NO: 435)
		(SEQ ID NO: 436)

		QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSESIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGTLVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHITCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTIS KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSPGGGGGGGGGQVQLVE SGGGVVPGRSLRLSCAASGFTLSSYGMHWVRQAPGKCLEW VAVIWDGSNKYYADSVKGRVTSRDNSKNLTLYLQMNSLRA EDTAVYCTRDRGRNVYFDNWGQGTLVTVSSGGGGGGGGG SGGGGSQSALTQPRSVSGSPGQSVTISCTGTSSDVGGYIFVSW YQQHIPGKAPKLMYDVSKRPSGVPDFRFGSKSVNTASLTISGL QAEDETDYYCCSYAGNYTYVFGCGTKVTVLG (SEQ ID NO: 437)	DIQMTQSPSSVSASVGDRVTITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLTITSSLQPEDFAIYY CQQAQKSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQQLSSPVTKSFNRRGE C (SEQ ID NO: 438)
	A		
	A		

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	CTCGAGACCCCTCTCACTCACTGTGCCATCTCCGGGACA GTGTCTTAGCAGCCGTACTGCTTGGAACTGGATCAGGCA GTCCCATCGAGATGCCCTTGAGTGGCTGGGAAGGACATAC TACAGGTCCAAGTGGTATCATGATTATTCAGTATCTGTGAA AAGTCGAATCACCATCGACCCAGACACATCCAAGAACCCAG TTCTCCCTGCAGCTGAACCTGTGTGACTCCCCGAGGACACGGC TGTTTATTATTGTGCAAGAGGGGTGCTCCCTTTGACTACT GGGCCAGGGAACCTGGTCACCGTGTCTCAGGAGGGGG CGGATCTGGCGGGGAGGAAGCGGTGGGGCGGCTCCGA AATTGTGTGACGCAGTCTCCAGGCACCTGCTTTGTCTC CAGGGGAAGAGCCACCCTCTCTGCAGGGCCAGTCAGAG TGTTAGCAGCAACTACTTAGCTGGTACCAACAGAAACCT GGCCAGGCTCCCAGGGCCCTTATCTATGCTGCATCCAACA GGCCCGCTGGCATCTCAGACAGGTTCAGTGGCAGTGGGTC TGGGACAGACTTCACTCAACATCAGCAGACTGGAGCCT GAAGATTTGCAGTGTATTCTGTCAAGCAGTATGGTAGCTC ACCGTCACTTTCGGCTGCGGACTAAGGTGGAGATCAAA CGA	(SEQ ID NO: 439)
		(SEQ ID NO: 440)

	QVQLVESGGGLVKPGGSLRLSCAASGFTFSDYYMSWIRQAPG KGLEWISYISSEIIYYVDSVKGRTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGS ^H FDYWGQGT ^L VT ^V SSASTKGPSVFP LAPSSKSTSGGTAA ^L GCLVK ^D YFPEP ^V TVSWNSGALTSGVHT FPAVLQSSGLYSLSSV ^V TV ^P SSSLGTQ ^T YICNVN ^H K ^P SN ^T K ^V D KKVEPKSCDK ^T HTCP ^P CPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYYVDGVEVHNAKTKPCEEQ YGSTYRCVSV ^L TVLHQDWLNGKEYKCKVSNKALPAPIEK ^T IS A KAKGQPREPQVY ^T LP ^P SREEMTKNQVSLTCLVKGFYPSDIAV A EWESNGQPENNYK ^T TPPVLDSGDSFFLYSKLTVDKSRWQQG NVFSCSV ^M HEALHNHYTQKSLSPGGGGGGGGGSQVQLQQ SGPGLVKPSQ ^T LSLTC ^A ISGDSVSSRTA ^W NWIRQSPRCLEW LGR ^T YYRSKWYHDYSVSVKSRITIDP ^T SKNQ ^F SLQLNSVTPE DTAVYYCARGAAPFDYWGQGT ^L VT ^V SSGGGGGGGGGGGGGG GSEIVLTQSPGTLSLSPGERATLSCRASQSVSSNYLAWYQQKP GQAPRALIYAAASNRAAGISDRFSGSGGTDFTLTISRLEPEDFA VYFCQQY ^G SSPLTFGCGTKVEIKR (SEQ ID NO: 441)	DIQMTQSPSSVSASVGDRV ^T ITTCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLT ^T ISSLQPEDFAIYY CQQA ^K SFPR ^T FGQGT ^K VEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLN ^N FYP ^R EAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C
		(SEQ ID NO: 442)

	TGAGGGTCCCTGAGACTCTCCTGTGCAGCCTCTGAATTCA CCTTCAGTGACTACTACATGAGCTGGATCCGCCAGGCTCCA GGGAAGTGCCCTGGAGTGGGTTTCATATATTAGTCGAAGTG GTGATACCATCTACTACGAGACTCTGTGAAGGGCCGATT CACCATCTCAGGGACAACGCCAAGAACTCACTGTATCTG CAAATGAATGGCTGCGAGCCGAAGACACGGCCGTGTATT ACTGTGCGAGAGACTTAGCAGCAGGTGCTACAGGGGGCCT TGACTGCTGGGGCCAGGGAACCTGGTCACCGTGTCCCTCA GGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGGC GGCTCCTCCTATGAGCTGACTCAGCCACCTCAGTGTCCGT GTCCCAGGACAGACAGCCAGCATCACCTGCTCTGGAGAA AGGTTGGGAAATAATATATTGCTGGTATCAGCAGAAAGC CAGGCCAGTCCCTGTCTGGTCACTATCAAGATTTC AAG CGCCCTCAGGGATCCCTGAGCGATTCTCTGGCTCCAACTC TGGGATCAGAGCCACTCTGACCATCAGCGGAGCCACAGGCT ATGGATGAGGCTGACTATTACTGTGAGGCGTGGGACAGCA GAACTGTGGTATTGGCTGCGGGACCAAGCTGACCGTCT AGGT	(SEQ ID NO: 443)
		(SEQ ID NO: 444)

		<p>QVQLVESGGGLVKPGGSLRLSCAASGFTSDYYMSWIRQAPG KGLEWISYISSEIIYYVDSVKGRFTISRDNAKNSLYLQMNSL RAEDTAVYYCARDVGSHEFYWGQGTLVTVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHT FPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHITCPPCPAPELLGGPSVFLFPPKPKDTLMISR TPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQ YGSTYRCVSVLTVLHQDWLNGKEYCKVSNKALPAPIEKTIS KAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSVMEALHNHYTQKSLSPGGGGGGGGSQVQLVE SGGGLVKPGGSLRLSCAASEFTSDYYMSWIRQAPGKCLEW VSYISRSGDTIYADSVKGRFTISRDNAKNSLYLQMNGLRAE DTAVYYCARDLAAGATGGLDCWGQGTLVTVSSGGGGGGG SGGGGSSYELTQPPSVSPGQTASITCSGERLGNKYICWYQ QKPGQSPVLVIYQDFKRPSPGIPERFSGNSGITATLTISGTQAM DEADYYCQAWDSRTTVVFGCGTKLTVLG</p>	<p>DIQMTQSPSSVSASVGDRVVTITCRASQDISRWLAWYQQKPGK APKLLISAASRLQSGVPSRFSGSGGTDFLTITSSLPEDFAIYY CQQAQKSFPRTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGE C</p>
		<p>(SEQ ID NO: 445)</p>	<p>(SEQ ID NO: 446)</p>

	TGGGGCCTCAGTGAAAGGTGTCCTGCAAGGCTTCTGGATAC ACCTTCCCGGCTACTATATGCACTGGGTGCGACAGGCCCC TGGACAGTGCCTTGAGTGGATGGATGGATCAACCCTGAC AGTGGTGGCACAAAGTATACACAGAAAGTTTCAGGGCAGGG TCACCTTGACCAGGGACGCGTCAGCACAGCCTACAT TGACCTGAACACAGGCTGAGATCTGACGACACGGCCGTATAT TACTGTGCGAGAGAGAGGTGTAGGACTACCAACTGCTATT TGGACTACTGGGCCAGGGAAGTCTGGTCAACCGTGTCTC AGGAGGGGGCGGATCTGGCGCGGAGGAAGCGGTGGGG CGGCTCCAGTCTGCCCTGACTCAGCCTGCCCTCCGTCTG GGAGCCCTGGACAGTCGATCACCATCTCCTGCACTGGAAC CAGCAGTGATGTTGGGAATTATAACCTTGTCTCCTGGTACC AACAGCACCCAGGCAAGCCCCCAACTCATGATTATGA GGTCAATAGCGGCCCTCAGGGGTTTCTAATCGCTTCTCTG GCTCCAAGTCTGGCAACACGGCCTCCCTGACAACTCTCTGG GCTCCAGGCTGAGGACGAGGCTGAATATTACTGCTGCTCA TATGCAGGTAGAGACACTTTCGTGGTGTTCGGCTGCGGGA CCAAGCTGACCGTCCCTAGGT	(SEQ ID NO: 447)
		(SEQ ID NO: 448)

	QVQLVESGGGLVKPGGSLRLSCAASGFTSDY ^Y MSWIRQAPG KGLEWISYISSEII ^Y YVDSVKGRFTISRDN ^{AK} NSLYLQMNSL RAEDTAVYYCARDVGS ^H FDYWGQGT ^{LV} TVSSASTKGPSVFP LAPSSKSTSGGTAALGCLVKDYFPEP ^{VT} SVWNSGALTSGVHT FPAVLQSSGLYSLSSV ^{TV} TPSSSLGTQTYICNVN ^{HK} PSNTKVD KKVEPKSCDK TH TCPPCPAPELLGGPSVFLFPPKPKD ^{TL} MISR TPEVTCVVVDVSHEDPEVKFNW ^Y VDGVEVHN ^{AK} TKPCEEQ YGSTYRCVSV ^{LV} TVLHQDWLNGKEYCKVSNKALPAPIEK ^{TIS} KAKGQPREPQVY ^{TL} PPSREEMTKNQVSLTCLVKGFYPSDIAV EWESNGQPENNYK ^{TT} PPVLDSDGSFFLYSKLTVDKSRWQQG NVFSCSV ^{MHE} ALHNHYTQKSLSLSPGSGGGGGG ^{SV} QLVQ SGTEVKKPGASVK ^{SK} ASGYTPGY ^Y MHWVRQAPGQCLE WMGWINPDSGGTKY ^T QKFGQGRV ^{TL} TRDASV ^{TA} YIDLNRLR SDDTAVYYCARE ^{RCRT} NCYLDYWGQGS ^{LV} TVSSGGGGSG GGSGGGGGS ^Q SA ^L TQPASVSGSPGQSI ^{TS} CTGTSSDV ^{GN} YNL VSWYQQHPGKAPK ^{LM} MIYEVNRRPSGV ^{SN} RFSGKSGNTASLT ISGLQAEDEAEYYCCSYAGRD ^{TF} VVFGCGTK ^{LV} LG	(SEQ ID NO: 449)
A	DIQMTQSPSSVSASVGDRV ^{VT} ITCRASQDISRW ^{LA} WYQQKPGK APKLLISAASRLQSGVPSRFSGSGGT ^{DF} TLTISSLQPEDFAI ^{YY} CQQA ^K SFPRTFGQGT ^K VEIK ^{RTV} AAPSVFIFPPSDEQLKSG ^{TA} SVVCLLN ^{NF} YPREAKVQW ^K VDNALQSGNSQESVTEQDSK ^{DS} TYSLSST ^{LT} LSKADYEKHK ^{VY} ACEV TH QQLSSPV ^{TK} SFNRGE	(SEQ ID NO: 450)
A	C	

Table 27A: CD-40-MSLN IgG-Fab

iPS #	Ab	Description	Type	LC E	LC K	IgFab HCv1

269

270

271

		DIQMTQSPSSLSASVGDILITICRASQNI TTYLNWYQQKPGKAPNLLISAA SRLRS GVPSRFGSGGTDFTLTISSLQPVDF TFYCQQTFTTPWTFGQGTKEIKRTVA APSVFIIPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTE QDSKDSYSTYSLSTLTLSKADYEKHKVY ACEVTHQGLSSPVTKSFNRGEC	QSVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAIT GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLQPKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETITTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	QVQLVQSGAEVKKPGASVKVCKAS GYTFAGYIMHWVRQAPGQGLEWM GWINPDSGGTNFAQQFQGRVTMTDR TSISTAYMEVSRRLRSDDTAVFYCARE KITMTGIYFDYWGQGTLVTVSSAST KGPSVFPLAPSSKSTSGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFP AVLQSSGLYSLKSVTVVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDK THTCPPCPAPELLGGPSVFLFPPKPKD TLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPCEEQYGST YRCVSVLTVLHQDWLNGKEYKCKV SNKALPAPIEKTISKAKGQPREPQVY TLPPSRREEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTTPPVLD DGSFFLYSKLTVDKSRWQQGNVFS SVMHEALHNHYTQKSLSLSPGGGS GGGSGVQLQESGPGLVKPSSETLSLT CTVSGGSISSSSYYWGWIRPPGKGL EWIGSIYYSGITNYNPSLKRVTISVD TSKNQFSLKLSSTAAADTAVYYCAR SSNYDAFDIWGGQGTMTVTVSSASTKG PSVFPLAPSSKSTSGGTAALGCLVKD YFPEPVTVSWNSGALTSGVHTFP LQSSGLYSLESVVTVPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSC
	(SEQ ID NO: 454)	(SEQ ID NO: 455)	(SEQ ID NO: 456)	
A				
A				

273

274

[illegible]

		DIQMTQSPSSLSASVGDILITITCRASQNI TTYLNWYQQKPGKAPNLLISASRLRS GVPSRFGSGGTDFTLTISSLQPVDF TFYCQQTFTTPWTFGQGTKEIKRTVA APSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTE QDSKDSITYSLESTLTLSKADYEKHKVY ACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSVSASVGDRVTITCRAS QGITRWLAWYQQKPGKAPKLLIYAA SVLQSGVPSRFGSGGTDFTLTISSL QPEDFATYYCQQSNSEFPTFGQGTK VEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNAL QSGNSQESVTEQDSKDSITYSLKSTLT LSKADYEKHKVYACEVTHQGLSSPV TKSFNRGEC	QVQLVQSGAEVKKPGASVKVSCKAS GYTFAGYYMHWRAPQGQGLEWM GWINPDSGGTNFAQQFQGRVTMTDR TSISTAYMEVSRRLRSDDTAVFYCARE KITMTGIYFDYWGQGTLVTVSSAST KGPSVFPLAPSSKSTSGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFP AVLQSSGLYSLKSVTVTPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDK THTCPPCPAPELLGGPSVFLFPPKPKD TLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNATKPCPEEQYGST YRCVSVLTVLHQDWLNGKEYKCKV SNKALPAPIEKTISKAKGQPREPQVY TLPPSRREEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTTTPVLD DGSFFLYSKLTVDKSRWQQGNVFC SVMHEALHNHYTQKSLSLSPGSGGS GGGSGVQLVESGGGLVKPGGSLRL SCAASGFTFSDYYMTWIRQAPGKGL EWISYISSSGSTIYADSVKGRFTISR DNAKNSLYLQMNSLRAEDTAVYYC ARDRNSHFYWGQGTLVTVSSASTK GPSVFPLAPSSKSTSGGTAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPA VLQSSGLYSLESVTVTPSSSLGTQTYI CNVNHKPSNTKVDKKVEPKSC
				(SEQ ID NO: 460)
				(SEQ ID NO: 461)
				(SEQ ID NO: 462)

277

278

[illegible]

[illegible]

			GACATCCAGATGACCCAGTCTCCATC CTCCCTGTCTGCATCTGTAGGAGACA TTCTCACCATCACTTGCCGGCAAGT CAGAACATTACCACCTATTTAAATTG GTATCAGCAGAAACCAGGGAAGCC CCTAACCTCCTGATCTCTGCTGCATC CCGTTTGCAGAGTGGGTCCCATCAA GGTTCAAGTGGCAGTGGATCTGGGAC AGATTTCACTCTCAACATCAGCAGTC TGCAACCTGTAGATTTTACAACTTTC TACTGTCAACAGACTTTTCACTACCCC GTGGACGTTTCGGCCAAAGGACCAAG GTGGAGATCAAAGAACGGTGGCTG CACCATCTGTCTTCATCTTCCCGCCAT CTGATGAGCAGTTGAAATCTGGAAC GCCTCTGTTGTGCTGCTGAATAA CTTCTATCCAGAGAGGCCAAAGTAC AGTGAAGGTGGATAACGCCCTCCA ATCGGGTAACCTCCAGGAGAGTGT ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTGACG CTGAGCAAAAGCAGACTACGAGAAAC ACAAAGTCTACGCCTGCGAAGTCACC CATCAGGGCCTGAGCTCGCCCGTCAC AAAGAGCTTCAACAGGGGAGAGTGT																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																															
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282

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	DIQMTQSPSSLSASVGDRTTITCRASQD ISNNLAWFQQKPGPKPSLMYAASSLH SGVPSTFSGSGGTDFTFTISSLQPEDFA TYYCQQYNSYPLTFGGGKVEIRRTVA APSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTE QDSKDSYSTLESSTLTLSKADYEKHKVY ACEVTHQGLSSPVTKSFNRGEC	QSVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAIT GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPKKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	QVQLVQSGAEVTKPGASVKVCKKAS GYTFAGYYMHWVRAPQGQGLEWM GWINPHSGGTNYAQKFQDRVTMTTR DTSINTAYMELSRLLSDDTA VYYCA RERISMVRGVGHNWFAPWGQGTLY TVSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPTVTSWNSGALTS GVHTFPAVLQSSGLYSLKSVVTPSS SLGTQTYICNVNHKPSNTKVDKKVE PKSCDKHTHTCPCPAPELLGGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNG KEYCKKVSNAKALPAIEKTIKAKGQ PREPQVYTLPPSREEMTKNQVSLTCL VKGFYPSDIAVEWESNGQPENNYKT TPPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMHEALHNHYTQKSLSL SPGGGGSGGGGQVQLQESGPGLVK PSETLSLTCTVSGSISSSSYYWGWIR QPPGKGLEWIGSIYYSGITNYPNPSLKS RVTSISVDTSKNQFSLKLSVTAADTA VYYCARSSNYDAFDIWGQGGMVTV SSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPTVTSWNSGALTSGV HTFPAVLQSSGLYSLKSVVTPSSSL GTQTYICNVNHKPSNTKVDKKVEPK SC	(SEQ ID NO: 478)	(SEQ ID NO: 479)	(SEQ ID NO: 480)
A						
A						

[illegible]

290

291

[illegible]

294

					TTTGACTACTGGGGCCAGGGAACC CTGGTCACCGTGTCTCTCAGCAAGC ACGAAGGGGGCGTCCGTATTTC CG CTTGCGCCCTCGTCGAAGTCAACTT CGGGAGGGACCGCGGCACCTTGGCT GTCTTGTCAAAGATTACTTCCCTGA GCCAGTGACAGTCAGCTGGAATTTC CGGTGCCCTCACGTCAGGAGTACA TACATTCCCTGCGGTATTGCAGTCC TCCGGACTCTACTCCCTGGAGTCGG TGGTAACGGTGGCCAGCTCCAGCTT GGGGACCCAGACGTACATTGTAA CGTGAATCACAACCAAGCAATAC TAAGGTAGATAAGAAAGTAGAACC GAAGAGCTGC	(SEQ ID NO: 487)	(SEQ ID NO: 488)	(SEQ ID NO: 489)
3086								
1								

[illegible]

[illegible]

	C + [anti - <hu CD4 0> 21- 230- 29H 10 VL]: :huK LC- S176 E + [anti - <hu Mes othel in> 7G1 I VL]: :huK LC- S176 K (IgG - Fab) ; LM R ID:		AACTGGTACGTGGACGGCGTGGAG GTGCATAATGCCAAGACAAAGCCG TGCAGGAGCAGTACGGCAGCACG TACCGTTGCTCAGCTCCTCAACG TCCTGCACCCAGACTGGCTGAATG GCAAGGAGTACAAGTGCAAGGTGT CCAACAAAGCCCTCCACGCCCCCA TCGAGAAACCATCTCCAAGCCA AAGGGCAGCCCCGAGAACACACAG TGTAACCCCTGCCCCCATCCCCGGA GGAGATGACCAAGAACCAGGTGAG CCTGACCTGCC'TGGTCAAAAGGCTTC TATCCCAGCGACATCGCCGTGGAG TGGGAGAGCAATGGCGAGCCGGAG AACAAC'TACAAGACCACGCCTCCC GTGCTGGACTCCGACGGCTCCTTCT TCCTCTATAGCAAGCTCACCGTGGA CAAGAGCAGGTGGCAGCAGGGGA ACGTCTTCTCATGCTCCGTGATGCA TGAGGCTCTGCACAACCAC'TACAC GCAGAAAGAGCCTCTCCCTGCTCCG GGTGGTGGCGATCGGGAGGTGGC GGATCCGAGGTGCAGCTGGTCGAG TCTGGAGGAGGCTTGATCCAGCCT GGGGGGTCCCTGAGACTCTCCTGT GCAGTCTCTGGGTTCAACCGTCAAGTA GCAAGTTTCATGACTGGGTCCGCC AGGCTCCA GGGAAAGGGCTGGAGT GGGTGTCA GTTATTTATAGCGGTGG TAAGACATACTACGCAGACTCCGT GAAAGGCCGATTCAACCATCTCCAG AGACAA'TTCCAAGAACACCGTGTGA TCTTCAAATGAACAGCCTGAGAGC CGAGGACACGGCCGTGTATTA'CTG TGC'GAGAGATAGCGGTGGCTGGGG
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SS-3086 2				GTACTTTGACTACTGGGGCCAGGG AACCTGGTCAACCGTGCTCCTCAGCA AGCACGAAGGGGCCGTCCGTATTT CCGCTTGCGCCCTCGTCGAAAGTCAA CTTGGGAGGGACCGCGGCACCTTG GCTGTCTTGTCAAAGATTACTTCCC TGAGCCAGTGACAGTCAGCTGGAA TTCCGGTGCCCTCACGTCAGGAGTA CATACTTCCCTGCGGTATTGCAGT CCTCCGGACTCTACTCCCTGGAGTC GGTGTTAACGGTGCCCCAGCTCCAG CTTGGGACCCAGACGTACATTGT AACGTGAATCACAAACCAAGCAAT ACTAAGGTAGATAAGAAAGTAGAA CCGAAGAGCTGC	(SEQ ID NO: 493)	(SEQ ID NO: 494)	(SEQ ID NO: 495)
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[illegible]

301

302

303

	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	EVQLLESGGGLVQPGGSLRLSCAAS GFTFSRNAMSWVRQAPGKGLIEWVS ATGGSGITYYADSVKGRFTISRDN KNTLYLQMNSLRAEDTAVYYCARG YSNSWYFDYWGGTLVTVSSAST KGPSVFLAPSSKSTGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFP AVLQSSGLYSLKSVTVTPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDK THTCPPCPAPELLGGPSVFLFPPKPKD TLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAKTKPCEEQYGGST YRCVSVLTVLHQDWLNGKEYKCKV SNKALPAPIEKTISKAKGQPREPQVY TLPPSRREEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTTPPVLD DGSFFLYSKLTVDKSRWQQGNVFS SVMHEALHNHYTQKSLSLSPGGGGS GGGGSQVQLQESGPGLVKPSSETLSLT CTVSGGSISSSSYYWGWIRPPGKGL EWIGSIYYSGITNYPNPSLKSRTISVD TSKNQFSLKLSSTAAADTAVYYCAR SSNYDAFDIWGGQGTMTVTVSSASTKG PSVFLAPSSKSTGGTAALGCLVKD YFPEPVTVSWNSGALTSGVHTFP LQSSGLYSLESVTVTPSSSLGTQTYIC NVNHKPSNTKVDKKVEPKSC	
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 503)	(SEQ ID NO: 504)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
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	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
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	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
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	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDPDRFSGSKSGTSASLAI GLQAEDEADYYCQSYDSSLGGWVF GGGTKLTVLPQKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKKYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS	(SEQ ID NO: 502)	(SEQ ID NO: 503)
	QVLTQPPSVSGAPGQR		

			CAGTCTGCCCTGACTCAGCCTGCCTC CGTGTCTGGAGCCCTGGACAGTCGA TCACCATCTCTGCACCTGGAACCA AGTGATGTTGGGAATTATAACCTTGT CTCCTGTTACCAACAGCACCCAGGCA AAGCCCCAAACTCATGATTTTGGAG GTCAATCAGCGGCCCTCAGGGGTTTC TAATCGCTTCTCTGGCTCCAAGTCTG GCACCACGGCTCCTGACAAATCTCT GGGCTCCAGGCTGCGGACGAGGCTG ATTATTTCTGCTGCTCATATACAAC AGTAGCACTTATGTGATCTTCGGCGG AGGGACCAAGCTGACCGTCTCTAGGT CAGCCCAAGGCTGCACCCCTCGGTAC TCTGTCCCCGCCCTCTCTGAGGAGC TTCAAGCCCAACAAGGCCACACTGGT GTGTCTCATCAGTGACTTCTACCCGG GAGCCGTGACAGTGGCCTGGAAGGC AGATAGCAGCCCCGTCAAGCGGGGA GTGGAACCAACCAACCCCTCCAAC AAAGCAACAACAAGTACGCGGCCGA AAGCTATCTGAGCCTGACGCTGAGC AGTGGAAAGTCCCACAGAAAGCTACAG CTGCCAGGTCACGCATGAAGGGAGC ACCGTGGAGAAGACAGTGGCCCCCTA CAGAAATGTTCA		GACATTACAGATGACCCAGTCTCCAT CTTCCGTGTCTGCATCTGTAGGGGA CAGATCACCATCACTTGTCTGGGC GAGTCAGGTATTACAGGTGGTT AGCCTGGTATCAGCAGAAACCAGG GAAAGCCCCTAAGTCTCTGATCTAT GTGCATCCGTTTTCGAAAGTGGG GTCCCATCAAGGTTACAGCGGCAGT GGATCTGGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAATC AAACGGACGGTGGCTGCACCATCT GTCTTCACTTCCCGCCACTCTGATG AGCAGTTGAAATCTGGAACCTGCCT CTGTTGTGTGCTGCTGAATAACTTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACCTCCAGGAGAGTGT CACAGACAGGACAGCAAGGACA GCACCTACAGCCTCAAGAGCACCC TGACGCTGAGCAAAAGCAGACTACG AGAAACACAAGTCTACGCCCTGCG AAGTACCCCATCAGGGCCTGAGCT CGCCCGTCAAAAGAGCTTCAACA GGGGAGAGTGT	GAGTGCAGCTGCTGGAGTCTGGG GGAGGTTGTACAGCCTGGGGG TCCCTGAGACTCTCCTGTGCAGCCT CTGGATTCACCTTATAGTAGAAATGC CATGAGTTGGTCCGCCAGGCTCC AGGAAAGGGCTGGAGTGGGTGTC AGCTACTGGGTAGTGGTATTAG CACATACTACGACACTCCGTGAA GGCCCGTTCAACATCTCCAGAGA CAATTCAGAAACACGCTGTATCT GCAAATGAACAGTCTGAGAGCCGA GGACACGGCCGTATATTACTGTGC GAGAGTTATAGCAACAGCTGGTG GTACTTGTACTACTGGGGCCAGGG AACCCTGGTCAACGCTGCTCTCAGCC TCCACCAAGGCCCATCGGTCTTCC CCCTGGCACCTCTCTCCAAAGACA CCTCTGGGGCACAGCGGCCCTGG GCTGCTGGTCAAGGACTACTTCCC CGAACCGGTGACGGTGTCTGTGGAA CTCAGGCGCCTGACCAGCGGCGT GCACACCTTCCCGGCTGTCTACAG TCCTCAGGACTCTACTCCCTCAAGA GCGTGGTGACCGTGGCCCTCCAGCA GCTTGGCACCCAGACCTACATCT GCAACGTGAATCACAAAGCCAGCA ACACCAAGTGGACAAAGAAAGTTG AGCCCAATCTTGTGACAAAATC ACACATGCCACCGTGCCAGCAC CTGAACCTCTGGGGGACCGTCTCAG TCTTCTCTTCCCCCAAAACCCAA GGACACCTCATGATCTCCCGGAC CCCTGAGGTACATGCTGGTGGT GGACGTGAGCCACGAAGACCCCTGA GGTCAAGTTCAACTGGTACGTGGA					
i	21- P	230 S	30 A12	: Ig G ₂	5 7	6 2	233 6	anti- 4H 6	anti- hu 9	Mes othel in>	4H6 VH1: :huI gG1 z- CH1 - E::E PKS C+

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	QSALETQASVSGSPGQITISCTGTSSD VGNYNLVSWYQQHPGKAPKLMIFEVN QRPSGVSNRFGSGKSGTTASLTISGLQA ADEADYFCCSYTTSSYVIFGGGTKLT VLGQPKAAPSVTLFPPSSEELQANKAT LVCLISDFYPGAVTVAWKADSSPVKA GVETTTPSKQSNNKYAAESYLSLTPEQ WKSHRSYSCQVTHEGSTVEKTVAPTE CS	DIQMTQSPSSVSASVGDRVTITCRAS QGITRWLAWYQQKPGKAPKLLIYAA SVLQSGVPSRFGSGSGTDFTLTISSL QPEDFATYYCQQSNFPRTFGQGTK VEIKRTVAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNAL QSGNSQESVTEQDSKDSITYSLKSTLT LSKADYEKHKVYACEVTHQGLSSPV TKSFNRGEC	EVQLLESGGGLVQPGGSLRLSCAAS GFTFSRNAMSWVRQAPGKGLEWVS ATGSGISTYYADSVKGRFTISRDN KNTLYLQMNSLRAEDTAVYYCARG YSNSWYFDYWGGTLVTVSSAST KGPSVFPLAPSSKSTSGGTAALGCLV KDYFPEPVTVSWNSGALTSGVHTFP AVLQSSGLYSLKSVTVTPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSCDK THTCPCPAPPELLGGPSVFLFPPKPKD TLMISRTPEVTCVVVDVSHEDPEVKF NWYVDGVEVHNAAKTKPCEEQYGST YRCVSVLTVLHQDWLNGKEYKCKV SNKALPAPIEKTISKAKGQPREPQVY TLPPSRREEMTKNQVSLTCLVKGFYPS DIAVEWESNGQPENNYKTTTPVLD DGSFFLYSKLTVDKSRWQQGNVFS SVMHEALHNHYTQKSLSLSPGGGGS GGGGSQVQLVESGGGLVKPGGSLRL SCAASGFTFSDYYMTWIRQAPGKGL EWISYISSSGSTIYADSVKGRFTISR DNAKNSLYLQMNSLRAEDTAVYYC ARDRNSHFDYWGGTLVTVSSASTK GPSVFPLAPSSKSTSGGTAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPA VLQSSGLYSLEISVTVTPSSSLGTQTYI CNVNHKPSNTKVDKKVEPKSC		
A			(SEQ ID NO: 508)	(SEQ ID NO: 509)	(SEQ ID NO: 510)
A					

[illegible]

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[illegible]

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3086 7				TGCTTTTGATATCTGGGGCCAAAGGG ACAATGGTCACCGTGTCTCAGCA AGCACGAAAGGGCCGTCGGTATTT CCGCTTGCGCCCTCGTCGAAAGTCAA CTTCGGGAGGGACCGCGGCACCTTG GCTGTCTTGTCAAAGATTACTTCCC TGAGCCAGTGACAGTCAGCTGGAA TTCCGGTGCCCTCACGTCAGGAGTA CATACTTCCCTGCGGTATTGCACT CCTCCGGACTCTACTCCCTGGAGTC GGTGTTAACGGTGCCCCAGCTCCAG CTTGGGGACCCAGACGTACATTGT AACGTGAATCACAAACCAAGCAAT ACTAAGGTAGATAAGAAAGTAGAA CCGAAGAGCTGC	(SEQ ID NO: 523)	(SEQ ID NO: 524)	(SEQ ID NO: 525)
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[illegible]

i	21- 230 P	21- 33 S	5 : H6 IgG	5 7 21- 6 3 4H 6 Fab	hu anti- <hu CD4 0> 21- 230- 33H 6 VHJ: :hul gG1 zSE FL2 *GK - K:-(G4S 2):-] hu anti- <hu Mes othel in> 4H6 VHJ: :hul gG1 z- CH1 - E::E PKS C+	GACATCCAGATGACCCAGTCTCCATC CTCCCTGTCTGCATCTGTAGGAGACA GAGTCACCATCACTTGCCGGCAGGT CAGAACATTAGCAGGCATTTAAATTG CTATAGCAGCATCCAGGGAAGCC CCTAAGGTCCTGATCCATCTGCATC CAGTTTGCCAAAGTGGGTCCCGTCAA GGTTCAAGTGGCAGTGGAATCTGGGAC AGATTTCAAGTCTTACCATCAGCAGTC TGCAACCTGAAGATTTTGGAACTTAC TTCTGTCAACAGAGTTACAGTACCCC TCCCACTTTCGGCGGAGGGACCAAG GTGGAGCTCAAACGAACGGTGGCTG CACCATCTGTCTTCAATCTCCCGCCAT CTGATGAGCAGTTGAAATCTGGAACT GCCTCTGTTGTGCTGCTGCTGAATAA CTTCTATCCAGAGAGGCCAAAGTAC N A AGTGGAAAGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC ACAAAGTCTACGCTGCGAAGTCAAC CATCAGGGCCTGAGCTCGCCCGTCA AAAGAGCTTCAACAGGGGAGAGTG GACATTCAGATGACCCAGTCTCCAT CTCCGTGTCTGCATCTGTAGGGA CAGATCACCATCACTTGTCGGC GAGTCAGGTATTACCAAGTGTT AGCTGGTATCAGCAAAACAGG GAAAGCCCTAAGCTCTGATCTAT GCTGATCCGTTTTCGAAAGTGG GTCCCATCAAGGTTCAAGGCGCAT GGATCTGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAAATC AAACGGACGGTGGTGCACCATCT GTCTTCATCTTCCCGCCATCTGATG AGCAGTTGAAATCTGGAAGTGCCT CTGTTGTGTGCTGTGTAATACTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACTCCCAAGGAGAGTGTC ACAGAGCAGGACAGCAAGGACAGCA CCTACAGCCTCGAAAGCACCTTGACG CTGAGCAAGAGCAGACTACGAGAAAC
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3086 8				TTTGACTATTGGGGCCAGGGAACC CTGGTCACCGTGTCTCAGCAAGC ACGAAGGGGCCGTCCGTATTTCGG CTTGCGCCCTCGTCGAAAGTCAACTT CGGAGGGACCGCGGCACTTGGCT GTCTTGTCAAAGATTACTTCCCTGA GCCAGTGACAGTCAGCTGGAATTC CGGTGCCCTCACGTCAGGAGTACA TACATTCCCTGCGGTATTGCAGTCC TCCGGACTCTACTCCCTGGAGTCGG TGGTAACGGTGCCCAAGCTCCAGCTT GGGGAACCCAGACGTACATTGTAA CGTGAATCACAAACCAAGCAATAC TAAGGTAGATAAGAAAAGTAGAACC GAAGAGCTGC	(SEQ ID NO: 529)	(SEQ ID NO: 530)	(SEQ ID NO: 531)
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[illegible]

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SS-30870				GTACTTTGACTACTGGGGCCAGGG AACCTGGTCACCGTGTCTCTCAGCA AGCACGAAGGGGCCGTCCGTATTT CCGCTTGCGCCCTCGTCGAAAGTCAA CTTGGGAGGGACCGCGGCACCTTG GCTGTCTTGTCAAAGATTACTTCCC TGAGCCAGTGACAGTCAGCTGGAA TTCCGGTGCCCTCACGTCAGGAGTA CATACTTCCCTGCGGTATTGCAGT CCTCCGGACTCTACTCCCTGGAGTC GGTGTTAACGGTGCCCCAGCTCCAG CTTGGGACCCAGACGTACATTGT AACGTGAATCACAAACCAAGCAAT ACTAAGGTAGATAAGAAAGTAGAA CCGAAGAGCTGC	(SEQ ID NO: 541)	(SEQ ID NO: 542)	(SEQ ID NO: 543)
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		DIQMTQSPSSLASVGDRTITCRAGQ NISRHLNWYQQNPGKAPKVLHPASSL PSGVPSRFGSGGTDLSLTISLQPEDF GTYFCQQSYSTPTFGGGTKVELKRTV AAPSVFIFFPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVT EQDSKDSYSTLESSTLTLSEADYEKHKV YACEVTHQGLSSPVTKSFNRGEC	DIVMTQSPDSLAVSLGERATINCKSS QSVLYSSNNKNYLAWYQQKPGQPP KLLIYWASTRESGVDPDRFSGSGGTD FTLTISLQAEDVAVYYCQQYYSTPP TFGGTKVEIKRTVAAPSVFIFFPSDE QLKSGTASVVCLLNNFYPREAKVQW KVDNALQSGNSQESVTEQDSKDSY SLKSTLTLSEADYEKHKVYACEVTH QGLSSPVTKSFNRGEC	QVQLVQSGAEVKKPGASVKVSCKAS GYTPGYYMYWLRAPGGQGLEWM GWINPDSGDTNYAQKFQGRVTMTTR DTSISTAFMELSRLRSDDTAVYYCAR EKPRYFDSFYFYLMDVWGQGTTVT VSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPTVSWNSGALTS GVHTFPAVLQSSGLYSLSKSVTVTPSS SLGTQTYICNVNHNKPSNTKVDKKVE PKSCDKHTHTCPCPAPELLGGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNG KEYCKKVSINKALPAPIEKTISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCL VKGFYPSDIAVEWESNGQPENNYKT TPPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMEALHNHYTQKSLSL SPGGGGGGGGSEVQLVESGGGLIQP GGSLRLSCAVSGFTVSSKFMWVRQ APGKLEWVSVIYSGGKTYADSVK GRFTISRDNKNTLYLQMNSLRAEDT AVYYCARDSDGGWGYFDYWGGGTL VTVSSASTKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPTVSWNSGALT SGVHTFPAVLQSSGLYSLSKSVTVTPS SSLGTQTYICNVNHNKPSNTKVDKKV EPKSC	(SEQ ID NO: 544)	(SEQ ID NO: 545)	(SEQ ID NO: 546)

			CAGGCTGTGCCGACTCAGCCCTCTTC CCTCTCTGCATCTCTCTGGAGCATCAG CCAGTCTCACTTGCACCTTACGCAGT GGCATCAATGTTGGTTCCTCCAGGAT CTATTGGTACCAGAGAACCCAGGG AGTCTCCCCAGTTTCTCTGAGGTA CACATCAGACTCAGATAAATTGCAG GGCTCTGGAGTCCCCAGCCGCTTCTC TGGATCCAAAGATGCTTCGGCCAATG CAGGACTTTTACTCATCTCTGGGCTC CAGTCTGAGGATGAGGCTGACTATTA CTGTATGATTTGGCACAGCAGCGCTG TGGTATTCCGGCGGAGGGACCAAACT GACCGTCTTAGGTACGCCAAGGCTG CACCTCGGTCACTCTGTTCCCGCCC TCCTCTGAGGAGCTTCAAGCCAAACA GGCCACACTGGTGTCTCATCAGTG NACTTCTACCCGGGAGCGGTGACAGTG GCCTGGAAGGCAGATAGCAGCCCCG TCAAGCGGGAGTGGAACCAACCAC ACCTCCAAACAAGCAACAACAAG TACGCGGCCGAAAGCTATCTGAGCCT GACGCTGAGCAGTGGAAATCCAC AGAAAGCTACAGTGCCAGGTACGC ATGAAGGGAGCACCGTGGAGAAGAC AGTGGCCCTACAGAATGTTCA		CAGTCAAGTGTGACGCAAGCGCCC TCAGTGTCTGGGCCCCCAGGGCAG AGGTCAACCATCTCTGCACCTGGG AGCAGCTCCAACATCGGGCAGGT TATGATGTTCACTGTTACAGCAG GTTCCAGGAACAGCCCCAAACT CTCATCTATGGTAACAGAAAGCG CCCTCAGGGTCCCTGACCGATTCT CTGGCTCCAAGTCTGGCACTCAGC CTCCCTGGCCATCACTGGGCTCCAG GCTGAGGATGAGGCTGATTATTAC TGCCAGTCTATGACAGCAGCCTG GGTGTGGGTGTTCCGCGGAGGG ACCAAGCTGACCGTCTTACAGCCC AAGGTGCACTCTGGTCACTCTGT TCCCGCCCTCTCTGAGGAGCTTCA AGCCAACAAGGCCACACTGGTGTG TCTCATCAGTGACTTCTACCCGGGA GCCGTGACAGTGGCTGGAAAGCA GATAGCAGCCCGTCAAGCGGGGA GTGGAACCAACACACCTCCAAA CAAGCAACAACAAGTACGCGGCC AAGAGCTATCTGAGCTGACGCT GAGCAGTGGAAAGTCCCAACAAGC TACAGTGGCAGGTACGCATGAA GGGAGCACCGTGGAGAAGACAGTG GCCCTACAGAATGTTCA		[hu anti- <hu CD4 0> 21- 230- 33H- 9 VH]; :hul gG1 zSE FL2 *GK - K:-(G4S)2::[hu anti- <hu Mes othel in> 4G1 2 VH]; :hul gG1 z- CH1 - E::E PKS
i	21- 230 H9- 33 IgG- 21- 233 4G 12- Fa b	P S : 5 7 6 3 2 2					

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3087 1				TGCTTTTGATATCTGGGGCCAAAGGG ACAATGGTCACCGTGTCTCAGCA AGCACGAAAGGGCCGTCGGTATTT CCGCTTGCGCCCTCGTCGAAAGTCAA CTTCCGGAGGGACCGCGGCACCTTG GCTGTCTTGTCAAAGATTACTTCCC TGAGCCAGTGACAGTCAGCTGGAA TTCCGGTGCCCTCACGTCAGGAGTA CATACTTCCCTGCGGTATTGCAGT CCTCCGGACTCTACTCCCTGGAGTC GGTGTTAACGGTGCCCCAGCTCCAG CTTGGGGACCCAGACGTACATTGT AACGTGAATCACAAACCAAGCAAT ACTAAGGTAGATAAGAAAGTAGAA CCGAAGAGCTGC	(SEQ ID NO: 547)	(SEQ ID NO: 548)	(SEQ ID NO: 549)
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[illegible]

			CAGGCTGTGCCGACTCAGCCCTCTTC CCTCTCTGCATCTCTCGAGCATCAG CCAGTCTACCTGCACCTTACGCAGT GGCATCAATGTTGGTCTCTCCAGGAT CTATTGGTACCAGCAGAACGCCAGGG AGTCCTCCCCAGTTTCTCTGAGGTA CACATCAGACTCAGATAAATTGCAG GGCTCTGGAGTCCCCAGCCGTTCTC TGGATCCAAAGATGCTTCGGCCAATG CAGGACTTTTACTCATCTCTGGGCTC CAGTCTGAGGATGAGGCTGACTATTA CTGTATGATTTGGCACAGCAGCGCTG TGGTATTCCGGCGGAGGGACCAAACT GACCGTCTAGGTACGCCAAGGCTG CACCCCTCGGTCACTCTGTTCGCCGCC TCCTCTGAGGAGCTTCAAGCCCAACAA GGCCACACTGGTGTGTTCTCATCAGTG ACTTCTACCCGGGAGCCGTGACAGTG GCCTGGAAGGCAGATAGCAGCCCCG TCAAGCGGGAGTGGAACCAACCAC ACCTCCAAACAAGCAACAACAAG TACGCGGCCGAAAGCTATCTGAGCCT GACGCTGAGCAGTGGAAATCCAC AGAAGCTACAGTGCCAGGTACGC ATGAAGGGAGCACCGTGGAGAAGAC AGTGGCCCTACAGAAATGTTCA		GACATTACAGATGACCCAGTCTCCAT CTTCCGTGTCTGCATCTGTAGGGGA CAGATCACCATCACTTGTCCGGGC GAGTCAGGTATTACAGGTGGTT AGCCTGGTATCAGCAGAAACCAGG GAAAGCCCCTAAGCTCTGTATCTAT GCTGCATCCGTTTTCGAAAGTGGG GTCCCATCAAGGTTACGCGGCAAT GGATCTGGGACAGATTTCACTCTCA CCATCAGCAGCCTGCAGCCTGAAG ATTTTGCAACTTACTATTGTCAACA GTCTAACAGTTTCCCTCGGACGTTT GGCCAAAGGACCAAGGTGGAATC AAACGGACGGTGGCTGCACCATCT GTCTTCACTTCCCGCATCTGATG AGCAGTTGAAATCTGGAACCTGCCT CTGTTGTGTGCTGCTGAATAACTTT CTATCCAGAGAGGCCAAAGTACA GTGGAAGGTGGATAACGCCCTCCA ATCGGTAACCTCCAGGAGAGTGT CACAGCAGGACAGCAAGGACA GCACCTACAGCCTCAAGAGCACCC TGACGCTGAGCAAAAGCAGACTACG AGAAACACAAGTCTACGCCCTGCG AAGTACCCATCAGGGCCTGAGCT CGCCCGTCACAAAGAGCTTCAACA GGGGAGAGTGT		CAGGTGACAGTTGGTGGAGTCTGGG GGAGGGTGGTCCAGCCTGGGAGG TCCCTGAGACTCTCTGTGCAGCGT CTGGATTCACTTCACTAGTACCATGG CATGCACTGGTCCGCCAACCTCC AGGCAAGGGCTGGAGTGGTGGC AGTTACTGTATGATGGAAGTAA TGAATACTATGGAGACTCCGTGAA GGCCGATTCACCATCTCCAGAGA CAATTCCAAGAAACACGCTGTATCT GCAATGAACAGCCTGAGAGTCGA GGACACGGCTGTGTATTACTGTAC GAGAGGGGGGGCCACTGGAACATA CGAGGGCCACTACTATGATGGA CGTCTGGGGCCAAAGGACCCACGGT CACCGTCTCTCAGCCTCCACCAAG GGCCCATCGTCTTCCCCCTGGCAC CCTCTCCAAGAGCACTCTGGGG GCACAGCGGCCCTGGGCTGCCTGG TCAAGGACTACTTCCCCGAACCCG TGACGGTGTCTGGAACCTCAGGCG CCCTGACCAGCGGCGTGCACACCT TCCCAGCTGTCTACAGTCTCAGG ACTCTACTCCCTCAAGAGCGTGGTG ACCGTGCCCTCCAGCAGCTTGGGC ACCCAGACCTACATCTGCAACGTG AATCACAAGCCAGCAACACCAAG GTGGACAAGAAAGTTGAGCCCCAAA TCTTGTGACAAAACTCACACATGCC CACCGTCCACAGCCTGAACCTCCT GGGGGACCGTCAGTCTTCTCTCTTC CCCCAAACCCCAAGGACACCCCTC ATGATCTCCCGGAGCCCTGAGGTC ACATGCGTGTGGTGACGTGAGC CACGAAGACCCTGAGGTCAAGTTC
i	21- P	anti- hu CD4 0> 21- 230- 33H- 9 VHJ; :huI gG1 zSE FL2 *GK - H9- IgG 5 7 233 3 6 2 7	21- 230- 33H- 9 VHJ; :huI gG1 zSE FL2 *GK - H9- IgG 5 7 233 3 6 2 7	anti- hu CD4 0> 21- 230- 33H- 9 VHJ; :huI gG1 zSE FL2 *GK - H9- IgG 5 7 233 3 6 2 7	anti- hu CD4 0> 21- 230- 33H- 9 VHJ; :huI gG1 zSE FL2 *GK - H9- IgG 5 7 233 3 6 2 7		

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[illegible]

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		QAVPTQPSSLSASPGASASLTCTLRSGI NVGSSRIYWYQQKPGSPQFLRLRYTSD SDKLQSGVPSRFSGSKDASANAALLL ISGLQSEDEADY YCMIWHSSAVVFGG GTKLTVLGQPKAAPSVTLFPPSSEELQ ANKATLVCLISDFYPGAVTVAWKADS SPVKAGVETTPSKQSNNKYAAESYLS LTPEQWKSHRSYSCQVTHEGSTVEKT VAPTECS	DIQMTQSPSSVSASVGDRVTITCRAS QDISRWLAWYQQKPGKAPKLLISAA SRLQSGVPSRFSGSGTDFTLTISL QPEDFAIYYCQAKSPRTFGQGTKV EIKRTVAAPSVFIAPPDEQLKSGTAS VVCLNNFYPREAKVQWKVDNALQ SGNSQESVTEQDKDSTYSLKSTLT SKADYEKHKVYACEVTHQGLSSPVT KSFNRGEC	QVQLVESGGGVVQPGSRSLRLSCAAS GFTSSHGHWVRQPPGKGLEWVA VIWYDGSNEYGDSVKGRFTISRDN SKNTLYQMNSLRVEDTAVYYCTR GGHWNYEGHYYGMDVWVGQGTITVT VSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPVTVSWNSGALTS GVHTFPAVLQSSGLYSLKSVVTPSS SLGTQTYICNVNHKPSNTKVDKKEVE PKSCDKTHTCTPCPAPPELLGGPSVFL FPPKPKDTLMISRTPEVTCVVDVSH EDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNG KEYCKKVSNAKALPAIEKTIKAKGQ PREPQVYTLPPSREEMTKNQVSLTCL VKGFYPSDIAVEWESNGQPENNYKT TPPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMEALHNHYTQKSLSL SPGGGGGGGGQVQLVESGGGLVK PGGSLRLSCAASGFTFSDYVMSWIRQ APGKLEWISYISSSESIYYVDSVKG RFTISRDNAKNSLYQMNSLRAEDT AVYYCARDVGSFHDYWGQGTILVTV SSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSKV HTFPAVLQSSGLYSLKSVVTPSSSL GTQTYICNVNHKPSNTKVDKKEVEPK SC	(SEQ ID NO: 562)	(SEQ ID NO: 563)	(SEQ ID NO: 564)
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QAVPTQPSSLSASPGASASLTCTLRSGI NVGSSRIYWYQQKPGSPQFLRLRYTSD SDKLQSGVPSRFSKSDASANAAGLL ISGLQSEDEADYYCMIWHSSAVVFGG GTKLTVLGQPKAAPSVTLFPPSSEELQ ANKATLVCLISDFYPGAVTVAWKADS SPVKAGVETTTTPSKQSNNKYAAESYLS LTPEQWKSHRSYSCQVTHEGSTVEKT VAPTECS	DIVMTQSPDSLAVSLGERATNCKSS QSVLYSSNNKNYLAWYQQKPGQPP KLLIYWASTRESGVDRFSGSGSGTD FTLTISLQAEDVAVYYCQYYSTPP TFGQGTKVEIKRTVAAPSVFIFFPPSDE QLKSGTASVVCLLNNFYPREAKVQW KVDNALQSGNSQESVTEQDSKDSY SLKSTLTLSKADYEKHKVYACEVTH QGLSSPVTKSFNRGEC	QVQLVESGGGVVQPGRSRLRLSCAAS GFTFSHGMHWVRQPPGKGLEWVA VIWYDGSNEYGDSVKGRFTISRDN SKNTLYLQMNLSRVEDTAVYYCTR GGHWNYEGHYGMDVWGQGTITV VSSASTKGPSVFPLAPSSKSTSGGTA ALGCLVKDYFPEPTVSWNSGALTS GVHTFPAVLQSSGLYSLKSVVTPVPS SLGTQTYICNVNHNKPSNTKVDKKVE PKSCDKTHTCTPCPAPELLGGPSVFL FPPKPKDTLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNG KEYCKKVSINKALPAIEKTIISKAKGQ PREPQVYTLPPSREEMTKNQVSLTCL VKGFYPSDIAVEWESNGQPENNYKT TPPVLDSDGSFFLYSKLTVDKSRWQ QGNVFSCSVMEALHNHYTQKSLSL SPGGGGGGGGSEVQLVESGGGLIQP GGSRLSCAVSGFTVSSKFMWVRQ APGKLEWVSVIYSGGKTYADSVK GRFTISRDNKNTLYLQMNSLRAEDT AVYYCARDSDGGWGYFDYWGGGTL VTVSSASTKGPSVFPLAPSSKSTSGGT AALGCLVKDYFPEPTVSWNSGALT SGVHTFPAVLQSSGLYSLKSVVTPVPS SSLGTQTYICNVNHNKPSNTKVDKKV EPKSC	(SEQ ID NO: 568)	(SEQ ID NO: 569)	(SEQ ID NO: 570)
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A					

[illegible]

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		<p>QVQLVESGGGVVQPGRLSLRSCAAS GFTLSSYGMHWVRQAPGKGLEWVA VIWYDGSNKYYADSVKGRVTISRDN SKNTLYQMNSLR AEDTAVYYCTRD GRNYYVFDNWRGQGTLLVTVSSATK GPSVFLPAPSSKSTGGTAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPA VLQSSGLYSLKSVTVVPSSSLGTQTY ICNVNHPKSTNTKVDKKVEPKSCDKT HTCTPPCPAPELLGGPSVFLFPPKPKDT LMISRTPEVTCVVVDVSHEDPEVKFN WYVDGVEVHNAKTKPCEEQYGSTY RCVSVLTVTLHQDWLNGKEYCKCKVS NKAALPAPIEKTISKAKGQPREPQVYV LPPSREEMTKNQVSLTCLVKGFYPSD IAVEWESNGQPENNYKTTPPVLDSD GSFFLYSLKLTVDKSRWQQGNVFCSS VMHEALHNHYTQKSLSLSPGSGGSG GGGSQLQESGPGLVKPSETLSLTC TVSGGSISSSSYYWGWRQPPGKGLE WIGSIYSGITDYNPSLKSRVTISVDIT SKNQFSLKLSVTAADTAVYYCAR SNYDAFDIWWGQGTMTVTVSSASTKGP SVFPLAPSSKSTGGTAALGCLVKDY FPEPVTVSWNSGALTSGVHTFPAVL QSSGLYSLKSVTVVPSSSLGTQTYICN VNHKPSNTKVDKKVEPKSC</p>	(SEQ ID NO: 576)
		<p>QSVLTQPPSVSGAPQRVTISCTGSSS NIGAGYDVHWYQQVPGTAPKLLIYG NSKRPSGVDRFSGSKSGTSASLAIT GLQAEDEADYYCQSDSSLGGWVF GGGTKLTVLQPKAAPSVTLFPPSSEE LQANKATLVCLISDFYPGAVTVAWK ADSSPVKAGVETTTPSKQSNKYYAA KSYLSLTPEQWKSHRSYSCQVTHEG STVEKTVAPTECS</p>	(SEQ ID NO: 575)
		<p>QSALTPQPRSVSGSPGQSVTISCTGTSSD VGGYIFVSWYQQHPGKAPKLMYDVS KRPSGVDRFSGSKSVNTASLTISGLQA EDETIDYYCCSYAGNYTVVFGTGTIKVT VLGQPKAAPSVTLFPPSSEELQANKAT LVCLISDFYPGAVTVAWKADSSPVKA GVETTTPSKQSNKYYAAESYLSLTPEQ WKSHRSYSCQVTHEGSTVEKTVAPTE CS</p>	(SEQ ID NO: 574)
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			<p>QVQLVESGGGVVQPGRLRLSCAAS GFTLSSYGMHWVRQAPGKGLEWVA VIWYDGSNKYYADSVKGRVITSRDN SKNTLYLQMNSLRAEDTAVYYCTR GRNYVYFDNWVGQGLT VTVSSATK GPSVFLAPSSKSTSGGT AALGCLVK DYFPEPVTVSWNSGALTSGVHTFPA VLQSSGLYSLKSVTVTPSSSLGTQTY ICNVNHPKPSNTKVDKKVEPKSCDKT HTCPPCPAPELLGGPSVFLPPKPKD LMISRTPEVTCVVVDVSHEDPEVKFN WYVDGVEVHNAKTKPCEEQYGSTY RCVSVLT VLVHQQDWLNGKEYCKV NKALPAPIEKTIKAKGQPREPQVY LPPSRHEMTKNQVSLTCLVKGFYPSD IAVEWEESNGQPENNYKTTPVLDSD GSFFLYSKLTVDKSRWQQGNVFCSS VMHEALHNHYTQKSLSLSPGSGSG GGGQVQLVESGGGLVKPGGSLRLS CAASGFTFSDYMTWIRQAPGKGLE WISYISSSGSTIYYADSVKGRFTISR NAKNSLYLQMNSLRAEDTAVYYCA RDRNSHFDYWGQGLT VTVSSASTKG PSVFLAPSSKSTSGGT AALGCLVKD YFPEPVTVSWNSGALTSGVHTFPAV LQSSGLYSLESVTVTPSSSLGTQTYIC NVNHPKPSNTKVDKKVEPKSC</p>	(SEQ ID NO: 580)	(SEQ ID NO: 581)	(SEQ ID NO: 582)
			<p>DIQMTQSPSSVSASVGDRVTITCRAS QGITRWLA WYQQKPGKAPKLLIYAA SVLQSGVPSRFSGSGTDFLTITISL QPEDFATYYCQQSNFPRITFGQGTK VEIKRTVAAPSVEIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWVKVDNAL QSGNSQESVTEQDSKDSYSLKSTLT LSKADYEKHKVYACEVTHQGLSSPV TKSFNRGEC</p>			
			<p>QSALTQPRSVSGSPGQSVTISCTGTSSD VGGYIFVSWYQQHPGKAPKLMYDVS KRPSGVPDFRFGSKSVNTASLTISGLQA EDETIDYYCCSYAGNYTYVFGTGTKVT VLGQPKAAPSVTLFPPSSEELQANKAT LVCLISDFYPGAVTVAWKADSSPVKA GVEITTPSKQSNNKYAAESYLSLTPEQ WKSRRSYSCQVTHEGSTVEKTVAPTE CS</p>			
			<p>A A</p>			

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		QVQLVESGGGVVQPGRLRLSCAAS GFTLSSYGMHWVRQAPGKGLEWVA VIWYDGSNKYYADSVKGRVTISRDN SKNTLYLQMNSLRAEDTAVYYCTRD GRNYVYFDNWGQGTLLVTSSASTK GPSFPLAPSSKSTSGGTAALGCLVK DYFPEPTVSWNSGALTSGVHTFPA VLQSSGLYSLKSVVTPSSSLGTQTY ICNVNHKPSNTKVDKKVEPKSCDKT HTCPPCPAPELGGPSVFLPPKPKDT LMISRTPEVTCVVVDVSHEDPEVKFN WYVDGVEVHNAKTKPCEEQYGSTY RCVSVLTVLHQDWLNGKEYKCKVVS NKALPAPIEKITISKAKGQPREPQVYT LPPSREEMTKNQVSLTCLVKGFYPSD IAVEWESNGQPENNYKTTTPVLDSD GSFFLYSKLTVDKSRWQQGNVFCSS VMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVESGGGLVKPGGSLRLS CAASGFTFSDYIMSWIRQAPGKGLE WISYISSESIYYVDSVKGRFTISRDN AKNSLYLQMNSLRAEDTAVYYCAR DVGSHFDYWGQGTLLVTSSASTKGP SVFPLAPSSKSTSGGTAALGCLVKDY FPEPTVSWNSGALTSGVHTFPAVL QSSGLYSLESVVTVPSSSLGTQTYICN VNHKPSNTKVDKKVEPKSC
		DIQMTQSPSSVSASVGDRVTITCRAS QDISRWLAWYQQKPGKAPKLLISAA SRLQSGVPSRFSGSGTDFTLTISSL QPEDFAIYYCQQAQKSPRTFGQGTKV EIKRTVAAPSVFIFPPSDEQLKSGTAS VVCLLNNFYPREAKVQWKVDNALQ SGNSQESVTEQDSKDSITYSLKSTLTL SKADYEEKHKVYACEVTHQQLSSPVT KSFNRGEC
		QSALTPQRSVSGSPGQSVTISCTGTSSD VGGYIFVSWYQQHPGKAPKLMYDVS KRPSGVPDFRFGSKSVNTASLTISGLQA EDETIDYCCSYAGNYTYVFGTGKVT VLGQPKAAPSVTLFPPSSEELQANKAT LVCLISDFYPGAVTVAWKADSSPVKA GVETTTPSKQSNINKYAAESYLSLTPEQ WKSHRYSQCQVTHEGSTVEKTVAPTE CS
	A	
	A	
		(SEQ ID NO: 586)
		(SEQ ID NO: 587)
		(SEQ ID NO: 588)

361

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30878				GGGCCAGGGAAACCCTGGTCACCGT GTCCTCAGCAAGCACGAAGGGGCC GTCCGTATTCCCGCTTGC GCCCTCG TCGAAGTCAACTTCGGGAGGGACC GCGGCAC TTGGCTGTCTTGTCAAAG ATTACTTCCCTGAGCCAGTGACAGT CAGCTGGAATTCCGGTGCCCTCAC GTCAGGAGTACATACATTCCCTGC GGTATTGCAGTCCCTCCGGACTCTAC TCCCTGGAGTCGGTGGTAACGGTG CCCAGCTCCAGCTTGGGGACCCAG ACGTACATTTGTAACGTGAATCAC AAACCAAGCAATACTAAGGTAGAT AAGAAAGTAGAACCCGAAGAGCTGC	(SEQ ID NO: 589)	(SEQ ID NO: 590)	(SEQ ID NO: 591)
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[illegible]

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[illegible]

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C + [anti - <hu CD4 0> 21- 230- 39C 2 VL]: :huL LC2 -E + [anti - <hu Mes othel in> 7G1 1 VL]: :huK LC- S176 K (IgG - Fab) ; LM R ID: SS-		TGGACGGCGTGGAGGTGCATAATG CCAAGACAAAGCCGTGCGAGGAGC AGTAGGGCAGCACGTACCGTTGGG TCAGCGTCTCACCGTCTCGACCA GGACTGGCTGAATGGCAAGGAGTA CAAGTGCAAGGTGTCCAACAAGC CCTCCAGCCCCCATCGAGAAAAC CATCTCAAAGCCAAAGGGCAGCC CCGAGAACACAGGTGTACACCT GCCCCATCCCGGAGGAGATGAC CAAGAACCAGGTCAAGCTGACCTG CCTGGTCAAAGGCTTCTATCCCAGC GACATCGCCGTGGAGTGGGAGAGC AATGGCAGCCGGAGAACAACTAC AAGACCAGCCTCCCGTCTGGAC TCCGACGGCTCCTTCTTCTCTATA GCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGAACTCTTCT CATGCTCCGTGATGCATGAGGCTCT GCACAACCACTACACGCAGAAAGAG CCTCTCCCTGTCTCCGGTGGTGGC GGATCGGGAGGTGGCGGATCCGAG GTGCAGCTGGTCGAGTCTGGAGGA GGCTTGATCCAGCCTGGGGGTCC CTGAGACTCTCTGTGCAGTCTCTG GGTTCACCGTCAGTAGCAAGTTCTAT GACCTGGGTCCGCCAGGCTCCAGG GAAGGGGTGGAGTGGGTGTCACT TATTTATAGCGGTGGTAAGACATA CTACGCAGACTCCGTGAAGGGCCG ATTACCATCTCCAGAGACAATTCC AAGAACACGCTGTATCTTCAAATG AACAGCCTGAGAGCCGAGGACACG GCCGTGTATTACTGTGCGGAGAGAT AGCGGTGGCTGGGGGTACTTTGAC
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411

[illegible]

			CAAGACAAAAGCCGTGCGAGGAGCAG TACGGCAGCACGTACCGTTGGGTCA GCGTCTCACCGTCTGCACCAAGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCCTGGTCAAAGC TTCTATCCCAGGACATCGCCGTGGA GTGGGAGACAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGTCAACGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGAGGTGGCGGATCCCA GGTGCAGCTGGTGCAGTCTGGGGCT GAGGTGAAGAACCTGGGGCCTCAG TGAAGGTGCTCTGCAAAGGCTTCTGGA TACACCTTCGCCGGCTACTATATGCA CTGGGTGCGACAGGCCCTGGACAA GGGCTTGAGTGGATGGGATGGATCA ACCTGACAGTGGAGGCACAAACTT TGCACAGCAGTTTCAGGGCAGGGTC ACCATGACCAAGGATACGTCCATCA GCACAGCCTACATGGAGGTGAGCAG GCTGAGATCTGACGACACGGCCGTG TTTTACTGTGCGAGAGAGAGATCA CTATGACTGGTATTTACTTTGACTAT TGGGGCCAGGGAACCTGGTCAACCG TGTCCTCAGCAAGCACGAAGGGGCC
23	0_	4G	7
V	Hj:	hu	lg
G1	z-	C	HI
-	E::	EP	KS
C	+	[an	ti-
<h	u	Me	sot
hel	in>	4G	12
VL	l::	hu	LL
C2	-E	+	

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		QSVLTQPPSVSGAPQQRVTISCTGSSSN IGAGYDVHWYQQVPGTAPKLLIYGNS KRPSGVDRFSGSGTSASL/ATGLQ AEDEADYYCQSYDSSLGGWVFGGT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNINKYAAESYLSLTP EQWKSHRSYSCQVTHEGSTVEKTVAP TECS	DIQMTQSPSSLSASVGDIITITCRASQNI TTYLNWYQQKPGKAPNLLISAAASRLRS GVPSRFGSGSGTDFLTITSLQPVDFT TFYCQQFTFTTPWTFGQGTKEIKRTVA APSVFIFFPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTE QDSKDSITYSLKSTLTLSKADYEKHKV YACEVTHQGLSSPVTKSFNRGEC	QVQLQESGPGLVKPSSETLSLTCTVSGG SISSSYWGWIRQPPGKGLEWIGSIYY SGITNYPNLSKSRVTISVDTSKNQFSLK LSSVTAADTAVYYCARPSNYDAFDIW GQGTMTVTSSASTKGPSVFPLAPSSKS TSGTAALGCLVKDYFPEPTVSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHPKPSNTKVDKK VEPKSCDKTHTCPPCPAPELLGGPSVF LFPPKPKDITLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVQSGAEVKKPGASVKVSC KASGYTFAGYIMHWVRQAPGQGLE WMGWINPDSGGTNFAQQFQGRVTMT RDTISITAYMEVSRLSRSDDTAVFYCAR EKITMTGIYFDYWGQGTLVTVSSASTK GPSVFPLAPSSKSTSGGTAAALGCLVKD YFPEPVTVSWNSGALTSGVHTFPAVLQ SSGLYSLESVTVTPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSC
				(SEQ ID NO: 670)
				(SEQ ID NO: 671)
				(SEQ ID NO: 672)

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			TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACCAGGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGTGGTG GCGGATCGGAGAGGTGGCGGATCCCA GGTGCAACTGGTGCAGTCTGGGGCT GAGGTGACGAAGCCTGGGGCCTCAG TGAAGGTGTCTGCAAGGCTTCTGGA TACACCTTCGCCGGCTACTATATGCA CTGGGTGCGACAGGCCCTGGACAA GGGCTTGAGTGGATGGATGGATCA ACCTCAGAGTGGTGGCACAACTA TGCACAGAAAGTTTCAGGACAGGGTC ACCATGACCAAGGACACGTCCATCA ACACAGCCTACATGGAAGTGAAGCAG GCTGAGATCTGACGACACGGCCGTG TATTACTGTGCGAGAGAAAGTATTTC TATGGTTCGGGGAGTCGGGCACAAC TGTTTCGCCCCCTGGGGCCAGGGA CCCTGGTACCGGTGCTCTCAGCAAGC ACGAAGGGGCCGTCCTGATTTCGCGCT
0	29	HI	0
V	Hj:	hu	lg
G1	z-	C	HI
-	E::	EP	KS
C	+	lan	ti-
<h	u	Me	sot
hel	in>	4G	12
VL	l::	hu	LL
C2	-E	+	

[illegible]

		QSVLTQPPSVSGAPQQRVTITISCTGSSSN IGAGYDVHWYQQVPGTAPKLLIYGNS KRPSGVPDFRFGSKSGTSASLAIITGLQ AEDEADYYCQSYDSSLGGWVFGGTT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNKYYAAESYLSLTP EQWKSHRSYSCQVTHEGSTVEKTVAP TECS	DIQMTQSPSSLSASVGDRVTITCRASQ DISNNLAWFQQKPGPKPKSLMYAASS LHSGVPSTFSGSGGTDFITISSLQPE DFATYYCQQNSYPLTFGGGTKVEIRR TVAAPSVFIFFPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSYSLKSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QVQLQESGPGLVKPSSETLSLTCTVSGG SISSSYWGWIRQPPGKGLEWIGSIYY SGITYNPSLKSRTVISVDTSKNQFSLK LSSVTAADTAVYYCARPSNYDAFDIW GQGTMTVTSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPTVSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHKPSNTKVDKK VEPKSCDKTHTCPPCPAPELLGGPSVF LFPPKPKDITLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVQSGAEVTKPGASVKVSC KASGYTFAGYIMHWVVRQAPGQGLE WMGWINPHSGGTNYAQKFQDRVTMT RDTISNTAYMELSRLSDDTA VYYCA RERISMVRGVGHNWFAPWGQGTLVLT VSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPTVSWNSGALTSGVH TFPAVLQSSGLYSLESVVTVPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSC (SEQ ID NO: 678)
				(SEQ ID NO: 677)
				(SEQ ID NO: 676)

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TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGTGACACGAGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCC TGCTGGACTCCGACGGCTCCTTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGGAGGTGGCGGATCCGA GGTGCAAGTCTGGAGTCTGGGGGA GGCTTGGTACAGCCTGGGGGTCCCT GAGACTCTCTGTGCAAGCCTCTGGAT TCACCTTTAGTAGAAATGCCATGAGT TGGGTCCGCCAGGCTCCAGGGAAGG GGCTGGAGTGGGTGTCAGCTACTGG TGGTAGTGGTATTAGCACATACTACG CAGACTCCGTGAAGGGCCGGTTTAC CATCTCCAGAGACAAATCCAAGAAC ACGCTGTATCTGCAAAATGAACAGTCT GAGAGCCGAGGACACGGCCGTATAT TACTGTGCGAGAGGTTATAGCAACA GCTGTGGTACTTTGACTACTGGGGC CAGGGAACCTTGGTCAACCGTGTCTC AGCAAGCACGAAGGGGCCGTCCGTA TTTCCGCTTGCCCTCTGTCGAAGTC																																		

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		QSVLTQPPSVSGAPQQRVTITISCTGSSSN IGAGYDVHWYQQVPGTAPKLLIYGNS KRPSGVPDFRFGSKSGTASLAIITGLQ AEDEADYYCQSYDSSLGGWVFGGTT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNINKYAAESYLSLTP EQWKSHRSYSCQVTHEGSTVEKTVAP TECS	QSQALTOPASVSGSPGQSIITISCTGTSSD VGNYNLVSWYQOHPGKAPKLMIFEV NQRPSGVNRFSGSKSGTTASLTISGLQ AADEADYFCCSYTTTSSTYVIFGGGTKL TVLGGQPKAAPSVTLFPPSSEELQANKA TLVCLISDFYPGAVTVAWKADSSPVK AGVETTTPSKQSNINKYAAKSYLSLTPE QWKSHRSYSCQVTHEGSTVEKTVAPT ECS	QVQLQESGPGLVKPSSETLSLTCTVSGG SISSSYWGWIRQPPGKGLEWIGSIYY SGITNYPNPSLKSRTVISVDTSKNQFSLK LSSVTAADTAVYYCARPSNYDAFDIW GQGTMTVTSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPTVSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHPKPSNTKVDKK VEPKSCDKTHTCPPCPAPELLGGPSVF LFPKPCKDTLMISRTPETCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGSEVQLLESGGGLVQPGGSLRLSC AASGFTFSRNAMSWVRQAPGKGLEW VSATGGSGISTYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCARG YSNSWVWFYDWGQGTLVTVSSASTK GPSVFPLAPSSKSTSGGTAAALGCLVKD YFPEPTVSWNSGALTSGVHTFPAVLQ SSGLYSLESVVTVPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSC (SEQ ID NO: 684)
				(SEQ ID NO: 683)
				(SEQ ID NO: 682)

425

426

427

QSVLTQPPSVSGAPGQRVTISCTGSSSN IGAGYDVHWYQQVPGTAPKLLIYGNS KRPSGVPDRFSGSKSGTSASLAITGLQ AEDEADYYCQSYDSSSLGGWVFGGQT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTTPSKQSNNKYAAESYLSLTP EQWKSHRYSYSCQVTHEGSTVEKTVAP TECS	DIQMTQSPSSLSASVGDRVTITCRAGQ NISRHLNHWYQQNPCKAPKVLIIHPASSL PSGVPSRFGSGSGTDFSLTISSLQPEDF GTFFCQQSYSTPPTFGGGTKVELKRTV AAPSVFIIPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVT EQDSKDSITYSLKSTLTLSKADYEKHK VYACEVTHQGLSSPVTKSFNRGEC	QVQLQESGPGLVKPSETLSLTCTVSGG SISSSYWGWIRQPPGKGLEWIGSIYY SGITNYPNPSLKSRTVISVDTSKNQFSLK LSSVTAADTAVYYCARPSNYDAFDIW GQGTMTVTSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPTVTSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHKPSNTKVDKK VEPKSCDKTHTCTPCPAPELLGGPSVF LFPPKPKDITLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTPPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGGGGSG GGGSQVQLVQSGAEVKKPGASVKVSC KASGYTPGYMYWLRQAPGQGLEW MGWINPDSGDTNYAQKFQGRVTMTR DTSISTAFMELSRLLSDDTAVYYCARE KPRYFDSFYYYLMDVWGQGTITVTVSS ASTKGPSVFPLAPSSKSTSGGTAALGC LVKDYFPEPTVTSWNSGALTSGVHTFP AVLQSSGLYSLESVTVTPSSSLGTQTYI CNVNHKPSNTKVDKKVEPKSC	(SEQ ID NO: 688)	(SEQ ID NO: 689)	(SEQ ID NO: 690)
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[hu ant i- <h u Me sol hel in> 4G 23 12 3 4 HJ: hu Ig G1 zS EF L2 L2 *G K- K:: (G H 4S 9) 2): F u ant i- <h uC D4 0> 21- 23	CAGTCAAGTGTGACGAGCCGCGCTC AGTGTCTGGGCCCCAGGCGAGAGG GTCACCATCTCTGCACTGGGAGCAG CTCCAACATCGGGCAGGTTATGAT GTTCACTGGTACCAGCAGGTTCCAGG AACAGCCCCAAACTCCTCATCTATG GTAACAGCAAGCGCCCTCAGGGGT CCCTGACCGATTCTCTGGCTCCAAGT CTGGCACCTCAGCCTCCCTGGCCATC ACTGGGCTCCAGGCTGAGGATGAGG CTGATTATTACTGCCAGTCTCTATGAC AGCAGCTGGGTGGTGGGTGTTTCG GCGGAGGACCAAGCTGACCGTCTCT ACAGCCCCAAGGCTGCACCCCTCGGTC ACTCTGTTCCCGCCCTCTCTGAGGA GCTTCAAGCCAAACAAGGCCACACTG GTGTGTTCTCATCAGTGACTTCTACCC GGGAGCCGTGACAGTGGCCTGGAAG GCAGATAGCAGCCCCGTCAAAGCGCG GAGTGGAAACCAACCAACCCCTCAA ACAAAGCAACAACAAGTACGCGGCC GAAAAGCTATCTGAGCCTGACGCGCTG AGCAGTGGAAATCCACAGAAAGCTA CAGCTGCCAGGTACGCATGAAGGG AGCACCGTGGAGAGACAGTGGCCC CTACAGAATGTTCA	CAGGCTGTGCCGACTCAGCCCTCTTC CCTCTTGCATCTCCTGGAGCATCAG CCAGTCTCACCTGCACCTTACGCAGT GGCATCAATGTTGGTCTCTCCAGGAT CTATTGGTACCAGCAAGAACCCAGGG AGTCTCCCAAGTTTCTCCTGAGGTA CACATCAGACTCAGATAAATTGCAG GGCTCTGGAGTCCCCAGCCGCTTCTC TGGATCCAAAGATGCTTCGGCCAAT GCAGGACTTTTACTCATCTCTGGGCT CCAGTCTGAGGATGAGGCTGACTATT ACTGTATGATTTGGCACAGCAGCGCT GTGGTATTCTGGCGGAGGGACCAAAC TGACCGTCTTAGGTACGCCCAAGGCT GCACCCCTCGGTCACTCTGTTCCCGCC CTCCTCTGAGGAGCTTCAAAGCCAACA AGGCCACACTGTGTGTCTCATCAGT GACTTCTACCCGGGAGCCGTGACAG TGGCCTGGAAAGCAGATAGCAGCCC CGTCAAGGCGGAGTGGAAACCAACC ACACCCCTCCAAACAAGCAACAACA AGTACGCGGCCAAGAGCTATCTGAG CCTGACGCTGAGCAGTGGAAAGTCC CACAGAAGCTACAGCTGCCAGGTCA CGCATGAAGGAGGACACCGTGGAGAA GACAGTGGCCCCCTACAGAATGTTCA	CAGGTGCAGCTGCAGGAGTCGGGCC CAGGACTGGTGAAGCCTTCGGAAAC CCTGTCCCTACCTGCAGTGTCTCTG GTGGCTCCATCAGCAGTAGTAGTTAC TACTGGGCTGGATCAGCAGGCCCC CAGGAAAGGGCTGGAGTGGATTGG GAGTATCTATTATAGTGGATCACCA ACTACAACCCGTCCCTCAAGAGTCG AGTACCATCTCCGTAGACACGTCCA AGAACCAAGTCTCCCTGAAGCTGAGT TCTGTGACCCGCCGACAGACAGGCCG TGTATTACTGTGCGAGGCCCAAGTAAC TACGATGCTTTTGATATCTGGGGCCA AGGACAATGGTCAACCGTCTCTCA GCCTCCACCAAGGCCCATCTGGTCTT CCCCCTGGCACCCCTCTCTCCAAGACA CCTCTGGGGCACAGCGGCCCTGGG CTGCTGGTCAAGGACTACTTCCCCCG AACCGGTGACGGTGTCTGTGGAATC AGCGCCCTGACCAGCGCGCTGCAC ACCTTCCCGGTGTCTACAGTCTCTC AGGACTCTACTCCCTCAAGAGCGGTG GTGACCGTCCCTCCAGCAGCTTGGG CAGCCAGACCTACATCTGCAACGTG AATCACAAGCCCAAGCAACACCAAGG TGGACAAGAAAGTTGAGCCCCAAATC TTGTGACAAAACCTCACACATGCCAC CGTGCCCAAGCACCTGAATCTCTGGG GGGACCGTCACTCTCTCTTCCCCC CAAAACCAAGGACACCCCTCATGAT CTCCCGGACCCCTGAGGTACATGCG TGGTGGTGGAGCTGAGCCACGAAGA CCCTGAGGTCAAGTTCAACTGGTACG TGGACGGCTGGAGGTGCATAATGC CAAGACAAAGCCGTGCGAGGAGCAG
i P S : Ig zS EF L2 L2 *G K- K:: (G H 4S 9) F ab	21 - 23 3 4 HJ: hu Ig G1 zS EF L2 L2 *G K- K:: (G H 4S 9) F u ant i- <h uC D4 0> 21- 23		

0_33 H9 V Hj: .hu lg G1 z- C H1 - E:: EP KS C + [an ti- <h u Me sot hel in> 4G 12 VL j:: hu LL C2 -E + [an	TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACCAAGGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCC TGCTGGACTCCGACGGCTCCTTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGGAGGTGCGGATCCCA GGTGCAAGTTGGTGGAGTCTGGGGGA GGCGTGGTCCAAGCCTGGGAGGTCCC TGAGACTCTCTGTGCGACCGTCTGGA TTCACCTTCAGTAGCCATGGCATGCA CTGGGTCCGCCAACCTCCAGGCAAG GGGTGGAGTGGGTGGCAGTTATCT GGTATGATGGAAGTAATGAATACTA TGGAGACTCCGTGAAGGGCCGATTTC ACCATCTCCAGAGACAATTCCAAGA ACACGCTGTATCTGCAAAATGAACAG CCTGAGAGTCGAGGACACGGCTGTG TATTACTGTACGAGAGGGGGGGGCC ACTGGAACACGAGGGCCACTACTA TGGTATGGACGTCTGGGGCCAAGGG ACCACGGTCACCGTGTCTCTCAGCAA GCACGAAAGGGCCGTCCGTATTTC
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[illegible]

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				(SEQ ID NO: 695)
				(SEQ ID NO: 694)

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	QSVLTQPPSVSGAPGQRVTISCTGSSNN IGAGYDVHWYQQVPGTAPKLLIYGN KRPSGV/PDRFSGSKGTSASLAITGLQ AEDEADYYCQSYDSSLGGWVFGGKT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNNKYAAESYLSLTP EQWKSHRYSYSCQVTHEGSTVEKTVAP TECS	QSVLTQPPSVSGSPGQSVTISCTGTSSD VGGYIFVSWYQQHPGKAPKLMYDVS KRPSGV/PDRFSGSKSVNTASLTISGLQ AEDETDYYCCSYAGNYTYVFGTGTKV TVLGGPKAAPSVTLFPPSSEELQANKA TLVCLISDFYPGAVTVAWKADSSPVK AGVETTTPSKQSNNKYAAKSYLSLTPE QWKSHRYSYSCQVTHEGSTVEKTVAPT ECS	QVQLQESGPGLVKPSETLSLTCTVSGG SISSSSYYWGWIRPPGKGLEWIGSIYY SGITNYNPSLKSRSVTISVDTSKNQFSLK LSSVTAADTAVYCARPSNYDAFDIW QGQTMVTVSSASTKGPSPFLAPSSKS TSGGTAALGCLVKDYFPEPVTVSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHNKPSNTKVDKK VEPKSCDKTHTCTPCPAPELLGGPSVF LFPPKPKDTLMISRTPETCVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTPPVLD SDGSFFLYSKLTVDKSRWQQGNVFC SVMHEALHNYHTQKSLSLSPGGGGSG GGGSQVQLVESGGGVVQPGRSLRLSC AASGFTLSSYGMHWVRQAPGKGLEW VAWIYDGSNKYYADSVKGRVTTISRD NSKNTLYLQMNLSRAEDTAVYYCTRD GRNYVYFDNWGGQGLTVTVSSASTKGP SVFPLAPSSKSTGGYTAALGCLVKDYF PEPVTVSWNSGALTSGVHTFPAVLQSS GLYSLESVVTVPPSSSLGTQTYICNVNH KPSNTKVDKKVEPKSC	(SEQ ID NO: 700)	(SEQ ID NO: 701)	(SEQ ID NO: 702)
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				TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACCAAGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGGAAGGTGGCGGATCCCA GGTACAGCTGCAACAGTCAGGTCCA GGACTGGTGAAGCCCTCGCAGACCC TCTCACTACCTGTGCCATCTCCGGG GACAGTGTCTTAGCAGCCGTACTGC TTGGAACTGGATCAGGCAGTCCCCAT CGAGAGGCCCTTGAGTGGCTGGGAAG GACATACTACAGGTCCAAGTGGTAT CATGATTATTCAGTATCTGTGAAAAG TCGAATCACCATCGACCCAGACACA TCCAAGAACCAGTTCTCCCTGCAGCT GAACCTGTGTACTCCCGAGGACACG GCTGTTTATTATTGTGCAAGAGGGGC TGCTCCCTTTGACTACTGGGGCCAGG GAACCCCTGGTACCGTGTCTCTCAGCA AGCACGAAGGGGCCGTCCTGATTTTC CGCTTGCGCCCTCGTCTGGAAGTCAACT
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				TCGGGAGGACCGCGGCACTTGGCT GTCTTGTCAAAGATTACTTCCCTGAG CCAGTGACAGTCAGCTGGAATTCCG GTGCCCTCACGTCAGGAGTACATAC ATTCCCTGCGGTATTGCAGTCCCTCCG GACTCTACTCCCTGGAGTCGGTGGTA ACGGTGCCCAAGCTCCAGCTTGGGGA CCCAGACGTACATTGTAACTGAAT CACAAACCAAGCAATACTAAGGTAG ATAAGAAAGTAGAACCCGAAGAGCTG C
				(SEQ ID NO: 705)
				(SEQ ID NO: 704)
				(SEQ ID NO: 703)
ti- <h uC D4 0> 21- 23 0_ 36 F3 VL J:: hu KL C- SI 76 K (Ig G- Fa b); L M R ID: SS - 30 82 5				

			CAGTCAGTGTTGACGAGCCGCCCTC AGTGTCTGGGCCCCAGGCAGAGG GTCACCATCTCTGCACTGGAGCAG CTCCAACATCGGGCAGGTTATGAT GTTCACTGGTACCAGAGGTTCCAGG AACAGCCCCAAACTCCTCATCTATG GTAACAGCAAGCGGCCCTCAGGGGT CCCTGACCGATTCTCTGGCTCCAAGT CTGGCACCTCAGCCTCCCTGGCCATC ACTGGGCTCCAGGCTGAGGATGAGG CTGATTATTACTGCCAGTCTCTATGAC AGCAGCCTGGGTGGTGGTGTTCG GCGGAGGGACCAAGCTGACCCGTCT ACAGCCCCAAGGCTGCACCCCTGGTC ACTCTGTTCCCGCCCTCTCTGAGGA GCTTCAAGCCAAACAAGGCCACACTG GTGTGTTCTATCAGTGACTTCTACCC GGGAGCCGTGACAGTGGCCTGGAAG GCAGATAGCAGCCCCGTCAAAGCGG GAGTGGAAACCAACCACACCTCCAA ACAAAGCAACAACAAGTACGCGGCC GAAAGCTATCTGAGCCTGACGCCGTG AGCAGTGGAAATGCCACAGAAAGCTA CAGCTGCCAGGTACGCATGAAGGG AGCACCGTGGAGAGACAGTGGCCC CTACAGAATGTTCA		TCCTATGAGCTGACTCAGCCACCCCTC AGTGTCCGTGTCCCAAGGACAGACA GCCAGCATCACTGCTCTGGAGAAA GGTTGGAAATAAATAATATTGCTGG TATCAGCAGAAGCCAGGCCAGTCCC CTGTTCTGGTCACTATCAAGATTTC AAGCGCCCTCAGGGATCCCTGAGC GATTCTCTGGCTCCAACCTCTGGGATC ACAGCCACTCTGACCATCAGCGGGA CCCAGGCTATGGATGAGGCTGACTA TTACTGTACGGCGTGGGACAGCAGA ACTGTGTTATTCGGCGGAGGGACCA AGCTGACCGTCTTAGTACGCCCAA GGCTGCACCCCTGGTCACTCTGTTC CGCCCTCTCTGAGGAGCTTCAAGCC AACAAAGGCCACACTGGTGTCTCAT CAGTGACTTCTACCCGGGAGCCGTG ACAGTGGCCTGGAAGGCAGATAGCA GCCCGTCAAAGCGGGAGTGGAAAC CACCACACCTCCAAACAAGCAAC AACAAAGTACGCGGCCAAGAGCTATC TGAGCCTGACGCTGAGCAGTGGAA GTCCCACAGAAAGCTACAGCTGCCAG GTCACGCATGAAGGAGCACCGTGG AGAAAGACAGTGGCCCCCTACAGAATG TTCA		CAGGTGCAGCTGCAGGAGTCGGGCC CAGGACTGGTGAAGCCTTCGGAAAC CCTGTCCCTCACTGCACCTGTCTCTG GTGGCTCCATCAGCAGTAGTAGTTAC TACTGGGGCTGGATCAGGACGCCCC CAGGAAAGGGCTGGAGTGGATTGG GAGTATCTATTATAGTGGATCACCA ACTACAACCCGTCCCTCAAGAGTCG AGTACCATCTCCGTAGACACGTCCA AGAACCAATCTCCCTGAAGCTGAGT TCTGTACCGCCGACAGACAGCGCCG TGTATTACTGTGCGAGGGCCAGTAAC TACGATGCTTTTGATATCTGGGGCCA AGGACAATGGTCAACCTGTCTCA GCCTCCACCAAGGGCCATCGGTCTT CCCCCTGGCACCTCTCTCCAAGAGCA CCTCTGGGGCACAGCGGCCCTGGG CTGCTGGTCAAGGACTACTTCCCCG AACCGGTGACGGTGTCTGTGAACTC AGCGCCCTGACCAGCGCGTGCAC ACCTTCCCGGTGTCTACAGTCTC AGGACTCTACTCCCTCAAGAGCGTG GTGACCGTGCCCTCCAGCAGCTTGGG CACCAGACCTACATCTGCAACGTG AATCACAAGCCCAGCAACACCAAGG TGGACAAGAAAGTTGAGCCCCAAATC TTGTGACAAAACCTCACACATGCCAC CGTGCCCAAGCACCTGAACCTCTGGG GGGACCGTCACTCTCTCTTCCCCC CAAAACCCAAAGGACACCCCTCATGAT CTCCCGGACCCCTGAGGTCACATGCG TGGTGGTGGACGTGAGCCACGAAGA CCCTGAGGTCAAGTTCAACTGTGACG TGGACGGCGTGGAGGTGCATAATGC CAAGACAAAGCCGTGCGGAGGAGCAG
i P S : 5 7 6 1 23 5 5	21 - 23	21 23 3 4 G 12 12 V HJ: hu Ig 12 G 2 1- 23 0 37 A 4S 6 F ab	21 23 3 4 G 12 12 V HJ: hu Ig 12 G 2 1- 23 0 37 A 4S 6 F ab	21 23 3 4 G 12 12 V HJ: hu Ig 12 G 2 1- 23 0 37 A 4S 6 F ab	21 23 3 4 G 12 12 V HJ: hu Ig 12 G 2 1- 23 0 37 A 4S 6 F ab		

				TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACCAAGGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGAGGTGGCGGATCCCA GGTGCAAGTGGTGGAGTCTGGGGGA GGCTTAGTCAAGCCTGGAGGGTCCCT GAGACTCTCTGTGCAGCCTCTGAAT TCACCTTCAGTGACTACTACATGAGC TGGATCCGCCAGGCTCCAGGGAAGG GGCTGGAGTGGGTTTCATATATAGT CGAAGTGGTGATACCATCTACTACGC AGACTCTGTGAAGGGCCGATTACCC ATCTCCAGGGACAACGCCAAGAACT CACTGTATCTGCAATGAATGGCCTG CGAGCCGAAGACACGGCCGTGTATT ACTGTGCGAGAGACTTAGCAGCAGG TGCTACAGGGGGCTTGACTGCTGG GGCCAGGGAAACCCCTGGTCACCGTGT CCTCAGCAAGCACGAAGGGGCCGTC CGTATTCCGCTTGCGCCCTCGTCGA
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				AGTCAACTTCGGGAGGGACCGCGGC ACTTGGCTGTCTTGTCAAAGATTACT TCCCTGAGCCAGTGACAGTCAGCTG GAATTCGGGTGCCCTCACGTCAGGA GTACATACATTCCCTGCGGTATTGCA GTCTCCGGACTCTACTCCCTGGAGT CGGTGGTAACGGTGCCAGCTCCAG CTTGGGGACCCAGACGTACATTGTGA ACGTGAATCACAAACCAGCAATAC TAAGGTAGATAAGAAAGTAGAACCG AAGAGCTGC
				(SEQ ID NO: 711)
				(SEQ ID NO: 710)
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ti- <h uC D4 0> 21- 23 0_ 37 A6 VL J:: hu LL C2 -K (lg G- Fa b); L M R ID: SS - 30 82 6				

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(SEQ ID NO: 712)	(SEQ ID NO: 713)	(SEQ ID NO: 714)

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			QSVLTQPPSVSGAPQQRVTITISCTGSSSN IGAGYDVHWYQQVPGTAPKLLIYGNS KRPSGVPDFRFGSKSGTASLAITGLQ AEDEADYYCQSYDSSLGGWVFGGTT KLTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNINKYAAESYLSLTP EQWKSHRSYSCQVTHEGSTVEKTVAP TECS	QSALETQASVSGSPGQITISCTGTSSD VGNYNLVSWYQQHPGKAPKLMIEV NRRPSGVSNRFGSKSGNTASLTISGLQ AEDEAEYYCCSYAGRDTFVVFGGGTK LTVLQPKAAPSVTLFPPSSEELQANK ATLVCLISDFYPGAVTVAWKADSSPV KAGVETTTPSKQSNINKYAAESYLSLTP EQWKSHRSYSCQVTHEGSTVEKTVAP TECS	QVQLQESGPGLVKPSSETLSLTCTVSGG SISSSYWGWIRQPPGKGLEWIGSIYY SGITNYPNPSLKSRTISVDTSKNQFSLK LSSVTAADTAVYYCARPSNYDAFDIW GQGTMTVTSSASTKGPSVFPLAPSSKS TSGGTAALGCLVKDYFPEPTVTSWNS GALTSGVHTFPAVLQSSGLYSLKSVVT VPSSSLGTQTYICNVNHPKPSNTKVDKK VEPKSCDKTHTCPPCPAPELLGGPSVF LFPPKPKDTLMISRTPEVTCVVVDVSH EDPEVKFNWYVDGVEVHNAKTKPCE EQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVQSGTEVKKPGASVKVSC KASGYTPGYYMHWVRQAPGQGGLW MGWINPDSGGTKYTKFQGRVTLTRD ASVSTAYIDLNRLSDDTAVYYCARE RCRTTNCLDYWGQGSLLTVSSASTK GPSVFPLAPSSKSTSGGTAALGCLVKD YFPEPVTVSWNSGALTSGVHTFPAVLQ SSGLYSLESVTVTPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSC
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AGCACGTACCGTTGCGTCAGCGTCCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGGAGTACAAGTGCAAGG TGTCACAACAAGCCCTCCAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACACACAGGT GTACACCTGCCCCCATCCCCGGGAG GAGATGACCAAGAACACAGGTCAGCC TGACCTGCCTGGTCAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGGCAGCCGGAGAACAA CTACAAGACCACGCTCCCGTGCTGG ACTCCGACGGCTCCTTCTTCTCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGTGGTGGCGGAT CGGAGGTGGCGGATCCCAGGTGCA ACTGGTGCAGTCTGGGCTGAGGTG ACGAAGCCTGGGGCTCAGTGAAGG TGTCTGCAAGGCTTCTGGATACACC TTCGCCGGCTACTATATGCACTGGGT GCGACAGGCCCTGGACAAGGGCTT GAGTGGATGGGATGGATCAACCCCTC ACAGTGGTGGCACAAACTATGCACA GAAGTTTCAGGACAGGGTCACCATG ACAGGGACACGTCCATCAACACAG CCTACATGGAAGTGAAGAGGCTGAG ATCTGACGACACGGCCGTGTATTACT GTGCGAGAGAACGTATTTCTATGGTT CGGGAGTCGGGCACAACTGGTTCTG CCCCCTGGGGCCAGGGAACCCCTGGT CACCGTGTCTCAGCAAGCACGAAG GGGCCGTCCGTATTTCCGCTTGCGCC																																		

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82	DIQMTQSPSSVSASVGDRVTITCRASQ GITRWLAWYQQKPGKAPKLLIYAASV LQSGVPSRFSGSGTDFTLTISSLQPE DFATYYCQQSNFPRTFGGQGTKVEIKR TVAAPSVFIFFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSITYSLESTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSLSASVGDRVTITCRASQ DISNNLAWFQQKPGKPKSLMYAASS LHSGVPSFSGSGGTDFTFITISLQPE DFATYYCQQNSYPLTFGGGKVEIRR TVAAPSVFIFFPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSITYSLKSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMTWIRQAPGKGLEWISYISS SGSTIYYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDRNSHFD YWGQGLVTVSSASTKGPSVFPLAPSS KSTGGTAALGCLVKDYFPEPVTVSW NSGALTSGVHTFPAVLQSSGLYSLKSV VTVPSSSLGTQTYICNVNHKPSNTKVD KKVEPKSCDKTHCTCPCPAPPELLGGPS VFLFPPKPKDTLMISRTPEVTCVVVDV SHEDPEVKFNWYVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGK EYKCKVSNKALPAPIEKTISKAKGQPR EPQVYITLPPSREEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPPVL DSDGSFFLYSKLTVDKSRWQQGNVFS CSVMHEALHNHYTQKSLSLSPGSGGS GGGGSQVQLVQSGAEVTKPGASVKVS CKASGYTFAGYIMHWVRQAPGQGLE WMGWINPHSGGTNYAQKFQDRVTMT RDTISNTAYMELSLRSDDTAVYYCA RERISMVRGVGHNWFAPWGQGTLVLT VSSASTKGPSVFPLAPSSKSTSGGTAAL GCLVKDYFPEPVTVSWNSGALTSGVH TFPAVLQSSGLYSLESVVTVPSSSLGTQ TYICNVNHKPSNTKVDKKVEPKSC (SEQ ID NO: 730)
9			(SEQ ID NO: 731)
			(SEQ ID NO: 732)

457

				AGCACGTACCGTTGCGTCAGCGTCCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGAGTACAAAGTCAAGG TGTCCAACAAGCCCTCCAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACCACAGGT GTACACCTGCCCCCATCCCCGGAG GAGATGACCAAGAACCAGGTCAGCC TGACCTGCCTGGTCAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGGCAGCCGGAGAAACA CTACAAGACCACGCTCCCGTGTGG ACTCCGACGGCTCCTCTTCCCTCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGGTGGTGGCGGAT CGGAGGTGGGGATCCGAGGTGCA GCTGCTGGAGTCTGGGGAGGCTTG GTACAGCCTGGGGGTCCCTGAGAC TCTCCTGTGCAGCCTCTGGATTACCC TTTAGTAGAAATGCCATGAGTTGGGT CCGCCAGGCTCCAGGGAAGGGGCTG GAGTGGGTGTACGCTACTGGTGGTA GTGGTATTAGCACATACTACGCAGA CTCCGTGAAGGGCCGGTTCAACATCT CCAGAGACAATTCCAAGAACACGCT GTATCTGCAATGAACAGTCTGAGA GCCGAGGACACGGCCGTATATTACT GTGCGAGAGGTTATAGCAACAGCTG GTGGTACTTTGACTACTGGGGCCAGG GAACCCCTGGTCAACCGTCTCCTCAGCA AGCACGAAGGGGCCGTCCGTATTTC CGCTTGCGCCCTCGTCGAAGTCAACT
0_	30	AI	2	V
Hj:	:hu	lg	G1	z-
C	HI	-	E::	EP
KS	C	+	[an	ti-
<h	u	Me	sot	hel
in>	4H	6	VL	l::
hu	KL	C-	SI	76

459

			DIQMTQSPSSVSA SVGDRVTITCRASQ GITRWLA WYQQKPGKAPKLLIYAASV LQSGVPSRFGSGGTDFTLTISSLQPE DFATYYCQQSNFPRTFGQGTKVEIKR TVAAPSVFIFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQJE SVTEQDSKDSYSLLESTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QSA LTQPA SVSGSPGQSITISCTGTSSD VGNYNL VSWYQQHPGKAPKLMIFEV NQRPSGVSNRFSGSKSGTTASLTISGLQ AADEADYFCCSYTTTSSTYVIFGGGTKL TVLGQPKAAPSVTLFPPSSEELQANKA TLVCLISDFYPGA VTVAWKADSSPVK AGVETTTPSKQSNNKYAAKSYLSLTPE QWKSHRSYSCQVTHEGSTVEKTVAPT ECS	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMTWIRQAPGKGLEWISYISS SGSTIYYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDRNSHFD YWGGQGLVTVSSASTKGPSVFPLAPSS KSTGGTAALGCLVKDYFPEPVTVSW NSGALTSGVHTFPAVLQSSGLYSLKSV VTVPSSSLGTQTYICNVNHKPSNTKVD KKVEPKSCDKTHTCPPCPAPPELLGGPS VFLFPPKPKDITLMISRTPEVTCVVVDV SHEDPEVKFNWYVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGK EYKCKVSNKALPAPIEKTISKAKGQPR EPQVYITLPSREEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPPVL DSDGSFFLYSKLTVDKSRWQQGNVFS CSVMHEALHNHYTQKSLSLSPGSGGS GGGSEVQLLESGGGLVQPGGSLRLS CAASGFTFSRNAMSWVRQAPGKGLE WV'SATGGSGISTYYADSVKGRFTISR NSKNTLYLQMNSLRAEDTAVYYCAR GYSNSWVYFDYWGGQGLVTVSSAST KGPSVFPLAPSSKSTSGGTAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPAVL QSSGLYSLESVTVTPSSSLGTQTYICNV NHKPSNTKVDKKVEPKSC
					(SEQ ID NO: 736)
					(SEQ ID NO: 737)
					(SEQ ID NO: 738)

[illegible]

0_	33	H6	V	Hj:	:hu	Ig	G1	z-	C	H1	-	E::	EP	KS	C	+	[an	ti-	<h	u	Me	sot	hel	in>	4H	6	VL	l::	hu	KL	C-	SI	76	E
AGCACGTACCGTTGCGTCAGCGTCCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGGAGTACAAGTGCAAGG TGTCCAACAAAGCCCTCCCAAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACACACAGGT GTACACCCTGCCCCATCCCCGGGAG GAGATGACCAAGAACACAGGTCAGCC TGACCTGCCTGGTCAAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGGCAGCCGGAGAACA CTACAAGACCACGCTCCCCGTGCTGG ACTCCGACGGCTCCTTCTTCTCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGGTGGTGGCGGAT CGGAGGTGGCGGATCCCAGGTGCA ACTGGTGCAGTCTGGGGCTGAAGTG AAGAAAGCTGGGGCTCAGTGAAGG TGTCTGCAAGGCTTCTGGATACACC TTCCCCGGCTACTATATGTACTGGTT GCGACAGGCCCCCTGGACAAGGACTT GAGTGGATGGGATGGATCAACCCCTG ACAGTGGTGACACAAACTATGCACA GAAGTTTCAGGGCAGGGTCACCATG ACCAGGGACACGTCCATCAGCACAG CCTTATGGAGCTGAGCAGGCTGAG ATCAGACGACACGGCCGTGTATTACT GTGCGAGAGAGAAAGCCAGATATTT TGACTCCTTCTACTACTACCTTATGG ACGTCTGGGGCCAAAGGGACCAACGGT CACCGTGTCTCAGCAAGCACGAAG GGGCCGTCCGTATTTCCGCTTGCGCC																																		

463

			DIQMTQSPSSVSASVGDRVTITCRASQ GITRWLAWYQQKPGKAPKLLIYAASV LQSGVPSRFGSGGTDFTLTISSLQPE DFATYYCQQSNFPRTFGGQGTKVEIKR TVAAPSVFIFFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQIE SVTEQDSKDSITYSLESTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSLSASVGDRVTITCRAGQ NISRHLNHWYQQNPCKAPKVLJHPASSL PSGVPSRFGSGGTDFTSLTISSLQPEDF GTFFCQQSYSTPPTFGGGTKVELKRTV AAPSVFIFFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQIESVT EQDSKDSITYSLKSTLTLSKADYEKHK VYACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMTWIRQAPGKGLEWISYISS SGSTIYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDNRSHFD YWGQGTLVTVSSASTKGPSVFLAPSS KSTSGTAALGCLVKDYFPEPTVSW NSGALTSGVHTFPAVLQSSGLYSLKSV VTVPSSSLGTQTYICNVNHKPSNTKVD KKVEPKSCDKTHTCPPCPAPPELLGGPS VFLFPPKPKDITLMISRTPEVTCVVVDV SHEDPEVKFNWYVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGK EYKCKVSNKALPAPIEKTISKAKGQPR EPQVYITLPPSREEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPPVL DSDGSFFLYSKLTVDKSRWQQGNVFS CSVMHEALHNHYTQKSLSLSPGSGGS GGGGSQVQLVQSGAEVKKPGASVKVS CKASGYTFPGYMYWLRQAPGQGLE WMGWINPDSGDTNYAQKFGQGRVTMT RDTISIAFMELSLRLSDDTAVYYCAR EKPRYFDSFYIYLMMDVWGQGTTVTS SASTKGPSVFPEPLAPSSKSTSGGTAALG CLVKDYFPEPTVTVSWNSGALTSGVHT FPAVLQSSGLYSLESVVTVPSSSLGTQT YICNVNHKPSNTKVDKKVEPKSC
			(SEQ ID NO: 742)	(SEQ ID NO: 743)	(SEQ ID NO: 744)

465

0_33	H9	V	HJ:	:hu	Ig	G1	z-	C	H1	-	E::	EP	KS	C	+	[an	ti-	<h	u	Me	sot	hel	in>	4H	6	VL	l::	hu	KL	C-	S1	76	E
AGCACGTACCGTTGCGTCAGCGTCTCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGGAGTACAAGTGCAAGG TGTCACAACAAGCCCTCCAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACCACAGGT GTACACCTGCCCCATCCCCGGGAG GAGATGACCAAGAACAGGTCAGCC TGACCTGCCTGGTCAAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGCAGCCGGAGAACAA CTACAAGACCACGCTCCCGTGTGG ACTCCGACGGCTCCTTCTTCTCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGGTGGTGGCGGAT CGGAGGTGGCGGATCCCAGGTGCA GTTGGTGGAGTCTGGGGAGGCGTG GTCCAGCCTGGGAGGTCCCTGAGAC TCTCCTGTGCAGCGTCTGGATTCCACC TTCAGTAGCCATGGCATGCACTGGGT CCGCCAACCTCCAGGCAAGGGGCTG GAGTGGGTGGCAGTTATCTGGTATG ATGGAAGTAATGAATACTATGGAGA CTCCGTGAAGGGCCGATTCACCATCT CCAGAGACAATTCCAAGAACACGCT GTATCTGCAAAATGAACAGCCTGAGA GTCGAGGACACGGCTGTGTATTACTG TACGAGAGGGGGGGCCACTGGAAC TACGAGGGCCACTACTATGGTATGG ACGCTGGGGCCAAAGGACCACGGT CACCGTGTCTCAGCAAGCACGAAG GGGCCGTCCGTATTTCCGCTTGCGCC																																	

467

		DIQMTQSPSSVSASVGDRVTITCRASQ GITRWLAWYQQKPGKAPKLLIYAASV LQSGVPSRFGSGGTDFTLTISSLQPE DFATYYCQQSNISFPRTFGQGTKVEIKR TVAAPSVFIFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQIE SVTEQDSKDSITYSLESTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QAVPTQPSSLSASPGASASLTCTLRSGI NVGSSRIYWYQQKPGSPQFLLRYTSD SDKLQSGVPSRFGSGKDSANAGLLL ISGLQSEDEADYYCMIWHSSAVVFGG GTKLTLVLGQPKAAPSVTLFPPSSEELQ ANKATLVCLISDFYPGAVTVAWKADS SPVKAQVETTTPSKQSNNKYAAKSYL SLTPEQWKSHRSYSCQVTHEGSTVEKT VAPTECS	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMTWIRQAPGKGLEWISYISS SGSTIYYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDRNSHFD YWGGQGLVTVSSASTKGPSVFLAPSS KSTSGGTAAALGCLVKDYFPEPTVSW NSGALTSGVHTFPAVLQSSGLYSLKSV VTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHTCPPCPAPPELLGGPS VFLFPPKPKDITLMISRTPEVTCVVVDV SHEDPEVKFNWYVDGVEVHNAKTKP CEEQYGSTYRCVSVLTVLHQDWLNGK EYKCKVSNKALPAPIEKTISKAKGQPR EPQVYITLPPSREEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPPVL DSDGSFFLYSKLTVDKSRWQQGNVFS CSVMHEALHNHYTQKSLSLSPGSGGS GGGGSQVQLVESGGGVVQPGRSLRLS CAASGFTFSHGMHWVRQPPGKGLE WVAWIWYDGSNEYYGDSVKGRFTISR DNSKNTLYLQMNSLRVEDTAVYYCTR GGGHWNYEGHYYGMDVWGQGTITV VSSASTKGPSVFLAPSSKSTSGGTAAAL GCLVKDYFPEPTVSWNSGALTSGVH TFPAVLQSSGLYSLESVVTVPSSSLGTQ TYICNVNHNKPSNTKVDKKEPKSC (SEQ ID NO: 748)
			(SEQ ID NO: 749)	(SEQ ID NO: 750)

[illegible]

0_35	F1	I	V	Hj:	:hu	Ig	G1	z-	C	H1	-	E::	EP	KS	C	+	[an	ti-	<h	u	Me	sot	hel	in>	4H	6	VL	::	hu	KL	C-	SI	76
AGCACGTACCGTTGCGTCAGCGTCCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGGAGTACAAGTGCAAGG TGTCCAACAAAGCCCTCCCAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACACACAGGT GTACACCTGCCCTCCCTATCCCGGGAG GAGATGACCAAGAACACAGGTCAGCC TGACCTGCCTGGTCAAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGGCAGCCGGAGAACAA CTACAAGACCACGCTCCCGTGCTGG ACTCCGACGGCTCCTTCTTCTCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGTGGTGGCGGAT CGGAGGTGGCGGATCCCAGTGCA GCTGTGGAGTCTGGGGAGGCGTG GTCCAGCCTGGGAGGTCCCTGAGAC TCTCCTGTGCAGCGTCTGGATTCAAC CTCAGTAGCTATGGCATGCCTGGGT CCGCCAGGCTCCAGGCAAGGGGCTG GAGTGGTGGCAGTTATCTGGTATG ATGGAAGTAATAAATACTATGCAGA CTCCGTGAAGGGCCGAGTCACCATCT CCAGAGACAATTCCAAGAACACGCT GTATCTGCAATGAATAGCCTGAGA GCCGAGGACACGGCTGTGTATTACT GTACGAGAGATGGCCGGAACACTACGT CTACTTTGACAACTGGGGCCAGGGA ACCCTGGTCACCGTGTCTCTCAGCAAG CAGGAAGGGGCCGTCCTGATTTCCGC TTGGCCCTCGTCGAAGTCAACTTCG																																	

E + [an ti-<h uC D4 0> 21- 23 0 35 F1 1 VL]:: hu LL C2 -K (lg G- Fa b); L M R ID: SS - 30 83 3				GGAGGACCGCGCACTTGGCTGTC TTGTCAAAGATTACTTCCCTGAGCCA GTGACAGTCAGCTGGAATTCCGGTG CCCTCACGTCAGGAGTACATACATTC CCTGCGGTATTGCAGTCTCCCGGACT CTACTCCCTGGAGTCGGTGGTAACGG TGCCCAGCTCCAGCTTGGGGACCCA GACGTACATTGTAACTGAATCACA AACCAAGCAATACTAAGGTAGATAA GAAAGTAGAACCCGAAGAGCTGC	(SEQ ID NO: 753)
					(SEQ ID NO: 752)
					(SEQ ID NO: 751)

473

474

[illegible]

		DIQMTQSPSSVSASVGDRVTITCRASQ GITRWLAWYQQKPGKAPKLLIYAASV LQSGVPSRFGSGGTDFTLTISLSQPE DFATYYCQQSNFPRTFGGQGTKVEIKR TVAAPSVFIFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSYSLSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	EIVLTQSPGTLSSLSPGERATLSCRASQ VSSNYLAWYQQKPGQAPRALIYAASN RAAGISDRFSGSGGTDFTLTISRLEPE DFAVYFCQQYGSSPLTFGGGTKVEIKR TVAAPSVFIFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSYSLKSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMTWIRQAPGKGLEWISYISS SGSTIYADSVKGRFTISRDNAKNSLY LQMNSLRAEDTAVYYCARDRNSHFD YWGGQGLVTVSSASTKGPSVFPLAPSS KSTSGGTAAAGCLVKDYFPEPVTVSW NSGALTSGVHTFPAVLQSSGLYSLKSV VTVPSSSLGTQTYICNVNHNKPSNTKVD KKVEPKSCDKTHTCPPCPAPPELLGGPS VFLFPPKPKDITLMISRTPEVTCVVVDV SHEDPEVKFNWYVDGVEVHNATKTP CEEQYGSTYRCVSVLTVLHQDWLNGK EYKCKVSNKALPAPIEKTISKAKGQPR EPQVYITLPPSREEMTKNQVSLTCLVKG FYPSDIAVEWESNGQPENNYKTTPPVL DSDGSFFLYSKLTVDKSRWQQGNVFS CSVMHEALHNHYTQKSLSLSPGSGGS GGGSQVQLQSGGPGGLVKPSQTLST CAISGDSVSSRTAWNWRQSPSRGLE WLGRTYYRSKWHYHDYSVVKSRTID PDTSKNQFSLQLNSVTPEDTAVYYCAR GAAPFDYWGGQGLTVTVSSASTKGPSV FPLAPSSKSTSGGTAALGCLVKDYFPE PVTVSWNSGALTSGVHTFPAVLQSSGL YSLESVTVTPSSSLGTQTYICNVNHNK SNTKVDKKVEPKSC
		(SEQ ID NO: 760)	(SEQ ID NO: 761)	(SEQ ID NO: 762)

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481

482

[illegible]

[illegible]

486

487

		DIQMTQSPSSVSASVGDRTITCRASQ DISRWLAWYQQKPGKAPKLLISAAAR LQSGVPSRFGSGGTDFTLTISSLQPE DFAIYCCQAKSFPRTFGGQGTKVEIKR TVAAPSVFIFFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQIE SVTEQDSKDSITYLESSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSLSASVGDILTITCRASQNI TTYLNWYQQKPGKAPNLLISAAARLRS GVPSRFGSGGTDFTLTISSLQPVDF TFYCQQFTTTPWTFGGQGTKVEIKRTVA APSVFIFFPPSDEQLKSGTASVVCLLNMF YPREAKVQWKVDNALQSGNSQIESVTE QDSKDSITYSLKSTLTLSKADYEKHKV YACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMSWIRQAPGKGLEWISYISS SESIYYVDSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDVGSHEFDY WGQGLTVTVSSASTKGPSVFPLAPSSK STGGTAALGCLVKDYFPEPTVSWN SGALTSGVHTFPAVLQSSGLYSLKSVV TVPSSSLGTQTYICNVNHKPSNTKVDK KVEPKSCDKTHTCPCPAPELLGGPSV FLPPKPKDITLMISRTPETVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFS SVMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVQSGAEVKKPGASVKVSC KASGYTFAGYIMHWVRQAPGQGLE WMGWINPDSGGTNFAQQFQGRVTMT RDTISITAYMEVSRRLSDDTAVFYCAR EKITMTGIYFDYWGQGLTVTVSSASTK GPSVFPLAPSSKSTSGGTAAALGCLVKD YFPEPTVTVSWNSGALTSGVHTFPAVLQ SSGLYSLESVTVTPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSC
		(SEQ ID NO: 778)	(SEQ ID NO: 779)	(SEQ ID NO: 780)

[illegible]

490

491

83	DIQMTQSPSSVSASVGDRVTITCRASQ DISRWLAWYQQKPKAPKLLISAASR LQSGVPSRFSGSGTDFTLTISSLQPE DFAIYYCQQAQKSFPRTFGGQGTKVEIKR TVAAPSVFIFFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSYLESSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSLSASVGDRVTITCRASQ DISNNLAWFQQKPKPKSLMYAASS LHSGVPSFTSGSGGTDFTFITISLQPE DFATYYCQQNSYPLTFGGGTKVEIRR TVAAPSVFIFFPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSYSLKSTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMSWIRQAPGKGLEWISYISS SESIYYVDSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDVGSHEFDY WGQGLTVTVSSASTKGPSVFPLAPSSK WGSGTAAALGCLVKDYFPEPVTVSWN SGALTSGVHTFPAVLQSSGLYSLKSVV TVPSSSLGTQTYICNVNHKPSNTKVDK KVEPKSCDKTHTCPCPAPELLGGPSV FLPPKPKDITLMISRTPETVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTISKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGQVQLVQSGAEVTKPGASVKVSC KASGYTFAGYIMHWVRQAPGQGLE WMGWINPHSGGTNYAQKFQDRVTMT RDTISNTAYMELSLRSDDTAVYYCA RERISMVRGVGHNWFAPWGQGTLVLT VSSASTKGPSVFPLAPSSKSTSGGTAAAL GCLVKDYFPEPVTVSWNSGALTSGVH TFPAVLQSSGLYSLESVVTVPSSSLGTQ TYICNVNHKPSNTKVDKKEPKSC (SEQ ID NO: 784)
8		(SEQ ID NO: 785)	(SEQ ID NO: 786)

[illegible]

				AGCACGTACCGTTGCGTCAGCGTCCT CACCGTCTGCACCAAGACTGGCTG AATGGCAAGAGTACAAAGTCAAGG TGTCACAACAAGCCCTCCAGCCCC ATCGAGAAAACCATCTCCAAAGCCA AAGGGCAGCCCCGAGAACCACAGGT GTACACCTGCCCCCATCCCCGGAG GAGATGACCAAGAACCAGGTCAGCC TGACCTGCCTGGTCAAGGCTTCTAT CCCAGCGACATCGCCGTGGAGTGGG AGAGCAATGGGCAGCCGGAGAAACA CTACAAGACCACGCTCCCGTGTGG ACTCCGACGGCTCCTTCTTCCCTAT AGCAAGCTCACCGTGGACAAGAGCA GGTGGCAGCAGGGGAACGTCTTCTC ATGCTCCGTGATGCATGAGGCTCTGC ACAACCACTACACGAGAAGAGCCT CTCCCTGTCTCCGGTGGTGGCGGAT CGGAGGTGGCGGATCCGAGGTGCA GCTGCTGGAGTCTGGGGAGGCTTG GTACAGCCTGGGGGTCCCTGAGAC TCTCCTGTGCAGCCTCTGGATTACCC TTTAGTAGAAATGCCATGAGTTGGGT CCGCCAGGCTCCAGGGAAGGGGCTG GAGTGGGTGTCAGCTACTGGTGGTA GTGGTATTAGCACATACTACGCAGA CTCCGTGAAGGGCCGGTTCAACATCT CCAGAGACAATTCCAAGAACACGCT GTATCTGCAATGAACAGTCTGAGA GCCGAGGACACGGCCGTATATTACT GTGCGAGAGGTTATAGCAACAGCTG GTGGTACTTTGACTACTGGGGCCAGG GAACCCCTGGTCAACCGTCTCTCAGCA AGCACGAAGGGGCCGTCCGTATTTC CGCTTGCGCCCTCGTCGAAGTCAACT
0_	30	AI	2	V
Hj:	:hu	lg	G1	z-
C	HI	-	E::	EP
KS	C	+	[an	ti-
<h	u	Me	sot	hel
in>	6F	4	VL	l::
hu	KL	C-	SI	76

[illegible]

[illegible]

0_	33	H6	V	HJ:	:hu	Ig	G1	z-	C	H1	-	E::	EP	KS	C	+	[an	ti-	<h	u	Me	sot	hel	in>	6F	4	VL	l::	hu	KL	C-	S1	76	E
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		DIQMTQSPSSVSASVGDRVTITCRASQ DISRWLAWYQQKPGKAPKLLISAAAR LQSGVPSRFGSGGTDFLTITSLQPE DFAIYYCQAKSFPRTFGGQGTKVEIKR TVAAPSVFIKPPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQAE SVTEQDSKDSYSLSESTLTLSKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	DIQMTQSPSSLSASVGDRVTITCRAGQ NISRHLNWIYQQNPGKAPKVLIIHPASSL PSGVPSRFGSGGTDFSLTISLQPEDF GTIFYCQQSYSTPTTFGGGTKVELKRTV AAPSVFIKPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVT EQDSKDSYSLKSTLTLSKADYEKHK VYACEVTHQGLSSPVTKSFNRGEC	QVQLVESGGGLVKPGGSLRLSCAASG FTFSDYYMSWIRQAPGKGLEWISYISS SESIYYVDSVKGRFTISRDNAKNSLYL QMNSLRAEDTAVYYCARDVGSHEFDY WGQGLTVTVSSASTKGPSVFPLAPSSK WGSGTAAAGCLVKDYFPEPTVSWN SGALTSGVHTFPAVLQSSGLYSLKSVV TVPSSSLGTQTYICNVNHKPSNTKVDK KVEPKSCDKTHTCPPCPAPELGGPSV FLPPPKDKDTLMISRTPETVTCVVVDVS HEDPEVKFNWYVDGVEVHNAKTKPC EEQYGSTYRCVSVLTVLHQDWLNGKE YKCKVSNKALPAPIEKTIKAKGQPRE PQVYTLPPSREEMTKNQVSLTCLVKGF YPSDIAVEWESNGQPENNYKTTTPVLD SDGSFFLYSKLTVDKSRWQQGNVFSK SVMHEALHNHYTQKSLSLSPGSGGSG GGGSOVQLVQSGAEVKKPGASVKVSC KASGYTPGYYMYWLRQAPGQGLEW MGWINPDSGDTNYAQKFQGRVTMTTR DTSISTAFMELSLRSDDTAVYYCARE KPRYFDSFYFYYLMDVWGQGTITVTVSS ASTKGPSVFPLAPSSKSTSGGTAAALGC LVKDYFPEPTVTVSWNSGALTSGVHTFP AVLQSSGLYSLESVTVTPSSSLGTQTYI CNVNHKPSNTKVDKKVEPKSC
		(SEQ ID NO: 796)	(SEQ ID NO: 797)	(SEQ ID NO: 798)

			GACATCCAGATGACCCAGTCTCCATC TTCCGTGTCTTCTGTCTGGAGACA GAGTCACCATCACTTGTGCGGCGAGT CAGGATATTAGCAGGTGGTTAGCCT GGTATCAGCAGAAACCAGGAAAGC CCCTAAGCTCTGATTCTGCTGCAT CCAGATTGCAAAGTGGAGTCCCATC AAGGTTACGCGGCAGTGGATCTGGG ACAGATTTCACTCTCACCATCAGCAG CCTGCAGCCTGAAGATTTTGCAATTT ACTATTGTCAACAGGCTAAAAGTTTT CCTCGGACGTTTCGGCCAAGGGACCA AGGTGGAATCAAAACGGACGGTGGC TGCACCATCTGTCTTTCATCTTCCCG CATCTGATGAGCAGTTGAAATCTGG AACTGCCTCTGTTGTGCTGCTGA ATAACTTCTATCCAGAGAGGCCAA AGTACAGTGGAAGGTGGATAACGCC CTCCAATCGGGTAACCTCCAGGAGA GTGTCACAGAGCAGGACAGCAAGGA CAGCACCTACAGCCTCGAAAGCACCC CTGACGCTGAGCAAAAGCAGACTACG AGAAACACAAAGTCTACGCCTGCGA AGTCACCCATCAGGGCCTGAGCTCG CCCGTCACAAAGAGCTTCAACAGGG GAGAGTGT		CAGGTGTGCCGACTCAGCCCCCTTC CCTCTTGCATCTCTGAGCATCAG CCAGTCTACCTGCACCTTACGCAGT GGCATCAATGTTGGTCTCTCCAGGAT CTATTGGTACCAGCAGAAAGCCAGG AGTCTCCCAAGTTTCTCTGAGGTA CACATCAGACTCAGATAAATTGCAG GGCTCTGGAGTCCCCAGCCCTTCTC TGGATCCAAAGATGCTTCGGCCAAT GCAGGACTTTTACTCATCTCTGGGT CCAGTCTGAGGATGAGGCTGACTATT ACTGTATGATTTGGCACAGCAGCGCT GTGGTATTCGGCGGAGGGACCAAAC TGACCGTCTAGGTACGCCAAGGCT GCACCTCGGTCACTCTGTTCCCGC CTCCTCTGAGGAGCTTCAAGCCAACA AGGCCACACTGGTGTGTCTCATCAGT GACTTCTACCCGGGAGCCGTGACAG TGGCCTGGAAGGCAGATAGCAGCCC CGTCAAGCGGGAGTGGAACCCACC ACACCTTCCAAACAAGCAACAACA AGTACGCGGCCAAGAGCTATCTGAG CCTGACGCTGAGCAGTGGAAAGTCC CACAGAAGCTACAGCTGCCAGGTCA CGCATGAAGGAGCACCGTGGAGAA GACAGTGGCCCCCTACAGAATGTTCA		CAGGTGCAGCTGTTGGAGTCTGGGG GAGGCTTGGTCAAGCCTGGAGGGTC CCTGAGACTCTCTGTGCAGCCTCTG GATTCACCTTCAGTACTACTACATG AGCTGGATCCGCCAGGCTCCAGGGA AGGGCTGAGTGGAGTTTCATACATT AGTAGACTGAAAGTATCATCTATT CGTAGACTCTGTGAAGGCCGATT ACCATCTCCAGGACAACGCCAAGA ACTACTGTATCTGCAATGAACAGC CTGAGAGCCGAGGACACGGCCGTGT ATTACTGTGCGAGAGATGTTGGGAG CCACTTTGACTACTGGGCCAGGGA ACCTGTGTACCCGTGCTCTCAGCCTC CACCAGGGCCCATCGGTCTTCCCC TGGCACCTCTCTCCAAAGACCTCT GGGGCACAAGGCCCTGGGTGCC TGGTCAAGGACTACTTCCCCGAAACG GTGACGGTGTGTTGGAACCTCAGGCG CCCTGACCAAGCGGTGACACCTTTC CCGGTGTCTACAGTCTCAGGACT CTACTCCCTCAAGAGCGTGGTGACCG TGCCCTCCAGCAGCTTGGGCACCCAG ACCTACATCTGCAACGTGAATCACA AGCCCAGCAACACCAAGGTGGACAA GAAAGTTGAGCCCCAAATCTTGTGAC AAAACTCACACATGCCACCCGTGCC CAGCACCTGAACCTCTGGGGGGACC GTCAGTCTTCTCTTCCCCCCTAAC CCAGGACACCCCTCATGATCTCCCG ACCCCTGAGGTACATGCGTGGTGGT GGACGTGAGCCACGAAGACCCCTGAG GTCAAGTTCAACTGGTACGTGGACG GCGTGGAGGTGCATAATGCCAAGAC AAAGCCGTGCGAGGAGCAGTACGGC																										
i	P	S	:	5	7	6	2	1-	23	1	0																						
ihu	ant	i-	<h	u	Me	sot	hel	in>	6F	4	V	Hj:	ihu	lg	Gl	zS	EF	L2	*G	K-	K::	(G	4S	)2:	u	ant	i-	<h	uC	D4	0>	21-	23

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+ [an ti- <h uC D4 0> 21- 23 0_33 H9 VL j:: hu LL C2 -K (lg G- Fa b); L M R ID: SS - 30 84 I				CTCGTCGAAAGTCAACTTCGGGAGGG ACCGGGCACTTGGCTGTCTTGTCAA AGATTACTTCCCTGAGCCAGTGACAG TCAGCTGGAATTCCGGTGCCCTCACG TCAGGAGTACATAACATTCCCTGCAGT ATTGCAGTCTCCGGACTCTACTCCC TGGAGTCGGTGGTAACGGTGCCCGAG CTCCAGCTTGGGGACCCAGACGTAC ATTTGTAACGTGAATCACAAACCAA GCAATACTAAGGTAGATAAGAAAGT AGAACCAGAAAGAGCTGC	(SEQ ID NO: 800) (SEQ ID NO: 801)
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0_35	F1	I	V	Hj:	:hu	Ig	G1	z-	C	H1	-	E::	EP	KS	C	+	[an	ti-	<h	u	Me	sot	hel	in>	6F	4	VL	::	hu	KL	C-	SI	76
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[illegible]

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[illegible]

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[illegible]

[illegible]

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[illegible]

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	+ [an ti- <h uC D4 0> 21- 23 0_ 4G 7 VL j:: hu KL C- SI 76 K (dg G- Fa b); L M R ID: SS . 30 84 6				CGAAGTCAACTTCGGGAGGGACCCGC GGCACTTGGCTGTCTTGTCAAAAGATT ACTTCCCCTGAGCCAGTGACAGTCAGC TGGAATTCGGGTGCCCTCACGTCAGG AGTACATACATTCCCTGCGGTATTGC AGTCCTCCGGACTCTACTCCCTGGAG TCGGTGGTAACGGTGCCCCAGCTCCA GCTTGGGGACCCAGACGTACATTGT AACGTGAATCACAAACCAAGCAATA CTAAGGTAGATAAGAAAGTAGAACC GAAGAGCTGC	(SFO ID NO.: 831)
						(SFO ID NO.: 830)
						(SFO ID NO.: 829)

		DIVMTQSPDSLAVSLGERATINCKSSQ SVLYSSNNKNYLAWYQOKPGQPPKLL IYWASTRESGVPDRFSGSGGTDFTLTI SSLQAEDVAVYYCQYYSTPPTFGQG TKVEIKRTVAAPSVFIFPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNAL QSGNSQESVTEQDSKDSITYSLESTLTL SKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC	DIQMTQSPSSLSASVGDILTITCRASQNI TTYLNWYQQKPGKAPNLLISAASRLRS GVPSRFGSGSGTDFTLTISLQPVDFT TFYCQQFTFTTPWTFGQGTKVEIKRTVA APSVFIFPPSDEQLKSGTASVVCLLNNF YPREAKVQWKVDNALQSGNSQESVTE QDSKDSITYSLKSTLTLKADYEKHKV YACEVTHQGLSSPVTKSFNRGEC	EVQLVESGGGLIQPGGSLRLSCAVSGF TVSSKFMTWVRQAPGKGLIEWVSVIYS GKNTYYADSVKGRFTISRDN SKNTLY LQMNSLR AEDTAVYYCARDSSGGWGY FDYWGGQGLTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVS WNSGALTSGVHTFPAVLQSSGLYSLSKS VVTVPSSSLGTQTYICNVNHHKPSNTKV DKKVEPKSCDKTHTCPPCPAPELGGP SVFLFPPKPKDITLMISRTPETVTCVVVD VSHEDPEVKFNWYVDGVEVHNAKTK PCEEQYGSTYRCVSVLTVLHQDWLNG KEYCKVSNKALPAPIEKTISKAKGQP REPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTPPV LDSGGSFFLYSKLTVDKSRWQQGNVF SCSVMH EALHNHYTQKSLSLSPGGGG SGGGGSQVQLVQSGAEVKKPGASVKV SCKASGYTFAGYYMHVWRQAPGQGL EWMGINPDSGGTNFAQQFQGRVTM TRDTSISTAYMEVSRLSRDDTAVFYCA REKITMTGIYFDYWGGQGLTVTVSSAST KGPSVFPLAPSSKSTSGGTAAALGCLVK DYFPEPVTVSWNSGALTSGVHTFPAVL QSSGLYSLESVTVTPSSSLGTQTYICNV NHHKPSNTKVDDKKVEPKSC
	(SEQ ID NO: 832)	(SEQ ID NO: 833)	(SEQ ID NO: 834)	
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E + lan ti- <h uC D4 0> 21- 23 0 _ 29 H1 0 VL]:: hu KL C- SI 76 K (lg G- Fa b); L M R ID: SS - 30				TGCGCCCTCGTCGAAGTCAACTTCGG GAGGACCGCGGCAC TTGGCTGTCTT GTCAAAGATTACTTCCC TGACCAGT GACAGTCAGCTGGAA TTCGGTGCC CTCACGTCAGGAGTACATATCCC TGCGGTATTGCAGTCTCCGGACTCT ACTCCC TGGAGTCGGTGTTAACGGT GCCCAGCTCCAGCTTGGGGACCCAG ACGTACATTTGTACGTGAATCACAA ACCAAGCAATACTAAGGTAGATAAG AAAGTAGAACCCGAAGAGCTGC	(SFO ID NO· 835)	(SFO ID NO· 836)	(SFO ID NO· 837)
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84	DIVMTQSPDSLAVSLGERATINCKSSQ SVLYSSNNKNYLAWYQOKPGQPPKLL IYWASTRESGVPDRFSGSGGTDFTLTI SSLQAEDVAVYYCQYYSTPPTFGQG TKVEIKRTVAAPSVFIFFPSDEQLKSGT ASVVCLLNFFYPREAKVQWKVDNAL QSGNSQESVTEQDSKDSITYSLESTLTL SKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC	DIQMTQSPSSLSASVGDRVTITCRASQ DISNNLAWFQQKPGKPKSLMYAASS LHSGVPSFSGSGGTDFFTITSSLQPE DFATYYCQYNSYPLTFGGGTKVEIRR TVAAPSVFIFFPSDEQLKSGTASVVCLL NNFYPREAKVQWKVDNALQSGNSQE SVTEQDSKDSITYSLKSTLTLTKADYEK HKVYACEVTHQGLSSPVTKSFNRGEC	EVQLVESGGGLIQPGGSLRLSCAVSGF TVSSKFMTWVRQAPGKGLEWVSVIYS GKMYYADSVKGRFTISRDNKSKNTLY LQMNSLRAEDTAVYYCARDSSGGWGY FDYWGGQGLTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVS WNSGALTSGVHTFPAVLQSSGLYSLSKS VVTVPSSSLGTQTYICNVNHKPSNTKV DKKVEPKSCDKTHTCPPCPAPELGGP SVFLFPPKPKDITLMISRTPETVTCVVVD VSHEDPEVKFNWYVDGVEVHNAKTK PCEEQYGSTYRCVSVLTVLHQDWLNG KEYCKVSNKALPAPIEKTISKAKGQP REPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTPPV LDSGGSFFLYSKLTVDKSRWQQGNVF SCSVMHREALHNHYTQKSLSLSPGGGG SGGGGSQVQLVQSGAEVTKPGASVKV SCKASGYTFAGYYMHVWRQAPGQGL EWMGWNPVHSGGTNYAQKFQDRVTM TRDTSINTAYMELSLRSLDDTAVYYC ARERISMVRGVGHNWFAPWGQGTLV TVSSASTKGPSVFPLAPSSKSTSGGTAA LGCLVKDYFPEPVTVSWNSGALTSGV HTFPAVLQSSGLYSLESVVTVPSSSLGT QTYICNVNHKPSNTKVDDKKVEPKSC (SEQ ID NO: 840)
7		(SEQ ID NO: 839)	
	(SEQ ID NO: 838)		
	A		
	A		

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E + [an ti-<h uC D4 0> 21- 23 0 _ 30 A1 2 VL]:: hu LL C2 -K (lg G- Fa b); L M R ID: SS - 30 84 8				AAC TTCGGAGGACCGGCACTT GGCTGTCTTGTCAAAGATTACTCCC TGAGCCAGTGACAGTCAGCTGGAAT TCCGGTGCCCTCACGTCAGGAGTACA TACATTCCCTGCGGTATTGCAGTCCT CCGGACTCTACTCCCTGGAGTCGGTG GTAACGGTGCCCAAGCTCCAGCTTGG GGACCCAGACGTACATTTGTAACTG GAATCACAAACCAAGCAATACTAAG GTAGATAAGAAAGTAGAACCCGAAGA GCTGC	(SEQ ID NO: 843)
					(SEQ ID NO: 842)
					(SEQ ID NO: 841)

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0	33	H9	V	HJ:
				:hu
				Ig
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[illegible]

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			(SEQ ID NO: 857)	(SEQ ID NO: 858)

542

E + [an ti- <h uC D4 0> 21- 23 0 35 F1 1 VL]:: hu LL C2 -K (lg G- Fa b); L M R ID: SS - 30 85 1				AAC TTC GGA GGA CCG GCA CTT GGT GTCTTGTCAAAGATTACTTCCC TGAGCCAGTGACAGTCAGCTGGAAT TCCGGTGCCCTCACGTCAGGAGTACA TACATTCCCTGCGGTATTGCAGTCCT CCGGACTCTACTCCCTGGAGTCGGTG GTAA CGGTGCCCAGCTCCAGCTTGG GGACCCAGACGTACATTTGTAACTG GAATCACAAACCAAGCAATACTAAG GTAGATAAGAAAGTAGAACCGAAGA GCTGC	(SEQ ID NO: 861)
					(SEQ ID NO: 860)
					(SEQ ID NO: 859)

		DIVMTQSPDSLAVSLGERATINCKSSQ SVLYSSNNKNYLAWYQOKPGQPPKLL IYWASTRESGVPDRFSGSGGTDFTLTI SSLQAEDVAVYYCQYYSTPPTFGQG TKVEIKRTVAAPSVFIPPSDEQLKSGT ASVVCLLNNFYPREAKVQWKVDNAL QSGNSQESVTEQDSKDSITYSLESTLTL SKADYEKHKVYACEVTHQGLSSPVTK SFNRGEC	QSALETQPRSVSGSPGQSVTISCTGTSSD VGGYIFVSWYQQHPGKAPKLMYDVS KRPSGVPDFRSGSKSVNTASLTISGLQ AEDETDYYCCSYAGNYTYVFGTGTKV TVLGGPKAAPSVTLFPPSSEELQANKA TLVCLISDFYPGAVTVAWKADSSPVK AGVETTTPSKQSNNKYAAKSYLSLTPE QWKSHRSYSCQVTHEGSTVEKTVAPT ECS	EVQLVESGGGLIQPGSRLSCAVSGF TVSSKFMTWVRQAPGKGLEWVSVIYS GKNTYYADSVKGRFTISRDNKSKNTLY LQMNSLRAEDTAVYYCARDSSGGWGY FDYWGGQGLTVTVSSASTKGPSVFPLAP SSKSTSGGTAALGCLVKDYFPEPVTVS WNSGALTSGVHTFPAVLQSSGLYSLSK VVTVPSSSLGTQTYICNVNHNKPSNTKV DKKVEPKSCDKTHTCPPCPAPELGGP SVFLFPPKPKDITLMISRTPETVTCVVVD VSHEDPEVKFNWYVDGVEVHNNAKTK PCEEQYGSTYRCVSVLTVLHQDWLNG KEYCKVSNKALPAPIEKTISKAKGQP REPQVYTLPPSREEMTKNQVSLTCLVK GFYPSDIAVEWESNGQPENNYKTTPPV LDSGGSFFLYSKLTVDKSRWQQGNVF SCSVMHREALHNHYTQKSLSLSPGGGG SGGGGSQVQLVESGGGVVQPGRSRLR SCAASGFTLSSYGMHWVRQAPGKGLE WVAWIWYDGSNKYYADSVKGRVTISR DNSKNTLYLQMNSLRAEDTAVYYCTR DGRNYYVFDNWGGQGLTVTVSSASTK GPSVFPLAPSSKSTSGGTAALGCLVKD YFPEPVTVSWNSGALTSGVHTFPAVLQ SSGLYSLESVTVPPSSSLGTQTYICNVN HKPSNTKVDKKVEPKSC
	(SEQ ID NO: 862)	(SEQ ID NO: 863)	(SEQ ID NO: 864)	
A				
A				

545

				TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACACAGGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGGAAGGTGGCGGATCCCA GGTACAGCTGCAACAGTCAGGTCCA GGACTGGTGAAGCCCTCGCAGACCC TCTCACTACCTGTGCCATCTCCGGG GACAGTGTCTTAGCAGCCGTACTGC TTGGAACTGGATCAGGCAGTCCCCAT CGAGAGGCCCTTGAGTGGCTGGGAAG GACATACTACAGGTCCAAGTGGTAT CATGATTATTCAGTATCTGTGAAAAG TCGAATCACCATCGACCCAGACACA TCCAAGAACCAGTTCTCCCTGCAGCT GAACCTGTGTACTCCCGAGGACACG GCTGTTTATTATTGTGCAAGAGGGGC TGCTCCCTTTGACTACTGGGGCCAGG GAACCCCTGGTACCGTGTCTCTCAGCA AGCACGAAGGGGCCGTCCGTATTTC CGCTTGCGCCCTCGTCGAAAGTCAACT
0	36	F3	V	HJ:
				:hu
				Ig
				G1
				z-
				C
				H1
				-
				E::
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				VL
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				KL
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				SI
				76
				E

	+ [an ti- <h uC D4 0> 21- 23 0_ 36 F3 VL j:: hu KL C- SI 76 K (dg G- Fa b); L M R ID: SS . 30 85 2			TGGGAGGGACCGGGCACTTGGCT GTCTTGTCAAAGATTACTTCCTGAG CCAGTGACAGTCAGCTGGAATCCG GTGCCCTCACGTCAGGAGTACATAC ATTCCCCTGCGGTATTGCAGTCCTCCG GACTCTACTCCCTGGAGTCGGTGGTA ACGGTGCCCAAGCTCCA GCTTGGGGA CCCAGACGTACATTTGTAACGTGAAT CACAAACCAAGCAATACTAAGGTAG ATAAGAAAAGTAGAACCCGAAGAGCTG C	(SFO ID NO· 867)
					(SFO ID NO· 866)
					(SFO ID NO· 865)

549

				TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACCAGGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGAGGTGGCGGATCCCA GGTGCAAGTGGTGGAGTCTGGGGGA GGCTTAGTCAAGCCTGGAGGGTCCCT GAGACTCTCTGTGCAGCCTCTGAAT TCACCTTCAGTGACTACTACATGAGC TGGATCCGCCAGGCTCCAGGGAAGG GGCTGGAGTGGGTTTCATATATAGT CGAAGTGGTGATACCATCTACTACGC AGACTCTGTGAAGGGCCGATTACCC ATCTCCAGGGACAACGCCAAGAACT CACTGTATCTGCAATGAATGGCCTG CGAGCCGAAGACACGGCCGTGTATT ACTGTGCGAGAGACTTAGCAGCAGG TGCTACAGGGGGCTTGACTGCTGG GGCCAGGGAAACCCCTGGTCACCGTGT CCTCAGCAAGCACGAAGGGGCCGTC CGTATTCCGCTTGCGCCCTCGTCGA
37	A6	V	HJ:	:hu
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[illegible]

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			TACGGCAGCACGTACCGTTGCGTCA GCGTCTCACCGTCTGCACGAGAC TGGCTGAATGGCAAGGAGTACAAGT GCAAGGTGTCCAACAAGCCCTCCC AGCCCCATCGAGAAACCATCTCC AAAGCCAAAGGGCAGCCCCGAGAAC CACAGGTGTACCCCTGCCCCCATCC CGGAGGAGATGACCAAGAACCAGG TCAGCCTGACCTGCCTGGTCAAAGGC TTCTATCCCAGCGACATCGCCGTGGA GTGGGAGAGCAATGGGCAGCCGGAG AACAACTACAAGACCACGCCCTCCCC TGCTGGACTCCGACGGCTCTCTTTC CTCTATAGCAAGCTCACCGTGGACA AGAGCAGGTGGCAGCAGGGGAACGT CTTCTCATGCTCCGTGATGCATGAGG CTCTGCACAACCACTACACGCAGAA GAGCCTCTCCCTGTCTCCGGGTGGTG GCGGATCGGAGAGGTGGCGGATCCCA GGTGCACTGGTGCAGTCTGGGACT GAGGTGAAGAAGCCTGGGGCCTCAG TGAAGGTGCTTGCAAGGCTTCTGGA TACACCTTCCCCGGCTACTATATGCA CTGGGTGCACAGGCCCTGGGACAG GGCTTGAGTGGATGGATGGATCA ACCTGACAGTGGTGACACAAGTA TACACAGAAAGTTTCAGGCGAGGGTC ACCTTGACCAAGGACGCCGTCCGTCA GCACAGCCTACATTGACCTGAACAG GCTGAGATCTGACGACACGGCCGTA TATTACTGTGCGAGAGAGAGGTGTA GGACTACCAACTGCTATTGGACTAC TGGGGCCAGGGAAGTCTGGTCACCG TGTCTCAGCAAGCACGAAGGGGCC GTCCGATTTCCGCTTGCGCCCTCGT
39	C2	V	HJ:
			:hu
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RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLESTLTLSKADYEKHKVYACEVTHQGLSS
PVTKSFNRGEC (SEQ ID NO: 926)

RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLESTLTLSKADYEKHKVYACEVTHQGLSS
PVTKSFNRGEC (SEQ ID NO: 927)

Lambda CL

(SEQ ID NO: 884)

GQPKAAPSVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSNNKYAASSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTecs

GQPKAAPSVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSNNKYAAKSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTecs (SEQ ID NO: 928)

GQPKAAPSVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSNNKYAAESYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTecs (SEQ ID NO: 929)

Common CHI-hinge-CH2-CH3

(SEQ ID NO: 885)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKK
VEPKSCDKTHCTCPAPELGGPSVFLPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYVTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK

Common CHI-hinge-CH2-CH3 (K590G)

(SEQ ID NO: 886)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVTPSSSLGTQTYICNVNHKPSNTKVDKK

VEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYVTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGG

(SEQ ID NO: 887)

GGSGGGGS

(SEQ ID NO: 888)

GGGGSGGGGS

(SEQ ID NO: 889)

GGGGSGGGSGGGGS

(SEQ ID NO: 890)

GGGGSGGGSGGGSGGGGS

(SEQ ID NO: 891)

GGGGQGGGGQ

(SEQ ID NO: 892)
GGGGQGGGGQGGGGQ

(SEQ ID NO: 893)
GGGGQGGGGQGGGGQGGGGQ

MDMRVPAQLLGLLLWLRGARC (SEQ ID NO: 894)
MAWALLLTLLTQGTGSWA (SEQ ID NO: 895)
MTCSPLLTLLIHCTGSWA (SEQ ID NO: 896) MEAPAQLLFLLLLWLPDTTG (SEQ ID NO: 897)
MEWTWRVFLVAAATGAHS (SEQ ID NO: 898)
METPAQLLFLLLWLPDTTG (SEQ ID NO: 899)
METPAQLLFLLLWLPDTTG (SEQ ID NO: 900)
MKHLWFFLLLVAAPRWVLS (SEQ ID NO: 901)
MEWSWVFLFFLSVTTGVHS (SEQ ID NO: 902)

(SEQ ID NO: 903)
Human MSLN v2 NM 013404
MALPTARPLLGSCGTPALGSLFLFLSLGWVQPSRTLGETGQEAAPLDGVLANPPNISSSPRQLLGFFPCAEEVSGLSTERVRELAVALAQNVLST
EQLRCLAHRLSEPPEDLDALPLDLLFLNPDFAFSGPQACTRFFSRITKANVDLLPRGAPERQRLPAALACWGVRSLLSEADVRLGGLACDLPGR

FVAESA EVLLPRLVSCPGPLDQDQQAARAALQGCGPPYPGPSTWSVSTMDALRGLLPVLGQPIRSIPQGI VAAWRQRSSRDPSWRQPERILRPFR
 REVEKTACPSGKKAREIDESLIFYKKWELEACVDAAALLATQMDRVNAIPFTYEQLDVLKHKLDLDELYPQGYPESVIQHLGYLFLKMSPEDIRKWNVT
 SLETLKALLEVNKGHEMSPQAPRRPLPQVATLIDRFVKGRGQDKDLDLTAFYPGYLCSLSPEELSSVPPSSIWA VVRPQDLDTC DPRQLDVL YPK
 ARLAFQNMNGSEYFVKIQSFLGGAPTEDLKALSQQNVSMDLATFMKLRITDAVLPLTVAEVQKLLGPHVEGLKA
 EERHRPV RDWILRQRQDDLDLTLGLQGGINGYLVLDLSMQEALSGTPCLLGP GPVLTVLALLLASTLA

(SEQ ID NO: 904)

Human MSLN v6 AY743922

MALPTARPLLGSCGTPALGSLFLFLFSLGWVQPSRTLGETGQEAAPLDGVLANPPNISSLSRQLLGFPCA EVSGLSTERVRELAV ALAQKNVKLST
 EQLRCLAHRLSEPPEDLDALPLDLLFLNPD AFSGPQACTHFFSRITKANVDLLPRGAPERQRLPAAALACWGVRSLLSEADVRALGGLACDLPGR
 FVAESA EVLLPRLVSCPGPLDQDQQAARAALQGCGPPYPGPSTWSVSTMDALRGLLPVLGQPIRSIPQGI VAAWRQRSSRDPSWRQPERILRPFR
 REVEKTACPSGKKAREIDESLIFYKKWELEACVDAAALLATQMDRVNAIPFTYEQLDVLKHKLDLDELYPQGYPESVIQHLGYLFLKMSPEDIRKWNVT
 SLETLKALLEVNKGHEMSPQVATLIDRFVKGRGQDKDLDLTAFYPGYLCSLSPEELSSVPPSSIWA VVRPQDLDTC DPRQLDVL YPKARLAFQNM
 NGSEYFVKIQSFLGGAPTEDLKALSQQNVSMDLATFMKLRITDAVLPLTVAEVQKLLGPHVEGLKAEERHRPV R
 DWILRQRQDDLDLTLGLQGGINGYLVLDLSVQEALSGTPCLLGP GPVLTVLALLLASTLA

(SEQ ID NO: 905)

cyno MSLN macaque v1 LMR C52457

MALPMARPLSGSCGTPALGSLFLFLFSLGWVQPSRVLAGETRQEAAPLDGILTNAPDIASLSRQLLGFCTVEVSGLSTELVQELAV ALGQKNVKLS
 AEQLRCLAHRLSEPPEDLDALPLDLLFLNPD AFSGPQACTHFFSRVAKANVDLLPRGAPERQRLPAAALTCWGVRSLLSEADVRALGGLACDLP
 GRFVAESA EWLPRLVRCLGPLDQDQQAARAALQRGGPPYPGPSTWSISTLDDLQSLLPVLGQPVHSIPQGI LAAWRQRSSRDPSWQQPEQTVLRP
 RFRRDVERITTCPEKEVHEIDENLIFYKKRELEACVDAAALLAQMDRVD AIPFTYEQLDVLKHKLDLDELYPQGYPESVIRHLGHLFLKMSPEDIRKWN
 VTSLETLKALLKVSKGHEMSAQVATLIDRVWGRGQDKDLDLTAFCPGCLCSLSPERLSSVPPSVIGAVVRPQDLDTCGPRQLDVL YPKARLAFQ
 NMSGSEYFVKIRPFLGGAPTEDLKALSQQNVSMDLATFMKLRREAVLPLTVAEVQKLLGPHVEGLKVVEEQHSPVRDWILKQRQDDLDLTLGLGLQG
 GIPNGYLILDL SVREALSGTPCLLGP GPVLTILALLLASTLA

Kappa CL S176E

(SEQ ID NO: 906)
RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLESTLTLSKADYEKHKVYACEVTHQGLSS
PVTKSFNRGEC

Kappa CL S176K
(SEQ ID NO: 907)
RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLESTLTLSKADYEKHKVYACEVTHQGLSS
PVTKSFNRGEC

Lambda CL S176K
(SEQ ID NO: 908)
GQPKAAPSVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSNNKYAAKSYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTSCS

Lambda CL S176E
(SEQ ID NO: 909)
GQPKAAPSVTLFPPSSEELQANKATLVCLISDFYPGAVTVAWKADSSPVKAGVETTTTPSKQSNNKYAAESYLSLTPEQWKSHRSYSCQVTHEGSTV
EKTVAPTSCS

(SEQ ID NO: 910)
> XM_005590816.2 (without N-terminal 11aa) cyno Mesothelin
MALPMARPLSGSCGTPALGSLFLFLFSLGWVQPSRVLAGETRQEAAPLDGILTNAPDIASLSPRQLLGFTTCVEVSGLSTELVQELAVALGQKNVKLS
AEQLRCLAHRLSEPPEDLDALPLDLLFLNPDAFSGPQACTHFFSRVAKANVDLLPRGAPERQRLPAAALTCWGVRGSLSEADVRLGGLACDLP
GRFVAESAEEVVLPRVLVRCGLPDDQDQEAARAALQRGGPPYGPSTWSISTLDDLQSLLPVLGQPVIIHSPQGILAAWRQRSSRDPSWQQPEQTVLR
PRFRDVERTTCPEKEVHEIDESLIFYKKRELEACVDAALLAAQMDRVDAIPFTYEQLDV/LKHKLDELYPQGYPESVIRHLGHLFLKMSPEDIRKW

NVTSLETLKALLKVSKGHEMSAQVATLIDRVVVGRLQDKDTADTLTAFCPGCLCSLSPERLSSVPPSIIGAVRPQDLDTGPRQLDVLVYPKARLAF
QNMGSSEYFVKIRPFLGGAPTEDLKALSQQNVSMDLATFMKLRREAVPLSVAEVQKLLGPHVEGLKVEEQHSPVRDWILKQRQDDLDLTLGLGLQ
GGIPNGYLILDLSVREALSGTPCLLGGPVLTVLALLASTLA

(SEQ ID NO: 911)

>XP_005569274.1 PREDICTED: tumor necrosis factor receptor superfamily member 5 isoform X1 [Macaca fascicularis]

MVRLPLQCVLWGCLLTAVYPEPTACREKQYLINSQCCSLCQPGQKLVSDCTEFTETECCLPCGESEFLDTWNETRCHQHKYCDPNLGLRVQKGT
SETDTICTCEEGLHCTSESCVPHRSCLPGFGVKQIATGVSDTICEPCPVGFFSNVSSAFCKRPWTSCTKDLVVOAGTNKTDVVCGPQDRQRA
LVVPICLGILFVILLVLFVISESEKVAKKPNDKVPHPKQEPQENFPDDLPGSNPAAPVQETLHGCPVTQEDGKESISVQERQ

(SEQ ID NO: 912)

Common CHI1-hinge-CH2-CH3 (S176K)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLKSVVTVPSSSLGTQTYICNVNHKPSNTKVDKK
VEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYVTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVVFSCVMHEALHNHYTQKSLSLSPGG

(SEQ ID NO: 913)

Common CHI1-hinge-CH2-CH3 (S176E)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLKSVVTVPSSSLGTQTYICNVNHKPSNTKVDKK
VEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH

QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGG

(SEQ ID NO: 914)

Common CHI-hinge-CH2-CH3 (S176K/K590G)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLKSVVTVPSSSLGTQTYICNVNHKPSNTKVDKK
VEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGK

(SEQ ID NO: 915)

Common CHI-hinge-CH2-CH3 (S176E/K590G)

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLKSVVTVPSSSLGTQTYICNVNHKPSNTKVDKK
VEPKSCDKTHITCPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPCEEQYGSTYRCVSVLTVLH
QDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFL
YSKLTVDKSRWQQGNVFCFSVMHEALHNHYTQKSLSLSPGK

CLAIMS

We claim:

1. A multispecific antibody construct comprising:

(i) a first antibody comprising two light chains and two heavy chains, wherein

the light chains comprise a first variable region (VL1) and a light chain constant region (CL);

the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and

CH3 regions;

the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and

(ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,

wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and

wherein the first antibody specifically binds to and agonizes human CD40 (SEQ ID NO: 1) and the scFv specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2).

2. The antibody construct of claim 1, wherein the two light chains are identical and the two heavy chain fusion proteins are identical.

3. The antibody construct according to any preceding claim, wherein the heavy chain comprises an amino acid substitution selected from the group consisting of:

(i) N297G or N297A;

(ii) L234A and L235A; and

(iii) R292C and V302C;

wherein the amino acid numbering is EU numbering according to Kabat.

4. The antibody construct according to claim 3, wherein the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

5. The antibody construct of any preceding claim, wherein:

the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 58, 59, and 60, respectively;
SEQ ID NOs: 64, 65, and 66, respectively;
SEQ ID NOs: 70, 71, and 72, respectively;
SEQ ID NOs: 76, 77, and 78, respectively;
SEQ ID NOs: 82, 83, and 84, respectively;
SEQ ID NOs: 88, 89, and 90, respectively;
SEQ ID NOs: 94, 95, and 96, respectively;
SEQ ID NOs: 100, 101, and 102, respectively;
SEQ ID NOs: 106, 107, and 108, respectively;
SEQ ID NOs: 112, 113, and 114, respectively;
SEQ ID NOs: 118, 119, and 120, respectively;
SEQ ID NOs: 124, 125, and 126, respectively; and
SEQ ID NOs: 130, 131, and 132, respectively;

the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

SEQ ID NOs: 136, 137, and 138, respectively;
SEQ ID NOs: 142, 143, and 144, respectively;
SEQ ID NOs: 148, 149, and 150, respectively;
SEQ ID NOs: 154, 155, and 156, respectively;
SEQ ID NOs: 160, 161, and 162, respectively;
SEQ ID NOs: 166, 167, and 168, respectively;
SEQ ID NOs: 172, 173, and 174, respectively;
SEQ ID NOs: 178, 179, and 180, respectively;
SEQ ID NOs: 184, 185, and 186, respectively;
SEQ ID NOs: 190, 191, and 192, respectively;
SEQ ID NOs: 196, 197, and 198, respectively;
SEQ ID NOs: 202, 203, and 204, respectively; and
SEQ ID NOs: 208, 209, and 210, respectively;

the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 230, 231, and 232, respectively;
SEQ ID NOs: 236, 237, and 238, respectively;
SEQ ID NOs: 242, 243, and 244, respectively; and
SEQ ID NOs: 248, 249, and 250, respectively;

and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

SEQ ID NOs: 254, 255, and 256, respectively;
SEQ ID NOs: 260, 261, and 262, respectively;

SEQ ID NOs: 266, 267, and 268, respectively; and

SEQ ID NOs: 272, 273, and 274, respectively.

6. The antibody construct of any preceding claim, wherein

1) the VL1 and VH1 are selected from the group consisting of:

a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;

and

2) the VL2 and VH2 are selected from the group consisting of:

a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a

CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

7. The antibody construct of any preceding claim, wherein:

the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

8. The antibody construct of any preceding claim, wherein:

1) the VL1 and VH1 are selected from the group consisting of:

- a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;
 - b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;
 - c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;
 - d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;
 - e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;
 - f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;
 - g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;
 - h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;
 - i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;
 - j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;
 - k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;
 - l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and
 - m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;
- and

2) the VL2 and VH2 are selected from the group consisting of:

- a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;
- b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;
- c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and
- d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.

9. The antibody construct according to any preceding claim, wherein the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.
10. The antibody construct according to any preceding claim, wherein the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.
11. The antibody construct according to any preceding claim, wherein the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.
12. The antibody construct according to any preceding claim, wherein the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.
13. The antibody construct according to any preceding claim, wherein the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.
14. The antibody construct according to any preceding claim, wherein
the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 286, 290, 294, 298, 302, 306, 310, 314, 318, 322, 326, 330, 336, 342, 346, 350, 354, 358, 362, 366, 370, 374, and 378; and
the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID Nos: 285, 289, 293, 297, 301, 305, 309, 313, 317, 321, 325, 329, 333, 337, 341, 345, 349, 353, 357, 361, 365, 369, 373, and 377.
15. The antibody construct according to any preceding claim, wherein the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:
SEQ ID NO: 286 and SEQ ID NO: 285, respectively;
SEQ ID NO: 290 and SEQ ID NO: 289, respectively;
SEQ ID NO: 294 and SEQ ID NO: 293, respectively;
SEQ ID NO: 298 and SEQ ID NO: 297, respectively;
SEQ ID NO: 302 and SEQ ID NO: 301, respectively;
SEQ ID NO: 306 and SEQ ID NO: 305, respectively;
SEQ ID NO: 310 and SEQ ID NO: 309, respectively;
SEQ ID NO: 314 and SEQ ID NO: 313, respectively;
SEQ ID NO: 318 and SEQ ID NO: 317, respectively;
SEQ ID NO: 322 and SEQ ID NO: 321, respectively;
SEQ ID NO: 326 and SEQ ID NO: 325, respectively;

SEQ ID NO: 330 and SEQ ID NO: 329, respectively;
SEQ ID NO: 334 and SEQ ID NO: 333, respectively;
SEQ ID NO: 338 and SEQ ID NO: 337, respectively;
SEQ ID NO: 342 and SEQ ID NO: 341, respectively;
SEQ ID NO: 346 and SEQ ID NO: 345, respectively;
SEQ ID NO: 350 and SEQ ID NO: 349, respectively;
SEQ ID NO: 354 and SEQ ID NO: 353, respectively;
SEQ ID NO: 358 and SEQ ID NO: 357, respectively;
SEQ ID NO: 362 and SEQ ID NO: 361, respectively;
SEQ ID NO: 366 and SEQ ID NO: 365, respectively;
SEQ ID NO: 370 and SEQ ID NO: 369, respectively;
SEQ ID NO: 374 and SEQ ID NO: 373, respectively; and
SEQ ID NO: 378 and SEQ ID NO: 377, respectively.

16. A polynucleotide encoding the light chain of the antibody construct according to any preceding claim.

17. A polynucleotide encoding the heavy chain fusion protein of the antibody construct according to any preceding claim.

18. A vector comprising the polynucleotide according to claim 16, the polynucleotide according to claim 17, or both.

19. A host cell transformed or transfected with the polynucleotide according to claim 16 and the polynucleotide according to claim 17.

20. A process for producing the antibody construct according to any one of claims 1-15, said process comprising culturing a host cell comprising a polynucleotide encoding the light chain and also comprising a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

21. A pharmaceutical composition comprising the antibody construct according to any one of claims 1-15 and a carrier, stabilizer, excipient, diluent, solubilizer, surfactant, emulsifier, preservative or adjuvant.

22. A method for treating or ameliorating a solid tumor disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof an effective amount of the antibody construct according to any one of claims 1-15.

23. The method according to claim 22, wherein the solid tumor disease is selected from the group consisting of: ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer disease or a metastatic cancer disease derived from any of the foregoing.

24. A kit comprising the antibody construct according to any one of claims 1-15 and, optionally, directions for use.

25. A multispecific antibody construct comprising:

(i) a first antibody comprising two light chains and two heavy chains, wherein

the light chains comprise a first variable region (VL1) and a light chain constant region (CL);

the heavy chains comprise a first heavy variable region (VH1) and CH1, hinge, CH2, and

CH3 regions;

the heavy chains comprise at least one amino acid substitution that results in a reduced binding affinity of the heavy chain for human Fc gamma RI receptor as compared with an unsubstituted heavy chain; and

(ii) a scFv comprising a second light chain variable region (VL2) and a second heavy chain variable region (VH2) of a second antibody, wherein the VL2 and the VH2 are connected via a first peptide linker,

wherein the scFv is fused at its amino terminus to the carboxyl terminus of each of the heavy chains through a second peptide linker such that a heavy chain fusion protein is formed; and

wherein the first antibody specifically binds to human mesothelin (MSLN) (SEQ ID NO: 2) and the scFv specifically binds to and agonizes human CD40 (SEQ ID NO: 1).

26. The antibody construct of claim 25, wherein the two light chains are identical and the two heavy chain fusion proteins are identical.

27. The antibody construct according to any one of claims 25-26, wherein the heavy chain comprises an amino acid substitution selected from the group consisting of:

(i) N297G or N297A;

(ii) L234A and L235A; and

(iii) R292C and V302C;

wherein the amino acid numbering is EU numbering according to Kabat.

28. The antibody construct according to claim 27, wherein the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

29. The antibody construct of any one of claims 25-28, wherein:

the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 230, 231, and 232, respectively;

SEQ ID NOs: 236, 237, and 238, respectively;

SEQ ID NOs: 242, 243, and 244, respectively; and

SEQ ID NOs: 248, 249, and 250, respectively;

the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting

of:

SEQ ID NOs: 254, 255, and 256, respectively;

SEQ ID NOs: 260, 261, and 262, respectively;

SEQ ID NOs: 266, 267, and 268, respectively; and

SEQ ID NOs: 272, 273, and 274, respectively;

the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 58, 59, and 60, respectively;

SEQ ID NOs: 64, 65, and 66, respectively;

SEQ ID NOs: 70, 71, and 72, respectively;

SEQ ID NOs: 76, 77, and 78, respectively;

SEQ ID NOs: 82, 83, and 84, respectively;

SEQ ID NOs: 88, 89, and 90, respectively;

SEQ ID NOs: 94, 95, and 96, respectively;

SEQ ID NOs: 100, 101, and 102, respectively;

SEQ ID NOs: 106, 107, and 108, respectively;

SEQ ID NOs: 112, 113, and 114, respectively;

SEQ ID NOs: 118, 119, and 120, respectively;

SEQ ID NOs: 124, 125, and 126, respectively; and

SEQ ID NOs: 130, 131, and 132, respectively;

and

the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting

of:

SEQ ID NOs: 136, 137, and 138, respectively;

SEQ ID NOs: 142, 143, and 144, respectively;

SEQ ID NOs: 148, 149, and 150, respectively;

SEQ ID NOs: 154, 155, and 156, respectively;

SEQ ID NOs: 160, 161, and 162, respectively;

SEQ ID NOs: 166, 167, and 168, respectively;
SEQ ID NOs: 172, 173, and 174, respectively;
SEQ ID NOs: 178, 179, and 180, respectively;
SEQ ID NOs: 184, 185, and 186, respectively;
SEQ ID NOs: 190, 191, and 192, respectively;
SEQ ID NOs: 196, 197, and 198, respectively;
SEQ ID NOs: 202, 203, and 204, respectively; and
SEQ ID NOs: 208, 209, and 210, respectively.

30. The antibody construct of any one of claims 25-29, wherein

1) the VL1 and VH1 are selected from the group consisting of:

a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;

and

2) the VL2 and VH2 are selected from the group consisting of:

a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

31. The antibody construct of any one of claims 25-30, wherein:

the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

32. The antibody construct of any one of claims 25-31, wherein:

1) the VL1 and VH1 are selected from the group consisting of:

- a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;
 - b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;
 - c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and
 - d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;
- and

2) the VL2 and VH2 are selected from the group consisting of:

- a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;
- b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;
- c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;
- d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;
- e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;
- f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;
- g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;
- h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;
- i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;
- j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;

- k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;
- l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and
- m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.

33. The antibody construct according to any one of claims 25-32, wherein the CL of the light chain is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

34. The antibody construct according to any one of claims 25-33, wherein the CH1-hinge-CH2-CH3 of the heavy chain is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

35. The antibody construct according to any one of claims 25-34, wherein the first peptide linker is selected from the group consisting of SEQ ID NOs: 888-893.

36. The antibody construct according to any one of claims 25-35, wherein the second peptide linker is selected from the group consisting of SEQ ID NOs: 887-893.

37. The antibody construct according to any one of claims 25-36, wherein the first peptide linker comprises SEQ ID NO: 889 and the second peptide linker comprises SEQ ID NO: 887.

38. The antibody construct according to any one of claims 25-37, wherein
the light chain comprises a sequence selected from the group consisting of SEQ ID NOs: 382, 386, 390, 394, 398, 402, 406, 410, 414, 418, 422, 426, 430, 434, 438, 442, 446, and 450; and
the heavy chain fusion protein comprises a sequence selected from the group consisting of SEQ ID NOs: 381, 385, 389, 393, 397, 401, 405, 409, 413, 417, 421, 425, 429, 433, 437, 441, 445, and 449.

39. The antibody construct according to any one of claims 25-38, wherein the light chain and the heavy chain fusion protein comprise polypeptides comprising amino acid sequences selected from the group consisting of:

- SEQ ID NO: 382 and SEQ ID NO: 381, respectively;
- SEQ ID NO: 386 and SEQ ID NO: 385, respectively;
- SEQ ID NO: 390 and SEQ ID NO: 389, respectively;
- SEQ ID NO: 394 and SEQ ID NO: 393, respectively;
- SEQ ID NO: 398 and SEQ ID NO: 397, respectively;
- SEQ ID NO: 402 and SEQ ID NO: 401, respectively;
- SEQ ID NO: 406 and SEQ ID NO: 405, respectively;
- SEQ ID NO: 410 and SEQ ID NO: 409, respectively;

SEQ ID NO: 414 and SEQ ID NO: 413, respectively;
SEQ ID NO: 418 and SEQ ID NO: 417, respectively;
SEQ ID NO: 422 and SEQ ID NO: 421, respectively;
SEQ ID NO: 426 and SEQ ID NO: 425, respectively;
SEQ ID NO: 430 and SEQ ID NO: 429, respectively;
SEQ ID NO: 434 and SEQ ID NO: 433, respectively;
SEQ ID NO: 438 and SEQ ID NO: 437, respectively;
SEQ ID NO: 442 and SEQ ID NO: 441, respectively;
SEQ ID NO: 446 and SEQ ID NO: 445, respectively; and
SEQ ID NO: 450 and SEQ ID NO: 449, respectively.

40. A polynucleotide encoding the light chain of the antibody construct according to any one of claims 25-39.

41. A polynucleotide encoding the heavy chain fusion protein of the antibody construct according to any one of claims 25-39.

42. A vector comprising the polynucleotide according to claim 40, the polynucleotide according to claim 41, or both.

43. A host cell transformed or transfected with the polynucleotide according to claim 40 and the polynucleotide according to claim 42.

44. A process for producing the antibody construct according to any one of claims 25-39, said process comprising culturing a host cell comprising a polynucleotide encoding the light chain and also comprising a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

45. A pharmaceutical composition comprising the antibody construct according to any one of claims 25-39 and a carrier, stabilizer, excipient, diluent, solubilizer, surfactant, emulsifier, preservative or adjuvant.

46. A method for treating or ameliorating a solid tumor disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof an effective amount of the antibody construct according to any one of claims 25-39.

47. The method according to claim 46, wherein the solid tumor disease is selected from the group consisting of: ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer disease or a metastatic cancer disease derived from any of the foregoing.

48. A kit comprising the antibody construct according to any one of claims 25-39 and, optionally, directions for use.

49. A multispecific antibody construct comprising:

a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

wherein:

the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

50. A multispecific antibody construct comprising:

a) two identical heavy chain fusion proteins each comprising a first heavy chain variable region (VH1) and a first CH1 domain, wherein the first CH1 domain is linked to a hinge-CH2-CH3 polypeptide, and wherein the hinge-CH2-CH3 polypeptide is linked to a second heavy chain variable region (VH2), wherein the VH2 is linked to a second CH1 domain; wherein

i) the VH1 or first CH1 domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 39, 44, and 183 using EU numbering; and

ii) the VH2 or second CH1 domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of a residue that corresponds to positions 39, 44, and 183 using EU numbering; and

b) a second polypeptide comprising a first light chain, wherein the first light chain comprises a first light chain variable region (VL1) and a first CL region; and wherein the VL1 or first CL domain comprises at least one amino acid substitution to introduce a positively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering; and

c) a third polypeptide comprising a second light chain, wherein the second light chain comprises a second light chain variable region (VL2) and a second CL region; and wherein the VL2 or second CL domain comprises at least one amino acid substitution to introduce a negatively charged amino acid at a residue selected from the group consisting of positions 38, 100, and 176 using EU numbering;

wherein the VH1 and VL1 interact to bind a first antigen and wherein the VH2 and VL2 interact to bind a second antigen;

wherein:

the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human mesothelin ("MSLN"; SEQ ID NO: 2); or

the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1).

51. The antibody construct according to any one of claims 49 or 50, wherein the hinge-CH2-CH3 polypeptide is linked to the VH2 via a peptide linker.

52. The antibody construct according to 51, wherein the peptide linker comprises a sequence selected from the group consisting of (Gly₃Ser)₂ (SEQ ID NO: 916), (Gly₄Ser)₂ (SEQ ID NO: 888), (Gly₃Ser)₃ (SEQ ID NO: 917), (Gly₄Ser)₃ (SEQ ID NO: 889), (Gly₃Ser)₄ (SEQ ID NO: 918), (Gly₄Ser)₄ (SEQ ID NO: 890), (Gly₃Ser)₅ (SEQ ID NO: 919), (Gly₄Ser)₅ (SEQ ID NO: 920), (Gly₃Ser)₆ (SEQ ID NO: 921), and (Gly₄Ser)₆ (SEQ ID NO: 922).

53. The antigen binding protein according to claim 49, wherein

- a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;
- b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;
- c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering; and
- d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering.

54. The antigen binding protein according to claim 53, wherein

- a) the first CH1 domain comprises a S183K mutation using EU numbering;
- b) the second CH1 domain comprises a S183E mutation using EU numbering;
- c) the first CL domain comprises a S176E mutation using EU numbering; and
- d) the second CL domain comprises a S176K mutation using EU numbering.

55. The antigen binding protein according to claim 53, wherein

- a) the VH1 comprises a Q39K mutation and the first CH1 domain comprises a S183K mutation using EU numbering;
- b) the VH2 comprises a Q39E mutation and the second CH1 domain comprises a S183E mutation using EU numbering;
- c) the VL1 comprises a Q38E mutation and the first CL domain comprises a S176E mutation using EU numbering; and
- d) the VL2 comprises a Q38K mutation and the second CL domain comprises a S176K mutation using EU numbering.

56. The antigen binding protein according to claim 53, wherein

- a) the first CH1 domain comprises G44K and S183K mutations using EU numbering;
- b) the second CH1 domain comprises G44E and S183E mutations using EU numbering;
- c) the first CL domain comprises G100E and S176E mutations using EU numbering; and
- d) the second CL domain comprises G100K and S176K mutations using EU numbering.

57. The antigen binding protein according to claim 50, wherein

- a) the VH1 or first CH1 domain comprises a mutation selected from the group consisting of Q39E, G44E, and S183E using EU numbering;
- b) the VH2 or second CH1 domain comprises a mutation selected from the group consisting of Q39K, G44K, and S183K using EU numbering;

- c) the VL1 or first CL domain comprises a mutation selected from the group consisting of Q38K, G100K, and S176K using EU numbering; and
- d) the VL2 or second CL domain comprises a mutation selected from the group consisting of Q38E, G100E, and S176E using EU numbering.

58. The antigen binding protein according to claim 57, wherein

- a) the first CH1 domain comprises a S183E mutation using EU numbering;
- b) the second CH1 domain comprises a S183K mutation using EU numbering;
- c) the first CL domain comprises a S176K mutation using EU numbering; and
- d) the second CL domain comprises a S176E mutation using EU numbering.

59. The antigen binding protein according to claim 57, wherein

- a) the VH1 comprises a Q39E mutation and the first CH1 domain comprises a S183E mutation using EU numbering;
- b) the VH2 comprises a Q39K mutation and the second CH1 domain comprises a S183K mutation using EU numbering;
- c) the VL1 comprises a Q38K mutation and the first CL domain comprises a S176K mutation using EU numbering; and
- d) the VL2 comprises a Q38E mutation and the second CL domain comprises a S176E mutation using EU numbering.

60. The antigen binding protein according to claim 57, wherein

- a) the first CH1 domain comprises G44E and S183E mutations using EU numbering;
- b) the second CH1 domain comprises G44K and S183K mutations using EU numbering;
- c) the first CL domain comprises G100K and S176K mutations using EU numbering; and
- d) the second CL domain comprises G100E and S176E mutations using EU numbering.

61. The antibody construct according to any one of claims 49-50, wherein the hinge-CH2-CH3 polypeptide comprises an amino acid substitution selected from the group consisting of:

- (i) N297G or N297A;
- (ii) L234A and L235A; and
- (iii) R292C and V302C;

wherein the amino acid numbering is EU numbering according to Kabat.

62. The antibody construct according to claim 61, wherein the hinge-CH2-CH3 polypeptide comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

63. The antibody construct of any one of claims 49-62, wherein the first antigen is human CD40 (SEQ ID NO: 1) and the second antigen is human MSLN (SEQ ID NO: 2); and

the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 58, 59, and 60, respectively;
SEQ ID NOs: 64, 65, and 66, respectively;
SEQ ID NOs: 70, 71, and 72, respectively;
SEQ ID NOs: 76, 77, and 78, respectively;
SEQ ID NOs: 82, 83, and 84, respectively;
SEQ ID NOs: 88, 89, and 90, respectively;
SEQ ID NOs: 94, 95, and 96, respectively;
SEQ ID NOs: 100, 101, and 102, respectively;
SEQ ID NOs: 106, 107, and 108, respectively;
SEQ ID NOs: 112, 113, and 114, respectively;
SEQ ID NOs: 118, 119, and 120, respectively;
SEQ ID NOs: 124, 125, and 126, respectively; and
SEQ ID NOs: 130, 131, and 132, respectively;

the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

SEQ ID NOs: 136, 137, and 138, respectively;
SEQ ID NOs: 142, 143, and 144, respectively;
SEQ ID NOs: 148, 149, and 150, respectively;
SEQ ID NOs: 154, 155, and 156, respectively;
SEQ ID NOs: 160, 161, and 162, respectively;
SEQ ID NOs: 166, 167, and 168, respectively;
SEQ ID NOs: 172, 173, and 174, respectively;
SEQ ID NOs: 178, 179, and 180, respectively;
SEQ ID NOs: 184, 185, and 186, respectively;
SEQ ID NOs: 190, 191, and 192, respectively;
SEQ ID NOs: 196, 197, and 198, respectively;
SEQ ID NOs: 202, 203, and 204, respectively; and
SEQ ID NOs: 208, 209, and 210, respectively;

the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 230, 231, and 232, respectively;
SEQ ID NOs: 236, 237, and 238, respectively;
SEQ ID NOs: 242, 243, and 244, respectively; and
SEQ ID NOs: 248, 249, and 250, respectively;

and the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

SEQ ID NOs: 254, 255, and 256, respectively;

SEQ ID NOs: 260, 261, and 262, respectively;

SEQ ID NOs: 266, 267, and 268, respectively; and

SEQ ID NOs: 272, 273, and 274, respectively.

64. The antibody construct of claim 63, wherein

1) the VL1 and VH1 are selected from the group consisting of:

a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

e) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

f) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

g) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

h) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

i) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

j) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

k) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

l) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

m) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively;

and

2) the VL2 and VH2 are selected from the group consisting of:

a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH2 comprising a

CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively.

65. The antibody construct of claim 63, wherein:

the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53;

the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54;

the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226.

66. The antibody construct of claim 65, wherein:

1) the VL1 and VH1 are selected from the group consisting of:

- a) a VL1 comprising SEQ ID NO: 5 and a VH1 comprising SEQ ID NO: 6;
 - b) a VL1 comprising SEQ ID NO: 9 and a VH1 comprising SEQ ID NO: 10;
 - c) a VL1 comprising SEQ ID NO: 13 and a VH1 comprising SEQ ID NO: 14;
 - d) a VL1 comprising SEQ ID NO: 17 and a VH1 comprising SEQ ID NO: 18;
 - e) a VL1 comprising SEQ ID NO: 21 and a VH1 comprising SEQ ID NO: 22;
 - f) a VL1 comprising SEQ ID NO: 25 and a VH1 comprising SEQ ID NO: 26;
 - g) a VL1 comprising SEQ ID NO: 29 and a VH1 comprising SEQ ID NO: 30;
 - h) a VL1 comprising SEQ ID NO: 33 and a VH1 comprising SEQ ID NO: 34;
 - i) a VL1 comprising SEQ ID NO: 37 and a VH1 comprising SEQ ID NO: 38;
 - j) a VL1 comprising SEQ ID NO: 41 and a VH1 comprising SEQ ID NO: 42;
 - k) a VL1 comprising SEQ ID NO: 45 and a VH1 comprising SEQ ID NO: 46;
 - l) a VL1 comprising SEQ ID NO: 49 and a VH1 comprising SEQ ID NO: 50; and
 - m) a VL1 comprising SEQ ID NO: 53 and a VH1 comprising SEQ ID NO: 54;
- and

2) the VL2 and VH2 are selected from the group consisting of:

- a) a VL2 comprising SEQ ID NO: 213 and a VH2 comprising SEQ ID NO: 214;
- b) a VL2 comprising SEQ ID NO: 217 and a VH2 comprising SEQ ID NO: 218;
- c) a VL2 comprising SEQ ID NO: 221 and a VH2 comprising SEQ ID NO: 222; and

d) a VL2 comprising SEQ ID NO: 225 and a VH2 comprising SEQ ID NO: 226.

67. A polynucleotide encoding the first light chain of the antibody construct according to any one of claims 49-66 and 76-80.

68. A polynucleotide encoding the second light chain of the antibody construct according to any one of claims 49-66 and 76-80.

69. A polynucleotide encoding the heavy chain fusion protein of the antibody construct according to any one of claims 49-66 and 76-80.

70. A vector comprising:

- a) the polynucleotide according to claim 67,
- b) the polynucleotide according to claim 68,
- c) the polynucleotide according to claim 69, or
- d) any combination of a), b), and c).

71. A host cell transformed or transfected with the polynucleotide according to claim 67, the polynucleotide according to claim 68 and the polynucleotide according to claim 69.

72. A process for producing the antibody construct according to any one of claims 49-66 and 76-80, said process comprising culturing a host cell comprising a polynucleotide encoding the first light chain, a polynucleotide encoding the second light chain and a polynucleotide encoding the heavy chain fusion protein under conditions allowing the expression of the antibody construct, and recovering the produced antibody construct from the culture.

73. A pharmaceutical composition comprising the antibody construct according to any one of claims 49-66 and 76-80 and a carrier, stabilizer, excipient, diluent, solubilizer, surfactant, emulsifier, preservative or adjuvant.

74. A method for treating or ameliorating a solid tumor disease or a metastatic cancer disease, comprising the step of administering to a subject in need thereof an effective amount of the antibody construct according to any one of claims 49-66 and 76-80.

75. The method according to claim 74, wherein the solid tumor disease is selected from the group consisting of: ovarian cancer, pancreatic cancer, mesothelioma, lung cancer, gastric cancer and triple negative breast cancer disease or a metastatic cancer disease derived from any of the foregoing.

76. A kit comprising the antibody construct according to any one of claims 49-66 and 76-80 and, optionally, directions for use.

77. The antibody construct of any one of claims 49-62, wherein the first antigen is human MSLN (SEQ ID NO: 2) and the second antigen is human CD40 (SEQ ID NO: 1); wherein:

the VL1 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 230, 231, and 232, respectively;

SEQ ID NOs: 236, 237, and 238, respectively;

SEQ ID NOs: 242, 243, and 244, respectively; and

SEQ ID NOs: 248, 249, and 250, respectively;

the VH1 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting

of:

SEQ ID NOs: 254, 255, and 256, respectively;

SEQ ID NOs: 260, 261, and 262, respectively;

SEQ ID NOs: 266, 267, and 268, respectively; and

SEQ ID NOs: 272, 273, and 274, respectively;

the VL2 comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 58, 59, and 60, respectively;

SEQ ID NOs: 64, 65, and 66, respectively;

SEQ ID NOs: 70, 71, and 72, respectively;

SEQ ID NOs: 76, 77, and 78, respectively;

SEQ ID NOs: 82, 83, and 84, respectively;

SEQ ID NOs: 88, 89, and 90, respectively;

SEQ ID NOs: 94, 95, and 96, respectively;

SEQ ID NOs: 100, 101, and 102, respectively;

SEQ ID NOs: 106, 107, and 108, respectively;

SEQ ID NOs: 112, 113, and 114, respectively;

SEQ ID NOs: 118, 119, and 120, respectively;

SEQ ID NOs: 124, 125, and 126, respectively; and

SEQ ID NOs: 130, 131, and 132, respectively;

and

the VH2 comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting

of:

SEQ ID NOs: 136, 137, and 138, respectively;

SEQ ID NOs: 142, 143, and 144, respectively;

SEQ ID NOs: 148, 149, and 150, respectively;

SEQ ID NOs: 154, 155, and 156, respectively;
SEQ ID NOs: 160, 161, and 162, respectively;
SEQ ID NOs: 166, 167, and 168, respectively;
SEQ ID NOs: 172, 173, and 174, respectively;
SEQ ID NOs: 178, 179, and 180, respectively;
SEQ ID NOs: 184, 185, and 186, respectively;
SEQ ID NOs: 190, 191, and 192, respectively;
SEQ ID NOs: 196, 197, and 198, respectively;
SEQ ID NOs: 202, 203, and 204, respectively; and
SEQ ID NOs: 208, 209, and 210, respectively.

78. The antibody construct of claim 77, wherein

1) the VL1 and VH1 are selected from the group consisting of:

a) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 230, 231, and 232, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 254, 255, and 256, respectively;

b) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 236, 237, and 238, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 260, 261, and 262, respectively;

c) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 242, 243, and 244, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 266, 267, and 268, respectively; and

d) a VL1 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 248, 249, and 250, respectively; and a VH1 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 272, 273, and 274, respectively;

and

2) the VL2 and VH2 are selected from the group consisting of:

a) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

b) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH2 comprising a CDRH1, a

CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

c) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

d) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

e) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

f) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

g) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

h) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

i) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

j) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

k) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH2 comprising a

CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

l) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

m) a VL2 comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH2 comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

79. The antibody construct of claim 77, wherein:

the VL1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 213, 217, 221, and 225;

the VH1 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 214, 218, 222, and 226;

the VL2 comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

the VH2 comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

80. The antibody construct of claim 79, wherein:

1) the VL1 and VH1 are selected from the group consisting of:

- a) a VL1 comprising SEQ ID NO: 213 and a VH1 comprising SEQ ID NO: 214;
 - b) a VL1 comprising SEQ ID NO: 217 and a VH1 comprising SEQ ID NO: 218;
 - c) a VL1 comprising SEQ ID NO: 221 and a VH1 comprising SEQ ID NO: 222; and
 - d) a VL1 comprising SEQ ID NO: 225 and a VH1 comprising SEQ ID NO: 226;
- and

2) the VL2 and VH2 are selected from the group consisting of:

- a) a VL2 comprising SEQ ID NO: 5 and a VH2 comprising SEQ ID NO: 6;
- b) a VL2 comprising SEQ ID NO: 9 and a VH2 comprising SEQ ID NO: 10;
- c) a VL2 comprising SEQ ID NO: 13 and a VH2 comprising SEQ ID NO: 14;
- d) a VL2 comprising SEQ ID NO: 17 and a VH2 comprising SEQ ID NO: 18;
- e) a VL2 comprising SEQ ID NO: 21 and a VH2 comprising SEQ ID NO: 22;
- f) a VL2 comprising SEQ ID NO: 25 and a VH2 comprising SEQ ID NO: 26;
- g) a VL2 comprising SEQ ID NO: 29 and a VH2 comprising SEQ ID NO: 30;
- h) a VL2 comprising SEQ ID NO: 33 and a VH2 comprising SEQ ID NO: 34;

- i) a VL2 comprising SEQ ID NO: 37 and a VH2 comprising SEQ ID NO: 38;
- j) a VL2 comprising SEQ ID NO: 41 and a VH2 comprising SEQ ID NO: 42;
- k) a VL2 comprising SEQ ID NO: 45 and a VH2 comprising SEQ ID NO: 46;
- l) a VL2 comprising SEQ ID NO: 49 and a VH2 comprising SEQ ID NO: 50; and
- m) a VL2 comprising SEQ ID NO: 53 and a VH2 comprising SEQ ID NO: 54.

81. An antigen binding protein that specifically binds and agonizes human CD40 (SEQ ID NO: 1), the antigen binding protein comprising a light chain variable region (VL) and a heavy chain variable region (VH), wherein:

the VL comprises a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of:

SEQ ID NOs: 58, 59, and 60, respectively;
SEQ ID NOs: 64, 65, and 66, respectively;
SEQ ID NOs: 70, 71, and 72, respectively;
SEQ ID NOs: 76, 77, and 78, respectively;
SEQ ID NOs: 82, 83, and 84, respectively;
SEQ ID NOs: 88, 89, and 90, respectively;
SEQ ID NOs: 94, 95, and 96, respectively;
SEQ ID NOs: 100, 101, and 102, respectively;
SEQ ID NOs: 106, 107, and 108, respectively;
SEQ ID NOs: 112, 113, and 114, respectively;
SEQ ID NOs: 118, 119, and 120, respectively;
SEQ ID NOs: 124, 125, and 126, respectively; and
SEQ ID NOs: 130, 131, and 132, respectively;

and

the VH comprises a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of:

SEQ ID NOs: 136, 137, and 138, respectively;
SEQ ID NOs: 142, 143, and 144, respectively;
SEQ ID NOs: 148, 149, and 150, respectively;
SEQ ID NOs: 154, 155, and 156, respectively;
SEQ ID NOs: 160, 161, and 162, respectively;
SEQ ID NOs: 166, 167, and 168, respectively;
SEQ ID NOs: 172, 173, and 174, respectively;
SEQ ID NOs: 178, 179, and 180, respectively;
SEQ ID NOs: 184, 185, and 186, respectively;
SEQ ID NOs: 190, 191, and 192, respectively;
SEQ ID NOs: 196, 197, and 198, respectively;

SEQ ID NOs: 202, 203, and 204, respectively; and

SEQ ID NOs: 208, 209, and 210, respectively.

82. The antibody construct of claim 81, wherein

the VL and VH are selected from the group consisting of:

a) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 58, 59, and 60, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 136, 137, and 138, respectively;

b) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 64, 65, and 66, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 142, 143, and 144, respectively;

c) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 70, 71, and 72, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 148, 149, and 150, respectively;

d) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 76, 77, and 6780, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 154, 155, and 156, respectively;

e) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 82, 83, and 84, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 160, 161, and 162, respectively;

f) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 88, 89, and 90, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 166, 167, and 168, respectively;

g) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 94, 95, and 96, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 172, 173, and 174, respectively;

h) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 100, 101, and 102, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 178, 179, and 180, respectively;

i) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 106, 107, and 108, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 184, 185, and 186, respectively;

j) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 112, 113, and 114, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 190, 191, and 192, respectively;

k) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 118, 119, and 120, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 196, 197, and 198, respectively;

l) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 124, 125, and 126, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 202, 203, and 204, respectively; and

m) a VL comprising a CDRL1, a CDRL2, and a CDRL3 selected from the group consisting of SEQ ID NOs: 130, 131, and 132, respectively; and a VH comprising a CDRH1, a CDRH2, and a CDRH3 selected from the group consisting of SEQ ID NOs: 208, 209, and 210, respectively.

83. The antibody construct of claim 81, wherein:

the VL comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 5, 9, 13, 17, 21, 25, 29, 33, 37, 41, 45, 49, and 53; and

the VH comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 6, 10, 14, 18, 22, 26, 30, 34, 38, 42, 46, 50, and 54.

84. The antibody construct of claim 83, wherein:

the VL2 and VH2 are selected from the group consisting of:

- a) a VL comprising SEQ ID NO: 5 and a VH comprising SEQ ID NO: 6;
- b) a VL comprising SEQ ID NO: 9 and a VH comprising SEQ ID NO: 10;
- c) a VL comprising SEQ ID NO: 13 and a VH comprising SEQ ID NO: 14;
- d) a VL comprising SEQ ID NO: 17 and a VH comprising SEQ ID NO: 18;
- e) a VL comprising SEQ ID NO: 21 and a VH comprising SEQ ID NO: 22;
- f) a VL comprising SEQ ID NO: 25 and a VH comprising SEQ ID NO: 26;
- g) a VL comprising SEQ ID NO: 29 and a VH comprising SEQ ID NO: 30;
- h) a VL comprising SEQ ID NO: 33 and a VH comprising SEQ ID NO: 34;

- i) a VL comprising SEQ ID NO: 37 and a VH comprising SEQ ID NO: 38;
- j) a VL comprising SEQ ID NO: 41 and a VH comprising SEQ ID NO: 42;
- k) a VL comprising SEQ ID NO: 45 and a VH comprising SEQ ID NO: 46;
- l) a VL comprising SEQ ID NO: 49 and a VH comprising SEQ ID NO: 50; and
- m) a VL comprising SEQ ID NO: 53 and a VH comprising SEQ ID NO: 54.

85. The antibody construct according to any one of claims 81-84, wherein the heavy chain comprises an amino acid substitution selected from the group consisting of:

- (i) N297G or N297A;
- (ii) L234A and L235A; and
- (iii) R292C and V302C;

wherein the amino acid numbering is EU numbering according to Kabat.

86. The antibody construct according to claim 85, wherein the heavy chain comprises N297G, R292C, and V302C mutations, wherein the amino acid numbering is EU numbering according to Kabat.

87. The antibody construct according to any one of claims 81-86, wherein the antigen binding protein further comprises a light chain CL polypeptide linked to the VL, wherein the CL polypeptide is selected from the group consisting of SEQ ID NO: 883 and SEQ ID NO: 884.

88. The antibody construct according to any one of claims 81-87, wherein the antigen binding protein further comprises a heavy chain CH1-hinge-CH2-CH3 polypeptide wherein the CH1-hinge-CH2-CH3 polypeptide is selected from the group consisting of SEQ ID NO: 885 and SEQ ID NO: 886.

89. The antibody construct according to any one of claims 81-88, wherein the antigen binding protein further comprises a second antigen binding portion that specifically binds a second antigen.

90. The antibody construct according to claim 89, wherein the second antigen is a tumor associated antigen (TAA).