

ANTI-PD-1 ANTIBODIES, ACTIVATABLE ANTI-PD-1 ANTIBODIES, AND METHODS OF USE THEREOF

Related Applications

[0001] This application claims the benefit of U.S. Provisional Application No. 62/191,902, filed July 13, 2015; U.S. Provisional Application No. 62/205,825, filed August 17, 2015; U.S. Provisional Application No. 62/295,314, filed February 15, 2016; U.S. Provisional Application No. 62/323,543, filed April 15, 2016 and U.S. Provisional Application No. 62/333,629, filed May 9, 2016; the contents of each of which are incorporated herein by reference in their entirety.

Field of the Invention

[0002] The invention relates generally to antibodies that specifically bind programmed cell death protein 1 (PD-1), activatable antibodies that specifically bind to PD-1 and methods of making and using these anti-PD-1 antibodies and anti-PD-1 activatable antibodies in a variety of therapeutic, diagnostic and prophylactic indications.

Background of the Invention

[0003] Antibody-based therapies have proven effective treatments for several diseases but in some cases, toxicities due to broad target expression have limited their therapeutic effectiveness. In addition, antibody-based therapeutics have exhibited other limitations such as rapid clearance from the circulation following administration.

[0004] Under conditions of chronic stimulation, T cells upregulate and sustain expression of the inhibitory receptor PD-1 to negatively regulate the quality and magnitude of T cell responses. The primary ligand for PD-1, PD-L1 is upregulated on many tumor cells and has been associated with inhibition of anti-tumor T-cell immunity via its engagement of PD-1 on tumor-infiltrating T cells. Clinical trials have confirmed the capacity of antibody blockade of either PD-1 or PD-L1 to restore the activity of durable tumor-specific immunity in patients across multiple tumor types. (Herbst et al, 2014; Lipson et al, 2015). However, because similar mechanisms control anti-tumor immunity and self-tolerance, systemic delivery of these checkpoint-targeted therapies can also induce systemic autoimmunity that can be exacerbated with combination treatments; such as nivolumab or pembrolizumab (anti-PD-1) and ipilimumab (anti-CTLA4).

New approaches are therefore needed that provide anti-tumor activity without deregulating systemic immunity.

[0005] In the realm of small molecule therapeutics, strategies have been developed to provide prodrugs of an active chemical entity. Such prodrugs are administered in a relatively inactive (or significantly less active) form. Once administered, the prodrug is metabolized *in vivo* into the active compound. Such prodrug strategies can provide for increased selectivity of the drug for its intended target and for a reduction of adverse effects.

[0006] Accordingly, there is a continued need in the field of antibody-based therapeutics for antibodies that mimic the desirable characteristics of the small molecule prodrug.

Summary of the Invention

[0007] The disclosure provides antibodies or antigen-binding fragments thereof that specifically bind programmed cell death protein 1 (PD-1), also known as CD279, SLEB2, and/or hSLE1. The use of the term “PD-1” is intended to cover any variation thereof, such as, by way of non-limiting example, PD1 and/or PD 1, all variations are used herein interchangeably. Aberrant expression and/or activity of PD-1 and PD-1-related signaling has been implicated in the pathogenesis of many diseases and disorders, such as cancer.

[0008] The present invention provides monoclonal antibodies (mAbs), activatable antibodies, and antigen-binding fragments thereof that specifically bind PD-1.

[0009] In some embodiments, the antibody includes an antibody or antigen-binding fragment thereof that specifically binds PD-1. In some embodiments, the antibody or antigen-binding fragment thereof that binds PD-1 is a monoclonal antibody, domain antibody, single chain, Fab fragment, a F(ab')₂ fragment, a scFv, a scAb, a dAb, a single domain heavy chain antibody, or a single domain light chain antibody. In some embodiments, such an antibody or antigen-binding fragment thereof that binds PD-1 is a mouse, other rodent, chimeric, humanized or fully human monoclonal antibody.

[00010] In some embodiments, the antibody includes an isolated antibody or antigen binding fragment thereof (AB) that specifically binds to mammalian PD-1, wherein the AB has one or more of the characteristics selected from the group consisting of: (a) the AB inhibits binding of mammalian PD-1 to mammalian PDL1 with an EC₅₀ value less than 5 nM; (b) the AB inhibits binding of mammalian PD-1 to mammalian PDL2 with an EC₅₀ value less than 5 nM; and (c) the AB specifically binds to human PD-1 and cynomolgus monkey PD-1.

[00011] In some embodiments, the antibody specifically binds to the mammalian PD-1 with a dissociation constant of 0.01 nM to 5 nM, 0.05 nM to 5 nM, 0.1 nM to 5 nM, 0.2 nM to 5 nM, 0.3 nM to 5 nM, 0.4 nM to 5 nM, 0.5 nM to 5 nM, 0.75 nM to 5 nM, 1 nM to 5 nM, 2 nM to 5 nM, 0.01 nM to 2 nM, 0.05 nM to 2 nM, 0.1 nM to 2 nM, 0.2 nM to 2 nM, 0.3 nM to 2 nM, 0.4 nM to 2 nM, 0.5 nM to 2 nM, 0.75 nM to 1 nM, 1 nM to 2 nM, 0.01 nM to 1 nM, 0.05 nM to 1 nM, 0.1 nM to 1 nM, 0.2 nM to 1 nM, 0.3 nM to 1 nM, 0.4 nM to 1 nM, 0.5 nM to 1 nM, 0.75 nM to 1 nM, 0.01 nM to 0.75 nM, 0.05 nM to 0.75 nM, 0.1 nM to 0.75 nM, 0.2 nM to 0.75 nM, 0.3 nM to 0.75 nM, 0.4 nM to 0.75 nM, 0.5 nM to 0.75 nM, 0.01 nM to 0.5 nM, 0.05 nM to 0.5 nM, 0.1 nM to 0.5 nM, 0.2 nM to 0.5 nM, 0.3 nM to 0.5 nM, 0.4 nM to 0.5 nM, 0.01 nM to 0.4 nM, 0.05 nM to 0.4 nM, 0.1 nM to 0.4 nM, 0.2 nM to 0.4 nM, 0.3 nM to 0.4 nM, 0.01 nM to 0.3 nM, 0.05 nM to 0.3 nM, 0.1 nM to 0.3 nM, 0.2 nM to 0.3 nM, 0.01 nM to 0.2 nM, 0.05 nM to 0.2 nM, 0.1 nM to 0.2 nM, 0.01 nM to 0.1 nM, 0.05 nM to 0.1 nM, or 0.01 nM to 0.05 nM.

[00012] In some embodiments, the mammalian PD-1 is selected from the group consisting of a human PD-1 and a cynomolgus monkey PD-1. In some embodiments, the mammalian PD-1 is a murine PD-1. In some embodiments, the antibody specifically binds to human PD-1 or cynomolgus monkey PD-1 with a dissociation constant of less than or equal to 1 nM. In some embodiments, the mammalian PD-1 is a human PD-1.

[00013] In some embodiments, the antibody or antigen binding fragment thereof specifically binds to the mammalian PD-1 with a dissociation constant is less than or equal to 0.01 nM, less than or equal to 0.05 nM, less than or equal to 0.1 nM, less than or equal to 0.2 nM, less than or equal to 0.3 nM, less than or equal to 0.4 nM, less than or equal to 0.5 nM, less than or equal to 0.75 nM, and less than or equal to 1 nM.

[00014] In some embodiments, the antibody has one or more of the characteristics selected from the group consisting of: (a) the AB specifically binds human PD-1 and cynomolgus monkey PD-1; (b) the AB inhibits binding of human PDL1 and human PDL2 to human PD-1; (c) the AB inhibits binding of cynomolgus monkey PDL1 and cynomolgus monkey PDL2 to cynomolgus monkey PD-1; (d) the AB specifically binds to murine PD-1; and (e) the AB inhibits binding of murine PDL1 and murine PDL2 to murine PD-1.

[00015] In some embodiments, the antibody blocks the ability of a natural ligand to bind to the mammalian PDL1 with an EC_{50} of 0.1 nM to 10 nM, 0.1 nM to 5 nM, 0.1 nM to 3 nM,

0.1 nM to 2 nM, 0.1 nM to 1 nM, 0.1 nM to 0.5 nM, 0.1 nM to 0.25 nM, 0.25 nM to 10 nM, 0.25 nM to 5 nM, 0.25 nM to 3 nM, 0.25 nM to 2 nM, 0.25 nM to 1 nM, 0.25 nM to 0.5 nM, 0.5 nM to 10 nM, 0.5 nM to 5 nM, 0.5 nM to 3 nM, 0.5 nM to 2 nM, 0.5 nM to 1 nM, 1 nM to 10 nM, 1 nM to 5 nM, 1 nM to 3 nM, 1 nM to 2 nM, 2 nM to 10 nM, 2 nM to 5 nM, 2 nM to 3 nM, 3 nM to 10 nM, 3 nM to 5 nM, or 5 nM to 10 nM. In some embodiments, the natural ligand is a mammalian PDL1 or a mammalian PDL2. In some embodiments, the natural ligand is selected from the group consisting of: a human PDL1, a human PDL2, a cynomolgus monkey PDL1, and a cynomolgus monkey PDL2. In some embodiments, the natural ligand is a murine PDL1 or a murine PDL2.

[00016] In some embodiments, the antibody blocks the ability of a natural ligand to bind to the mammalian PDL1 with an EC_{50} of less than or equal to 0.1 nM, less than or equal to 0.25 nM, less than or equal to 0.5 nM, less than or equal to 1 nM, less than or equal to 2 nM, less than or equal to 3 nM, less than or equal to 4 nM, less than or equal to 5 nM or less than or equal to 10 nM.

[00017] In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00018] In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00019] In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21. In some embodiments, the anti-PD-1 antibody includes a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is

derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45. In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 47.

[00020] In some embodiments, the anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 21. In some embodiments, the anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or of SEQ ID NO: 47. In some embodiments, the anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or 47. In some embodiments, the anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45. In some embodiments, the anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 47.

[00021] In some embodiments, the anti-PD-1 antibody includes: (a) a variable heavy chain complementarity determining region 1 (VH CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 653-657; (b) a variable heavy chain complementarity determining region 2 (VH CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 658-663; (c) a variable heavy chain complementarity determining region 3 (VH CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 664-668; (d) a variable light chain complementarity determining region 1 (VL CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 669-677; (e) a variable light chain complementarity determining region 2 (VL CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 678-682; and (f) variable light chain complementarity determining region 3 (VL CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 683-687.

[00022] In some embodiments, the anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein

as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664).

[00023] In some embodiments, the anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[00024] In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence,

a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[00025] In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1346, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 626.

[00026] In some embodiments, the anti-PD-1 antibody includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1514, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 638.

[00027] In some embodiments, the anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346. In some embodiments, the anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626. In some embodiments, the anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626.

[00028] In some embodiments, the anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity

determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707).

[00029] In some embodiments, the anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[00030] In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[00031] In some embodiments, the anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514. In some embodiments, the anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638. In some embodiments, the anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638.

[00032] In some embodiments, the anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity

determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713).

[00033] In some embodiments, the anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[00034] In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[00035] In some embodiments, the antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the antibody comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence

selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00036] In some embodiments, the antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the antibody comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 45. In some embodiments, the anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 47.

[00037] In some embodiments, the anti-PD-1 antibody includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1346, and a light chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 626.

[00038] In some embodiments, the anti-PD-1 antibody includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1514, and a light chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 638.

[00039] In some embodiments, the antibody comprises a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence shown in Table 8.

[00040] In some embodiments, the antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence that

includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR3 sequence shown in Table 8.

[00041] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the combination is a combination of the three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[00042] In some embodiments, the antibody comprises a light chain that comprises a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination is a combination of the three light chain CDR sequences (VL CDR1, VL CDR2, VL CDR3) shown in a single row in Table 8.

[00043] In some embodiments, the antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, and a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8.

[00044] In some embodiments, the antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, and a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding VH CDR sequence shown in a single row in Table 8 and the corresponding VL CDR shown in a single row in Table 8.

[00045] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in a combination of three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[00046] In some embodiments, the antibody comprises a light chain that comprises a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in

a combination of three light chain CDR sequences (VL CDR1, VL CDR2, VL CDR3) shown in a single row in Table 8.

[00047] In some embodiments, the antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707).

[00048] In some embodiments, the anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[00049] In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence

comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[00050] In some embodiments, the antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713).

[00051] In some embodiments, the anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[00052] In some embodiments, the anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID

NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[00053] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[00054] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00055] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00056] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00057] In some embodiments, the anti-PD-1 is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1346, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 626.

[00058] In some embodiments, the anti-PD-1 antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1514, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 638.

[00059] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[00060] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00061] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00062] In some embodiments, the antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least

90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00063] In some embodiments, the anti-PD-1 is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1346, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain t that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 626.

[00064] In some embodiments, the anti-PD-1 antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1514, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 638.

[00065] In some embodiments, the antibody is encoded by a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38. In some embodiments, the antibody is encoded by a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38.

[00066] In some embodiments, the antibody is encoded by a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the antibody is encoded by a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[00067] In some embodiments, the antibody is encoded by heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a light chain nucleic

acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the antibody is encoded by heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[00068] In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38. In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38.

[00069] In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[00070] In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%,

98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[00071] In some embodiments, the antibody or antigen-binding fragment thereof is incorporated in a multispecific antibody or antigen-binding fragment thereof, where at least one arm of the multispecific antibody specifically binds PD-1. In some embodiments, the antibody or antigen-binding fragment thereof is incorporated in a bispecific antibody or antigen-binding fragment thereof, where at least one arm of the bispecific antibody specifically binds PD-1.

[00072] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[00073] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00074] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain amino acid sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain amino acid sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00075] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from amino acid

sequence SEQ ID NO: 21. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 47.

[00076] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 21. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or of SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or 47. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 47.

[00077] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody includes: (a) a variable heavy chain complementarity determining region 1 (VH CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 653-657; (b) a variable heavy chain complementarity determining region 2 (VH CDR2)

comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 658-663; (c) a variable heavy chain complementarity determining region 3 (VH CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 664-668; (d) a variable light chain complementarity determining region 1 (VL CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 669-677; (e) a variable light chain complementarity determining region 2 (VL CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 678-682; and (f) variable light chain complementarity determining region 3 (VL CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 683-687

[00078] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664).

[00079] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific

antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO:676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[00080] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO:676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO:676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[00081] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from the amino acid

sequence of SEQ ID NO: 1346, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 626.

[00082] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1514, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 638.

[00083] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626.

[00084] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707).

[00085] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[00086] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[00087] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638.

[00088] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713).

[00089] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises

KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[00090] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[00091] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[00092] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00093] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or

99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[00094] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1346, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 626.

[00095] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1514, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 638.

[00096] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707).

[00097] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[00098] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence,

wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[00099] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713).

[000100] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[000101] In some embodiments, at least one arm of the multispecific antibody, *e.g.*, a bispecific antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence,

wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[000102] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence shown in Table 8.

[000103] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR1 sequence shown in Table 8; a VH CD2 sequence

that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR3 sequence shown in Table 8.

[000104] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination is a combination of the six CDR sequences (VH CDR1, VH CDR2, VH CDR3, VL CDR1, VL CDR2, and VL CDR3) shown in a single row in Table 8.

[000105] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the combination is a combination of the three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000106] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding VH CDR sequence shown in a single row in Table 8 and the corresponding VL CDR shown in a single row in Table 8.

[000107] In some embodiments, at least one arm of the multispecific antibody, e.g., a bispecific antibody, comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%,

98%, 99% or more identical to the corresponding CDR sequence in a combination of three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000108] The disclosure also provides activatable antibodies that include an antibody or antigen-binding fragment thereof that specifically binds PD-1 coupled to a masking moiety (MM), such that coupling of the MM reduces the ability of the antibody or antigen-binding fragment thereof to bind PD-1. In some embodiments, the MM is coupled via a cleavable moiety (CM) that includes sequence that functions as a substrate for a protease. The activatable anti-PD-1 antibodies of the disclosure are activated when the cleavable moiety is cleaved by a protease. For example, the protease is produced by a tumor that is in proximity to T cells that express PD-1. In some embodiments, the protease is produced by a tumor that is co-localized with T cells that express PD-1.

[000109] The activatable anti-PD-1 antibodies provided herein, also referred to herein as anti-PD-1 activatable antibodies or PD-1 activatable antibodies, are stable in circulation, activated at intended sites of therapy and/or diagnosis but not in normal, *e.g.*, healthy tissue or other tissue not targeted for treatment and/or diagnosis, and, when activated, exhibit binding to PD-1 that is at least comparable to the corresponding, unmodified antibody.

[000110] The invention also provides methods of treating, preventing and/or delaying the onset or progression of, or alleviating a symptom associated with aberrant expression and/or activity of PD-1 in a subject using antibodies or activatable antibodies that bind PD-1, particularly activatable antibodies that bind and neutralize or otherwise inhibit at least one biological activity of PD-1.

[000111] In some embodiments, the activatable anti-PD-1 antibody comprises an activatable antibody that, in an activated state, specifically binds to mammalian PD-1, wherein said activatable antibody comprises: an antibody or an antigen binding fragment thereof (AB) that specifically binds to mammalian PD-1; a masking moiety (MM) that inhibits the binding of the AB to mammalian PD-1 when the activatable antibody is in an uncleaved state; and a cleavable moiety (CM) coupled to the AB, wherein the CM is a polypeptide that functions as a substrate for a protease.

[000112] In some embodiments, the activatable anti-PD-1 antibody comprises an activatable antibody that, in an activated state, (a) specifically binds to mammalian PD-1; and (b) specifically blocks a natural ligand of PD-1 from binding to the mammalian PD-1, wherein the

activatable antibody comprises: an antibody or an antigen binding fragment thereof (AB) that specifically binds to mammalian PD-1; a masking moiety (MM) that inhibits the binding of the AB to mammalian PD-1 when the activatable antibody is in an uncleaved state; and a cleavable moiety (CM) coupled to the AB, wherein the CM is a polypeptide that functions as a substrate for a protease.

[000113] In some embodiments, the activatable antibody in an uncleaved state specifically binds to the mammalian PD-1 with a dissociation constant of 0.5 nM to 1 nM, 0.5 nM to 2 nM, 0.5 nM to 5 nM, 0.5 nM to 10 nM, 0.5 nM to 15 nM, 0.5 nM to 20 nM, 0.5 nM to 25 nM, 0.5 nM to 50 nM, 0.5 nM to 75 nM, 0.5 nM to 100 nM, 0.5 nM to 150 nM, 0.5 nM to 200 nM, 0.5 nM to 300 nM, 0.5 nM to 400 nM, 1 nM to 2 nM, 1 nM to 5 nM, 1 nM to 10 nM, 1 nM to 15 nM, 1 nM to 20 nM, 1 nM to 25 nM, 1 nM to 50 nM, 1 nM to 75 nM, 1 nM to 100 nM, 1 nM to 150 nM, 1 nM to 200 nM, 1 nM to 300 nM, 1 nM to 400 nM, 2 nM to 5 nM, 2 nM to 10 nM, 2 nM to 15 nM, 2 nM to 20 nM, 2 nM to 25 nM, 2 nM to 50 nM, 2 nM to 75 nM, 2 nM to 100 nM, 2 nM to 150 nM, 2 nM to 200 nM, 2 nM to 300 nM, 2 nM to 400 nM, 5 nM to 10 nM, 5 nM to 15 nM, 5 nM to 20 nM, 5 nM to 25 nM, 5 nM to 50 nM, 5 nM to 75 nM, 5 nM to 100 nM, 5 nM to 150 nM, 5 nM to 200 nM, 5 nM to 300 nM, 5 nM to 400 nM, 10 nM to 15 nM, 10 nM to 20 nM, 10 nM to 25 nM, 10 nM to 50 nM, 10 nM to 75 nM, 10 nM to 100 nM, 10 nM to 150 nM, 10 nM to 200 nM, 10 nM to 300 nM, 10 nM to 400 nM, 15 nM to 20 nM, 15 nM to 25 nM, 15 nM to 50 nM, 15 nM to 75 nM, 15 nM to 100 nM, 15 nM to 150 nM, 15 nM to 200 nM, 15 nM to 300 nM, 15 nM to 400 nM, 20 nM to 25 nM, 20 nM to 50 nM, 20 nM to 75 nM, 20 nM to 100 nM, 20 nM to 150 nM, 20 nM to 200 nM, 20 nM to 300 nM, 20 nM to 400 nM, 25 nM to 50 nM, 25 nM to 75 nM, 25 nM to 100 nM, 25 nM to 150 nM, 25 nM to 200 nM, 25 nM to 300 nM, 25 nM to 400 nM, 50 nM to 75 nM, 50 nM to 100 nM, 50 nM to 150 nM, 50 nM to 200 nM, 50 nM to 300 nM, 50 nM to 400 nM, 75 nM to 100 nM, 75 nM to 150 nM, 75 nM to 200 nM, 75 nM to 300 nM, 75 nM to 400 nM, 100 nM to 150 nM, 100 nM to 200 nM, 100 nM to 300 nM, 100 nM to 400 nM, 150 nM to 200 nM, 150 nM to 300 nM, 150 nM to 400 nM, 200 nM to 300 nM, 200 nM to 400 nM, or 300 nM to 400 nM.

[000114] In some embodiments, the activatable antibody in an activated state specifically binds to the mammalian PD-1 with a dissociation constant of 0.01 nM to 5 nM, 0.05 nM to 5 nM, 0.1 nM to 5 nM, 0.2 nM to 5 nM, 0.3 nM to 5 nM, 0.4 nM to 5 nM, 0.5 nM to 5 nM, 0.75 nM to 5 nM, 1 nM to 5 nM, 2 nM to 5 nM, 0.01 nM to 2 nM, 0.05 nM to 2 nM, 0.1 nM to

2 nM, 0.2 nM to 2 nM, 0.3 nM to 2 nM, 0.4 nM to 2 nM, 0.5 nM to 2 nM, 0.75 nM to 1 nM, 1 nM to 2 nM, 0.01 nM to 1 nM, 0.05 nM to 1 nM, 0.1 nM to 1 nM, 0.2 nM to 1 nM, 0.3 nM to 1 nM, 0.4 nM to 1 nM, 0.5 nM to 1 nM, 0.75 nM to 1 nM, 0.01 nM to 0.75 nM, 0.05 nM to 0.75 nM, 0.1 nM to 0.75 nM, 0.2 nM to 0.75 nM, 0.3 nM to 0.75 nM, 0.4 nM to 0.75 nM, 0.5 nM to 0.75 nM, 0.01 nM to 0.5 nM, 0.05 nM to 0.5 nM, 0.1 nM to 0.5 nM, 0.2 nM to 0.5 nM, 0.3 nM to 0.5 nM, 0.4 nM to 0.5 nM, 0.01 nM to 0.4 nM, 0.05 nM to 0.4 nM, 0.1 nM to 0.4 nM, 0.2 nM to 0.4 nM, 0.3 nM to 0.4 nM, 0.01 nM to 0.3 nM, 0.05 nM to 0.3 nM, 0.1 nM to 0.3 nM, 0.2 nM to 0.3 nM, 0.01 nM to 0.2 nM, 0.05 nM to 0.2 nM, 0.1 nM to 0.2 nM, 0.01 nM to 0.1 nM, 0.05 nM to 0.1 nM, or 0.01 nM to 0.05 nM.

[000115] In some embodiments, the activatable antibody comprises an AB that specifically binds to the mammalian PD-1 with a dissociation constant of 0.01 nM to 5 nM, 0.05 nM to 5 nM, 0.1 nM to 5 nM, 0.2 nM to 5 nM, 0.3 nM to 5 nM, 0.4 nM to 5 nM, 0.5 nM to 5 nM, 0.75 nM to 5 nM, 1 nM to 5 nM, 2 nM to 5 nM, 0.01 nM to 2 nM, 0.05 nM to 2 nM, 0.1 nM to 2 nM, 0.2 nM to 2 nM, 0.3 nM to 2 nM, 0.4 nM to 2 nM, 0.5 nM to 2 nM, 0.75 nM to 1 nM, 1 nM to 2 nM, 0.01 nM to 1 nM, 0.05 nM to 1 nM, 0.1 nM to 1 nM, 0.2 nM to 1 nM, 0.3 nM to 1 nM, 0.4 nM to 1 nM, 0.5 nM to 1 nM, 0.75 nM to 1 nM, 0.01 nM to 0.75 nM, 0.05 nM to 0.75 nM, 0.1 nM to 0.75 nM, 0.2 nM to 0.75 nM, 0.3 nM to 0.75 nM, 0.4 nM to 0.75 nM, 0.5 nM to 0.75 nM, 0.01 nM to 0.5 nM, 0.05 nM to 0.5 nM, 0.1 nM to 0.5 nM, 0.2 nM to 0.5 nM, 0.3 nM to 0.5 nM, 0.4 nM to 0.5 nM, 0.01 nM to 0.4 nM, 0.05 nM to 0.4 nM, 0.1 nM to 0.4 nM, 0.2 nM to 0.4 nM, 0.3 nM to 0.4 nM, 0.01 nM to 0.3 nM, 0.05 nM to 0.3 nM, 0.1 nM to 0.3 nM, 0.2 nM to 0.3 nM, 0.01 nM to 0.2 nM, 0.05 nM to 0.2 nM, 0.1 nM to 0.2 nM, 0.01 nM to 0.1 nM, 0.05 nM to 0.1 nM, or 0.01 nM to 0.05 nM.

[000116] In some embodiments, the mammalian PD-1 is selected from the group consisting of a human PD-1 and a cynomolgus monkey PD-1. In some embodiments, the AB specifically binds to human PD-1 or cynomolgus monkey PD-1 with a dissociation constant of less than or equal to 1 nM. In some embodiments, the mammalian PD-1 is a human PD-1. In some embodiments, the AB has one or more of the characteristics selected from the group consisting of: (a) the AB specifically binds human PD-1 and cynomolgus monkey PD-1; (b) the AB inhibits binding of human PDL1 and human PDL2 to human PD-1; and (c) the AB inhibits binding of cynomolgus monkey PDL1 and cynomolgus monkey PDL2 to cynomolgus monkey PD-1.

[000117] In some embodiments, the mammalian PD-1 is mouse PD-1. In some embodiments, the activatable antibody comprises an AB that specifically binds mouse PD-1 or inhibits binding of mouse PDL1 and mouse PDL2 to mouse PD1.

[000118]

[000119] I

[000120] In some embodiments, the activatable antibody in an uncleaved state specifically binds to the mammalian PD-1 with a dissociation constant greater than or equal to 0.5 nM, greater than or equal to 1 nM, greater than or equal to 2 nM, greater than or equal to 3 nM, greater than or equal to 4 nM, greater than or equal to 5 nM, greater than or equal to 10 nM, greater than or equal to 15 nM, greater than or equal to 20 nM, greater than or equal to 25 nM, greater than or equal to 50 nM, greater than or equal to 75 nM, greater than or equal to 100 nM, greater than or equal to 150 nM, greater than or equal to 200 nM, greater than or equal to 300 nM and/or greater than or equal to 400 nM.

[000121] In some embodiments, the activatable antibody in an activated state specifically binds to the mammalian PD-1 with a dissociation constant less than or equal to 0.01 nM, less than or equal to 0.05 nM, less than or equal to 0.1 nM, less than or equal to 0.2 nM, less than or equal to 0.3 nM, less than or equal to 0.4 nM, less than or equal to 0.5 nM, less than or equal to 0.75 nM, and less than or equal to 1 nM.

[000122] In some embodiments, the activatable antibody comprises an AB that specifically binds to the mammalian PD-1 with a dissociation constant less than or equal to 0.01 nM, less than or equal to 0.05 nM, less than or equal to 0.1 nM, less than or equal to 0.2 nM, less than or equal to 0.3 nM, less than or equal to 0.4 nM, less than or equal to 0.5 nM, less than or equal to 0.75 nM, and less than or equal to 1 nM.

[000123] In some embodiments, the activatable antibody comprises an AB blocks the ability of a natural ligand to bind to the mammalian PDL1 with an EC_{50} of 0.1 nM to 10 nM, 0.1 nM to 5 nM, 0.1 nM to 3 nM, 0.1 nM to 2 nM, 0.1 nM to 1 nM, 0.1 nM to 0.5 nM, 0.1 nM to 0.25 nM, 0.25 nM to 10 nM, 0.25 nM to 5 nM, 0.25 nM to 3 nM, 0.25 nM to 2 nM, 0.25 nM to 1 nM, 0.25 nM to 0.5 nM, 0.5 nM to 10 nM, 0.5 nM to 5 nM, 0.5 nM to 3 nM, 0.5 nM to 2 nM, 0.5 nM to 1 nM, 1 nM to 10 nM, 1 nM to 5 nM, 1 nM to 3 nM, 1 nM to 2 nM, 2 nM to 10 nM, 2 nM to 5 nM, 2 nM to 3 nM, 3 nM to 10 nM, 3 nM to 5 nM, or 5 nM to 10 nM.

[000124] In some embodiments, the natural ligand is a mammalian PDL1 or a mammalian PDL2. In some embodiments, the natural ligand is selected from the group consisting of: a human PDL1, a human PDL2, a cynomolgus monkey PDL1, and a cynomolgus monkey PDL2.

[000125] In some embodiments, the activatable antibody has one or more of the following characteristics: (a) the AB induces type 1 diabetes in a non-obese diabetic (NOD) mouse; and (b) the activatable antibody in an uncleaved state inhibits the induction of type 1 diabetes in a NOD mouse.

[000126] In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 0.5 mg/kg to 15 mg/kg, 1 mg/kg to 15 mg/kg, 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, 10 mg/kg to 15 mg/kg, 0.5 mg/kg to 10 mg/kg, 1 mg/kg to 10 mg/kg, 2 mg/kg to 10 mg/kg, 3 mg/kg to 10 mg/kg, 5 mg/kg to 10 mg/kg, 0.5 mg/kg to 5 mg/kg, 1 mg/kg to 5 mg/kg, 2 mg/kg to 5 mg/kg, 3 mg/kg to 5 mg/kg, 0.5 mg/kg to 3 mg/kg, 1 mg/kg to 3 mg/kg, 2 mg/kg to 3 mg/kg, 0.5 mg/kg to 2 mg/kg, 1 mg/kg to 2 mg/kg, or 0.5 mg/kg to 1 mg/kg. In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 3 mg/kg to 10 mg/kg, 3 mg/kg or 10 mg/kg. In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 3 mg/kg to 10 mg/kg. In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 3 mg/kg. In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 10 mg/kg.

[000127]

[000128] In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 0.5 mg/kg to 15 mg/kg, 1 mg/kg to 15 mg/kg, 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, 10 mg/kg to 15 mg/kg, 0.5 mg/kg to 10 mg/kg, 1 mg/kg to 10 mg/kg, 2 mg/kg to 10 mg/kg, 3 mg/kg to 10 mg/kg, 5 mg/kg to 10 mg/kg, 0.5 mg/kg to 5 mg/kg, 1 mg/kg to 5 mg/kg, 2 mg/kg to 5 mg/kg, 3 mg/kg to 5 mg/kg, 0.5 mg/kg to 3 mg/kg, 1 mg/kg to 3 mg/kg, 2 mg/kg to 3 mg/kg, 0.5 mg/kg to 2 mg/kg, 1 mg/kg to 2 mg/kg, or 0.5 mg/kg to 1 mg/kg. In some embodiments, the AB induces

type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 1 mg/kg to 20 mg/kg, 1 mg/kg, 3 mg/kg, 5 mg/kg, 10 mg/kg, or 20 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 1 mg/kg to 20 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 1 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 3 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 5 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 10 mg/kg. In some embodiments, the AB induces type 1 diabetes in the NOD mouse after administration of the AB at a single dose of 20 mg/kg.

[000129]

[000130] In some embodiments, the activatable antibody has one or more of the following characteristics: (a) the activatable antibody in an uncleaved state does not induce type 1 diabetes in greater than 50% of a population of non-obese diabetic (NOD) mice, and (b) the AB induces type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of a population of NOD mice.

[000131] In some embodiments, the activatable antibody does not induce type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of the population of NOD mice after administration to each mouse in the population a single dose of the activatable antibody at a dosage of: 0.5 mg/kg to 15 mg/kg, 1 mg/kg to 15 mg/kg, 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, 10 mg/kg to 15 mg/kg, 0.5 mg/kg to 10 mg/kg, 1 mg/kg to 10 mg/kg, 2 mg/kg to 10 mg/kg, 3 mg/kg to 10 mg/kg, 5 mg/kg to 10 mg/kg, 0.5 mg/kg to 5 mg/kg, 1 mg/kg to 5 mg/kg, 2 mg/kg to 5 mg/kg, 3 mg/kg to 5 mg/kg, 0.5 mg/kg to 3 mg/kg, 1 mg/kg to 3 mg/kg, 2 mg/kg to 3 mg/kg, 0.5 mg/kg to 2 mg/kg, 1 mg/kg to 2 mg/kg, or 0.5 mg/kg to 1 mg/kg. In some embodiments, the activatable antibody inhibits the induction of type 1 diabetes in the NOD mouse after administration of the activatable antibody at a single dose of 3 mg/kg to 10 mg/kg, 3 mg/kg or 10 mg/kg. In some embodiments the dose is a single

dose of 3 mg/kg to 10 mg/kg. In some embodiments the dose is a single dose of 3 mg/kg. In some embodiments the dose is a single dose of 10 mg/kg.

[000132]

[000133] In some embodiments, the activatable antibody has one or more of the following characteristics: (a) the activatable antibody in an uncleaved state does not induce type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of a population of non-obese diabetic (NOD) mice when administered at a single dose of 3 mg/kg; and (b) the AB induces type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of a population of NOD mice, when administered at a single dose of 3 mg/kg.

[000134] In some embodiments, the activatable antibody has one or more of the following characteristics: (a) the activatable antibody in an uncleaved state does not induce type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of a population of non-obese diabetic (NOD) mice when administered at a single dose of 10 mg/kg; and (b) the AB induces type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of a population of NOD mice, when administered at a single dose of 10 mg/kg.

[000135] In some embodiments, the AB induces type 1 diabetes in greater than 50%, greater than 60%, greater than 70%, greater than 80%, between 50% to 100%, between 50% and 75%, or between 70% and 90% of the population of the NOD mice after administration to each mouse in the population a single dose of the AB at a dosage of: 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, 10 mg/kg to 15 mg/kg, 2 mg/kg to 10 mg/kg, 3 mg/kg to 10 mg/kg, 5 mg/kg to 10 mg/kg, 2 mg/kg to 5 mg/kg, 3 mg/kg to 5 mg/kg, or 2 mg/kg to 3 mg/kg. In some embodiments the dose is a single dose of 3 mg/kg to 20 mg/kg, 3 mg/kg, 5 mg/kg, 10 mg/kg, or 20 mg/kg. In some embodiments the dose is a single dose of 3 mg/kg to 20 mg/kg. In some embodiments the dose is a single dose of 3 mg/kg. In some embodiments the dose is a single dose of 5 mg/kg. In some embodiments the dose is a single dose of 10 mg/kg. In some embodiments the dose is a single dose of 20 mg/kg.

[000136]

[000137] In some embodiments, the activatable antibody in an uncleaved state does not induce type 1 diabetes in greater than 50% of a population of non-obese diabetic (NOD) mice, wherein the population of NOD mice are concurrently dosed with an anti-CTLA4 antibody.

[000138] In some embodiments, the population of NOD mice are each administered a single dose of the activatable antibody at a dosage of 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, 10 mg/kg to 15 mg/kg, 2 mg/kg to 10 mg/kg, 3 mg/kg to 10 mg/kg, 5 mg/kg to 10 mg/kg, 2 mg/kg to 5 mg/kg, 3 mg/kg to 5 mg/kg, or 2 mg/kg to 3 mg/kg concurrently with a dose of an anti-CTLA4 antibody. In some embodiments, the population of NOD mice are each administered a single dose of the activatable antibody at a dosage of 10 mg/kg concurrently with a dose of an anti-CTLA4 antibody.

[000139] In some embodiments, the population of NOD mice is each administered a single dose of the anti-CTLA4 antibody at a dosage of 2 mg/kg to 15 mg/kg, 3 mg/kg to 15 mg/kg, 5 mg/kg to 15 mg/kg, or 10 mg/kg to 15 mg/kg concurrently with a dose of an anti-PD-1 activatable antibody. In some embodiments, the population of NOD mice is each administered a single dose of the anti-CTLA4 antibody at a dosage of 10 mg/kg concurrently with a dose of an anti-PD-1 activatable antibody.

[000140] In some embodiments, the population of NOD mice is each administered a single dose of the activatable antibody at a dosage of 5 mg/kg to 15 mg/kg and a single dose of the anti-CTLA4 antibody at a dosage of 5 mg/kg to 15 mg/kg. In some embodiments, the population of NOD mice is each administered a single dose of the activatable antibody at a dosage of 10 mg/kg concurrently with a single dose of an anti-CTLA4 antibody at a dosage of 10 mg/kg.

[000141] In some embodiments, the NOD mouse is a female NOD/ShiLtJ mouse substrain. In some embodiments, the population of NOD mice is each 5 weeks old when the activatable antibody is first administered. In some embodiments, the population of NOD mice is each 10 weeks old when the activatable antibody is first administered.

[000142] The activatable antibodies in an activated state bind PD-1 and include (i) an antibody or an antigen binding fragment thereof (AB) that specifically binds to PD-1; (ii) a masking moiety (MM) that, when the activatable antibody is in an uncleaved state, inhibits the binding of the AB to PD-1; and (c) a cleavable moiety (CM) coupled to the AB, wherein the CM is a polypeptide that functions as a substrate for a protease.

[000143] In some embodiments, the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM.

[000144] In some embodiments, the activatable antibody comprises a linking peptide between the MM and the CM.

[000145] In some embodiments, the activatable antibody comprises a linking peptide between the CM and the AB.

[000146] In some embodiments, the activatable antibody comprises a first linking peptide (LP1) and a second linking peptide (LP2), and wherein the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-LP1-CM-LP2-AB or AB-LP2-CM-LP1-MM. In some embodiments, the two linking peptides need not be identical to each other. . In some embodiments, each of LP1 and LP2 is a peptide of about 1 to 20 amino acids in length.

[000147]

[000148] In some embodiments, at least one of LP1 or LP2 comprises an amino acid sequence selected from the group consisting of (GS)_n, (GGS)_n, (GSGGS)_n (SEQ ID NO: 363) and (GGGS)_n (SEQ ID NO: 364), where n is an integer of at least one.

[000149] In some embodiments, at least one of LP1 or LP2 comprises an amino acid sequence selected from the group consisting of GGSG (SEQ ID NO: 365), GGSGG (SEQ ID NO: 366), GSGSG (SEQ ID NO: 367), GSGGG (SEQ ID NO: 368), GGGSG (SEQ ID NO: 369), and GSSSG (SEQ ID NO: 370).

[000150] In some embodiments, LP1 comprises the amino acid sequence GSSGGSGGSGGSG (SEQ ID NO: 371), GSSGGSGGSGG (SEQ ID NO: 372), GSSGGSGGSGGS (SEQ ID NO: 373), GSSGGSGGSGGSGGS (SEQ ID NO: 374), GSSGGSGGSG (SEQ ID NO: 375), GSSGGSGGSGS (SEQ ID NO: 376), GGGSSGGS (SEQ ID NO: 65), or GGGSSGG (SEQ ID NO: 1040).

[000151] In some embodiments, LP2 comprises the amino acid sequence GSS, GGS, GGGS (SEQ ID NO: 377), GSSGT (SEQ ID NO: 378) or GSSG (SEQ ID NO: 379).

[000152] In some embodiments, the AB has a dissociation constant (K_d) of about 100 nM or less for binding to PD-1.

[000153] In some embodiments, the activatable antibody includes an antibody or antigen-binding fragment thereof (AB) that specifically binds PD-1. In some embodiments, the antibody or antigen-binding fragment thereof that binds PD-1 is a monoclonal antibody, domain antibody, single chain, Fab fragment, a F(ab')₂ fragment, a scFv, a scAb, a dAb, a single domain heavy chain antibody, or a single domain light chain antibody. In some embodiments, such an antibody or antigen-binding fragment thereof that binds PD-1 is a mouse, other rodent, chimeric, humanized or fully human monoclonal antibody.

[000154] In some embodiments, the activatable antibody comprises a heavy chain amino acid sequence that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the activatable antibody comprises a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000155] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000156] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21. In some embodiments, the activatable anti-PD-1 antibody includes a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 47.

[000157] In some embodiments, the activatable anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 21. In some embodiments, the activatable anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or of SEQ ID NO: 47. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or 47. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 47.

[000158] In some embodiments, the activatable anti-PD-1 antibody includes: (a) a variable heavy chain complementarity determining region 1 (VH CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 653-657; (b) a variable heavy chain complementarity determining region 2 (VH CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 658-663; (c) a variable heavy chain complementarity determining region 3 (VH CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 664-668; (d) a variable light chain complementarity determining region 1 (VL CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 669-677; (e) a variable light chain complementarity determining region 2 (VL CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 678-682; and (f) variable light chain complementarity determining region 3 (VL CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 683-687.

[000159] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH

CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664).

[000160] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[000161] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises

GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[000162]

[000163] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1346, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 626.

[000164] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1514, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 638.

[000165] In some embodiments, the activatable anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346. In some embodiments, the activatable anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626.

[000166] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH

CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707).

[000167] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[000168] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[000169] In some embodiments, the activatable anti-PD-1 antibody includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514. In some embodiments, the activatable anti-PD-1 antibody includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638.

[000170] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH

CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713).

[000171] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[000172] In some embodiments, the activatable anti-PD-1 antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[000173] In some embodiments, the activatable antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the activatable antibody comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the activatable antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence amino acid sequence selected from

the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000174] In some embodiments, the activatable antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the activatable antibody comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the activatable antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 45. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 21, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 47.

[000175] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1346, and a light chain sequence that is at least 90%,

91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 626.

[000176] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1514, and a light chain sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 638.

[000177] In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a heavy chain amino acid sequence selected from the group consisting of the heavy chain sequences shown in Table 7. In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a light chain amino acid sequence selected from the group consisting of the light chain sequences shown in Table 7. In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a heavy chain amino acid sequence selected from the group consisting of the heavy chain sequences shown in Table 7 and a light chain amino acid sequence selected from the group consisting of the light chain sequences shown in Table 7.

[000178] In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of the heavy chain sequences shown in Table 7. In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a light chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of the light chain sequences shown in Table 7. In some embodiments, the AB of the activatable anti-PD-1 antibody comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of the heavy chain sequences shown in Table 7 and a light chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of the light chain sequences shown in Table 7.

[000179] In some embodiments, the AB of the activatable antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%,

97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707).

[000180] In some embodiments, the AB of the activatable antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[000181] In some embodiments, the AB of the activatable antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QQSSNWPRT (SEQ ID NO: 1710).

[000182] In some embodiments, the AB of the activatable antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence

comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713).

[000183] In some embodiments, the AB of the activatable antibody includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[000184] In some embodiments, the AB of the activatable antibody includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to QHSRDLPLT (SEQ ID NO: 1716).

[000185] In some embodiments, the activatable antibody comprises a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, a variable light

chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence shown in Table 8.

[000186] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR1 sequence shown in Table 8; a VH CD2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR3 sequence shown in Table 8.

[000187] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, and a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8.

[000188] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the combination is a combination of the three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000189] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in a combination of the three VH CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8 and in a combination of the three VL CDR sequences (VL CDR1, VL CDR2, and VL CDR3) shown in a single row in Table 8.

[000190] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in a combination of three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000191] In some embodiments, the activatable antibody comprises a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence shown in Table 9; a VH CDR2 sequence shown in Table 9; a VH CDR3 sequence shown in Table 9; a VL CDR1 sequence shown in Table 9; a VL CDR2 sequence shown in Table 9; and a VL CDR3 sequence shown in Table 9.

[000192] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence that includes a sequence that is at least 90%, 91%,

92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR1 sequence shown in Table 9; a VH CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR2 sequence shown in Table 9; a VH CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR3 sequence shown in Table 9; a VL CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR1 sequence shown in Table 9; a VL CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR2 sequence shown in Table 9; and a VL CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR3 sequence shown in Table 9.

[000193] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination is a combination shown in Table 9.

[000194] In some embodiments, the activatable antibody comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in a combination shown in Table 9.

[000195] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675 or SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664.

[000196] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675 or SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000197] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, , 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675 or SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human immunoglobulin heavy chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of 381, 382, 383, and 1807.

[000198] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1

sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675 or SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, a human immunoglobulin heavy chain constant region, and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807, and the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000199] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664.

[000200]

[000201] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, , 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human

immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000202] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human immunoglobulin heavy chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807.

[000203] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 675; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, a human immunoglobulin heavy chain constant region, and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807, and the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000204] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664.

[000205] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000206] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID

NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, and a human immunoglobulin heavy chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807.

[000207] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664, a human immunoglobulin heavy chain constant region, and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807, and the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000208] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain

comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000209] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000210] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, 99; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In

some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000211] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), which is referred to herein as mask PD034 or PD34, a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011, and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000212] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain

constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000213] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), and a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000214] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), which is referred to herein as mask PD034 or PD34, a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), which is also referred to herein as substrate 2012, and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000215] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a

cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000216] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), and a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000217] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), which is referred to herein as mask PD034 or PD34, a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011, and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3

sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000218] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000219] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), and a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination

further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000220] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), which is referred to herein as mask PD034 or PD34, a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), which is also referred to herein as substrate 2001, and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000221] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664. In some embodiments, the combination further comprises a human immunoglobulin

heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000222] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), and a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000223] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSGNH (SEQ ID NO: 361), which is also referred to herein as substrate 2002, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1

sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000224] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSANPRG (SEQ ID NO: 1092), which is also referred to herein as substrate 2003, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In

some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000225] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDDH (SEQ ID NO: 1095), which is also referred to herein as substrate 2006, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000226] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDIH (SEQ ID NO: 1096), which is also referred to herein as substrate 2007, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain

sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000227] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDQH (SEQ ID NO: 1097), which is also referred to herein as 2008, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human

immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000228] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDTH (SEQ ID NO: 1098), which is also referred to herein as substrate 2009, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000229] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDYH (SEQ ID

NO: 1099), which is also referred to herein as 2010, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000230] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSANI (SEQ ID NO: 1102), which is also referred to herein as substrate 2013, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1

sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000231] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising the amino acid sequence TSYCSIEHYPCNTHH (SEQ ID NO: 99), a cleavable moiety (CM) comprising the amino acid sequence ISSGLLSGRSDNI (SEQ ID NO: 1111), which is also referred to herein as substrate 2014, and a combination selected from the group consisting of (a) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 21 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 47; (b) a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VL CDR1 sequence of SEQ ID NO: 676; the VL CDR2 sequence of SEQ ID NO: 678; the VL CDR3 sequence of SEQ ID NO: 683; the VH CDR1 sequence of SEQ ID NO: 653; the VH CDR2 sequence of SEQ ID NO: 658; and the VH CDR3 sequence of SEQ ID NO: 664; and (c) a combination of a variable heavy chain (VH) sequence of SEQ ID NO: 21 and a variable light chain (VL) sequence of SEQ ID NO: 47. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In

some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000232] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1151; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000233] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1152; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000234] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1190; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000235] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1191; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000236] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1198; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000237] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1199; and a heavy chain comprising the amino acid sequence of SEQ ID NO:21.

[000238] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2055; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000239] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2054; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000240] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2057; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000241] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2056; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000242] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2059; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000243] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2058; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2052.

[000244] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2055; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000245] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2054; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000246] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2057; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000247] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2056; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000248] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2059; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000249] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 2058; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 2053.

[000250] In some embodiments, the activatable antibody comprises a masking moiety (MM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1362; a cleavable moiety (CM) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102,

1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 1514 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 638. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000251] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1362; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716). In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000252] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1362; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; the VH sequence of SEQ ID NO: 1514 and the VL sequence of SEQ ID NO: 638. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000253] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1217; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the combination comprises the VH CDR sequences of the variable heavy chain sequence of SEQ ID NO: 1346 and the VL CDR sequences of the variable light chain sequence of SEQ ID NO: 626. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000254] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1217; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; and a combination of a

VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710). In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000255] In some embodiments, the activatable antibody comprises a MM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1217; a CM comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157; the VH sequence of SEQ ID NO: 1346 and the VL sequence of SEQ ID NO: 626. In some embodiments, the combination further comprises a human immunoglobulin heavy chain constant region, a human immunoglobulin light chain constant domain, or both a human immunoglobulin heavy chain constant region and a human immunoglobulin light chain constant domain. In some embodiments, the human immunoglobulin heavy chain constant domain comprises an amino acid sequence selected from the group consisting of SEQ ID NO: 63, 381, 382, 383, and 1807. In some embodiments, the human immunoglobulin light chain constant domain is a human kappa constant domain comprising SEQ ID NO: 61 or SEQ ID NO: 1344.

[000256] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), which is also referred to herein as substrate 2012; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000257] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000258] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), which is also referred to herein as substrate 2001; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000259] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSGNH (SEQ ID NO: 361), which is also referred to herein as substrate 2002; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000260] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANPRG (SEQ ID NO: 1092), which is also referred to herein as substrate 2003; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000261] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDDH (SEQ ID NO: 1095), which is also referred to herein as substrate 2006; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000262] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDIH (SEQ ID NO: 1096), which is also referred to herein as substrate 2007; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000263] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDQH (SEQ ID NO: 1097), which is also referred to herein as substrate 2008; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000264] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDTH (SEQ ID NO: 1098), which is also referred to herein as substrate 2009; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000265] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDYH (SEQ ID NO: 1099), which is also referred to herein as substrate 2010; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000266] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANI (SEQ ID NO: 1102), which is also referred to herein as substrate 2013; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000267] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNI (SEQ ID NO: 1111), which is also referred to herein as substrate 2014; the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain (LC) sequence of SEQ ID NO: 543.

[000268] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), which is also referred to herein as substrate 2012; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000269] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000270] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), which is also referred to herein as substrate 2001; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000271] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSGNH (SEQ ID NO: 361), which is also referred to herein as substrate 2002; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000272] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANPRG (SEQ ID NO: 1092), which is also referred to herein as substrate 2003; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000273] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDDH (SEQ ID NO: 1095), which is also referred to herein as substrate 2006; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000274] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDIH (SEQ ID NO: 1096), which is also referred to herein as substrate 2007; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000275] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDQH (SEQ ID NO: 1097), which is also referred to herein as substrate 2008; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000276] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDTH (SEQ ID NO: 1098), which is also referred to herein as substrate 2009; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000277] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDYH (SEQ ID NO: 1099), which is also referred to herein as substrate 2010; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000278] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANI (SEQ ID NO: 1102), which is also referred to herein as substrate 2013; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000279] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNI (SEQ ID NO: 1111), which is also referred to herein as substrate 2014; the variable heavy chain region (VH) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the variable light chain region (VL) of the light chain (LC) sequence of SEQ ID NO: 543.

[000280] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANP (SEQ ID NO: 1101), which is also referred to herein as substrate 2012; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000281] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000282] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNH (SEQ ID NO: 214), which is also referred to herein as substrate 2001; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000283] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSGNH (SEQ ID NO: 361), which is also referred to herein as substrate 2002; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000284] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANPRG (SEQ ID NO: 1092), which is also referred to herein as substrate 2003; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000285] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDDH (SEQ ID NO: 1095), which is also referred to herein as substrate 2006; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000286] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDIH (SEQ ID NO: 1096), which is also referred to herein as substrate 2007; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000287] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDQH (SEQ ID NO: 1097), which is also referred to herein as substrate 2008; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000288] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDTH (SEQ ID NO: 1098), which is also referred to herein as substrate 2009; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000289] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDYH (SEQ ID NO: 1099), which is also referred to herein as substrate 2010; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000290] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSANI (SEQ ID NO: 1102), which is also referred to herein as substrate 2013; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000291] In some embodiments, the activatable antibody comprises a MM comprising the amino acid sequence ACRICQDHPATKWNS (SEQ ID NO: 549), a CM comprising the amino acid sequence ISSGLLSGRSDNI (SEQ ID NO: 1111), which is also referred to herein as substrate 2014; the heavy chain complementarity determining regions (CDRs) of the heavy chain (HC) sequence of SEQ ID NO: 546, and the light chain complementarity determining regions (CDRs) of the light chain (LC) sequence of SEQ ID NO: 543.

[000292] In some embodiments, the activatable antibody comprises a light chain comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 1808, 1810, 1812, 1814, 1816, 1818, 1820, 1822, 1824, 1826, 1828, and 1830; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000293] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1808; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000294] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1810; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000295] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1812; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000296] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1814; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000297] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1816; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000298] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1818; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000299] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1820; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000300] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1822; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000301] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1824; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000302] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1826; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000303] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1828; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000304] In some embodiments, the activatable antibody comprises a light chain comprising the amino acid sequence of SEQ ID NO: 1830; and a heavy chain comprising the amino acid sequence of SEQ ID NO: 546.

[000305] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence

that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[000306] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000307] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000308] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000309] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[000310] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least

90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000311] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000312] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that comprises a nucleic acid sequence encoding a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a nucleic acid sequence that comprises a nucleic acid sequence encoding a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000313] In some embodiments, the activatable antibody is encoded by a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38. In some embodiments, the activatable antibody is encoded by a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38.

[000314] In some embodiments, the activatable antibody is encoded by a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the activatable antibody is encoded by a

light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[000315] In some embodiments, the activatable antibody is encoded by heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the activatable antibody is encoded by heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[000316] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38.

[000317] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[000318] In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 2, 6, 10, 14, 18, 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 4, 8, 12, 16, 20, 40, 42, 44, 46, 48,

50, 52, 54, 56, 58, and 60. In some embodiments, the activatable antibody is encoded by a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a heavy chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 22, 24, 26, 28, 30, 32, 34, 36, and 38, and a nucleic acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to a light chain nucleic acid sequence selected from the group consisting of SEQ ID NO: 40, 42, 44, 46, 48, 50, 52, 54, 56, 58, and 60.

[000319] In some embodiments, the MM has a dissociation constant, i.e., dissociation constant at an equilibrium state, K_d for binding to the AB that is greater than the K_d for binding of the AB to PD-1.

[000320] In some embodiments, the MM has a K_d for binding to the AB that is no more than the K_d for binding of the AB to PD-1.

[000321] In some embodiments, the MM has a K_d for binding to the AB that is no less than the K_d for binding of the AB to PD-1.

[000322] In some embodiments, the MM has a K_d for binding to the AB that is approximately equal to the K_d for binding of the AB to PD-1.

[000323] In some embodiments, the MM has a K_d for binding to the AB that is less than the K_d for binding of the AB to PD-1.

[000324] In some embodiments, the MM has a K_d for binding to the AB that is no more than 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, or 1,000 fold greater than the K_d for binding of the AB to PD-1. In some embodiments, the MM has a K_d for binding to the AB that is between 1-5, 2-5, 2-10, 5-10, 5-20, 5-50, 5-100, 10-100, 10-1,000, 20-100, 20-1000, or 100-1,000 fold greater than the K_d for binding of the AB to PD-1.

[000325] In some embodiments, the MM has an affinity for binding to the AB that is less than the affinity of binding of the AB to PD-1.

[000326] In some embodiments, the MM has an affinity for binding to the AB that is no more than the affinity of binding of the AB to PD-1.

[000327] In some embodiments, the MM has an affinity for binding to the AB that is approximately equal of the affinity of binding of the AB to PD-1.

[000328] In some embodiments, the MM has an affinity for binding to the AB that is no less than the affinity of binding of the AB to PD-1.

[000329] In some embodiments, the MM has an affinity for binding to the AB that is greater than the affinity of binding of the AB to PD-1.

[000330] In some embodiments, the MM has an affinity for binding to the AB that is 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, or 1,000 less than the affinity of binding of the AB to PD-1. In some embodiments, the MM has an affinity for binding to the AB that is between 1-5, 2-5, 2-10, 5-10, 5-20, 5-50, 5-100, 10-100, 10-1,000, 20-100, 20-1000, or 100-1,000 fold less than the affinity of binding of the AB to PD-1. In some embodiments, the MM has an affinity for binding to the AB that is 2 to 20 fold less than the affinity of binding of the AB to PD-1. In some embodiments, a MM not covalently linked to the AB and at equimolar concentration to the AB does not inhibit the binding of the AB to PD-1.

[000331] In some embodiments, the MM does not interfere or compete with the AB for binding to PD-1 when the activatable antibody is in a cleaved state.

[000332] In some embodiments, the MM is a polypeptide of about 2 to 40 amino acids in length. In some embodiments, the MM is a polypeptide of up to about 40 amino acids in length.

[000333] In some embodiments, the MM is a polypeptide of no more than 40 amino acids in length.

[000334] In some embodiments, the MM polypeptide sequence is different from that of PD-1. In some embodiments, the MM polypeptide sequence is different from that of human PD-1. In some embodiments, the MM polypeptide sequence is no more than 50% identical to any natural binding partner of the AB. In some embodiments, the MM polypeptide sequence is different from that of PD-1 and is no more than 40%, 30%, 25%, 20%, 15%, or 10% identical to any natural binding partner of the AB. In some embodiments, the MM polypeptide sequence is no more than 50% identical to human PD-1. In some embodiments, the MM polypeptide sequence is different from that of PD-1 and is no more than 40%, 30%, 25%, 20%, 15%, or 10% identical to human PD-1.

[000335] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 66-213, 384-514, 548-571, 1206-1295, and 1351-1465. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 66-213, 384-514, 548-571, 1206-1295, and 1351-1465. In some embodiments, the MM comprises an amino acid sequence selected from the group

consisting of SEQ ID NOs: 66, 67, 70, 71, 74, 77, 81, 82, 84, 90, 91, 93, and 99. In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 66, 71, 74, 77, 82, 84, 90, 91, 93, and 99. In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1217. In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1362.

[000336] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 66-213. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 66-213.

[000337] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 384-514. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 384-514.

[000338] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 548-571. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 548-571.

[000339] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1295. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 1206-1295.

[000340] In some embodiments, the MM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1465. In some embodiments, the MM comprises an amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NOs: 1351-1465.

[000341] In some embodiments, the antibody or activatable antibody comprises a heavy chain comprising an amino acid selected from the group consisting of SEQ ID NOs: 2052 and 2053.

[000342] In some embodiments, the antibody or activatable antibody comprises a light chain comprising an amino acid selected from the group consisting of SEQ ID NOs: 2054 to 2059.

[000343] In some embodiments, the antibody or activatable antibody comprises a heavy chain comprising an amino acid selected from the group consisting of SEQ ID NOs: 2052 and 2053 and a light chain comprising an amino acid selected from the group consisting of SEQ ID NOs: 2054 to 2059.

[000344] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least two times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000345] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least five times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000346] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 10 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000347] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 20 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000348] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 40 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000349] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 100 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000350] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 1000 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000351] In some embodiments, the coupling of the MM to the AB reduces the ability of the AB to bind PD-1 such that the dissociation constant (K_d) of the AB when coupled to the MM towards PD-1 is at least 10,000 times greater than the K_d of the AB when not coupled to the MM towards PD-1.

[000352] In some embodiments, in the presence of PD-1, the MM reduces the ability of the AB to bind PD-1 by at least 90% when the CM is uncleaved, as compared to when the CM is cleaved when assayed *in vitro* using a target displacement assay such as, for example, the assay described in PCT Publication No. WO 2010/081173, the contents of which are hereby incorporated by reference in their entirety.

[000353] In some embodiments, the protease that cleaves the CM is active, e.g., up-regulated, in diseased tissue, and the protease cleaves the CM in the activatable antibody when the activatable antibody is exposed to the protease.

[000354] In some embodiments, the protease is produced by a tumor that is in proximity to T cells that express PD-1, and the protease cleaves the CM in the activatable antibody when the activatable antibody is exposed to the protease. In some embodiments, the protease is produced by a tumor that is co-localized with T cells that express PD-1, and the protease cleaves the CM in the activatable antibody when the activatable antibody is exposed to the protease.

[000355] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least twofold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state (i.e., when the activatable antibody is in the cleaved state), the AB binds PD-1.

[000356] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least fivefold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state (i.e., when the activatable antibody is in the cleaved state), the AB binds PD-1.

[000357] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 10-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state (i.e., when the activatable antibody is in the cleaved state), the AB binds PD-1.

[000358] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 20-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state (i.e., when the activatable antibody is in the cleaved state), the AB binds PD-1.

[000359] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 40-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state, the AB binds PD-1.

[000360] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 50-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state, the AB binds PD-1.

[000361] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 100-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state, the AB binds PD-1.

[000362] In some embodiments, the CM is positioned in the activatable antibody such that when the activatable antibody is in the uncleaved state, binding of the activatable antibody to PD-1 is reduced to occur with a dissociation constant that is at least 200-fold greater than the dissociation constant of an unmodified AB binding to PD-1, whereas in the cleaved state, the AB binds PD-1.

[000363] In some embodiments, the CM is a polypeptide of up to 15 amino acids in length.

[000364] In some embodiments, the CM is a polypeptide that includes a first cleavable moiety (CM1) that is a substrate for at least one matrix metalloprotease (MMP) and a second cleavable moiety (CM2) that is a substrate for at least one serine protease (SP). In some embodiments, each of the CM1 substrate sequence and the CM2 substrate sequence of the CM1-CM2 substrate is independently a polypeptide of up to 15 amino acids in length.

[000365] In some embodiments, the CM is a substrate for at least one protease that is or is believed to be up-regulated in cancer. In some embodiments, the CM is a substrate for at least one protease that is or is believed to be up-regulated in inflammation. In some embodiments, the CM is a substrate for at least one protease that is or is believed to be up-regulated in autoimmunity.

[000366] In some embodiments, the CM is a substrate for at least one protease selected from the group consisting of a matrix metalloprotease (MMP), thrombin, a neutrophil elastase, a cysteine protease, legumain, and a serine protease, such as matriptase (MT-SP1), and urokinase (uPA). Without being bound by theory, it is believed that these proteases are up-regulated in at least one of cancer, inflammation, and/or autoimmunity.

[000367] Exemplary substrates include but are not limited to substrates cleavable by one or more of the following enzymes or proteases listed in Table 3.

[000368] In some embodiments, the CM is selected for use with a specific protease, for example a protease that is known to be co-localized with the target of the activatable antibody. For example, the protease is produced by a tumor that is in proximity to T cells that express PD-1. In some embodiments, the protease is produced by a tumor that is co-localized with T cells that express PD-1.

[000369] In some embodiments, the CM is a substrate for at least one MMP. Examples of MMPs include the MMPs listed in the Table 3. In some embodiments, the CM is a substrate for a protease selected from the group consisting of MMP 9, MMP14, MMP1, MMP3, MMP13, MMP17, MMP11, and MMP19. In some embodiments the CM is a substrate for MMP9. In some embodiments, the CM is a substrate for MMP14.

[000370] In some embodiments, the CM is a substrate that includes the sequence TGRGPSWV (SEQ ID NO: 295); SARGPSRW (SEQ ID NO: 319); TARGPSFK (SEQ ID NO: 297); LSGRSDNH (SEQ ID NO: 294); GGWHTGRN (SEQ ID NO: 320); HTGRSGAL (SEQ ID NO: 321); PLTGRSGG (SEQ ID NO: 296); AARGPAIH (SEQ ID NO: 322);

RGPAFNPM (SEQ ID NO: 323); SSRGPAYL (SEQ ID NO: 324); RGPATPIM (SEQ ID NO: 325); RGPA (SEQ ID NO: 326); GGQPSGMWGW (SEQ ID NO: 327); FPRPLGITGL (SEQ ID NO: 328); VHMPGLGFLGP (SEQ ID NO: 302); SPLTGRSG (SEQ ID NO: 329); SAGFSLPA (SEQ ID NO: 330); LAPLGLQRR (SEQ ID NO: 331); SGGPLGVR (SEQ ID NO: 332); PLGL (SEQ ID NO: 333); LSGRSGNH (SEQ ID NO: 1157); SGRSANPRG (SEQ ID NO: 1158); LSGRSDDH (SEQ ID NO: 1161); LSGRSDIH (SEQ ID NO: 1162); LSGRSDQH (SEQ ID NO: 1165); LSGRSDTH (SEQ ID NO: 1166); LSGRSDYH (SEQ ID NO: 1169); LSGRSDNP (SEQ ID NO: 1520); LSGRSANP (SEQ ID NO: 1695); LSGRSANI (SEQ ID NO: 1696); and/or LSGRSDNI (SEQ ID NO: 1697).

[000371] In some embodiments, the CM comprises the amino acid sequence LSGRSDNH (SEQ ID NO: 294). In some embodiments, the CM comprises the amino acid sequence TGRGPSWV (SEQ ID NO: 295). In some embodiments, the CM comprises the amino acid sequence PLTGRSGG (SEQ ID NO: 296). In some embodiments, the CM comprises the amino acid sequence GGQPSGMWGW (SEQ ID NO: 327). In some embodiments, the CM comprises the amino acid sequence FPRPLGITGL (SEQ ID NO: 328). In some embodiments, the CM comprises the amino acid sequence VHMPGLGFLGP (SEQ ID NO: 302). In some embodiments, the CM comprises the amino acid sequence PLGL (SEQ ID NO: 333). In some embodiments, the CM comprises the amino acid sequence SARGPSRW (SEQ ID NO: 319). In some embodiments, the CM comprises the amino acid sequence TARGPSFK (SEQ ID NO: 297). In some embodiments, the CM comprises the amino acid sequence GGWHTGRN (SEQ ID NO: 320). In some embodiments, the CM comprises the amino acid sequence HTGRSGAL (SEQ ID NO: 321). In some embodiments, the CM comprises the amino acid sequence AARGPAIH (SEQ ID NO: 322). In some embodiments, the CM comprises the amino acid sequence RGPAFNPM (SEQ ID NO: 323). In some embodiments, the CM comprises the amino acid sequence SSRGPAYL (SEQ ID NO: 324). In some embodiments, the CM comprises the amino acid sequence RGPATPIM (SEQ ID NO: 325). In some embodiments, the CM comprises the amino acid sequence RGPA (SEQ ID NO: 326). In some embodiments, the CM comprises the amino acid sequence LSGRSGNH (SEQ ID NO: 1157). In some embodiments, the CM comprises the amino acid sequence SGRSANPRG (SEQ ID NO: 1158). In some embodiments, the CM comprises the amino acid sequence LSGRSDDH (SEQ ID NO: 1161). In some embodiments, the CM comprises the amino acid sequence LSGRSDIH (SEQ ID NO: 1162). In

some embodiments, the CM comprises the amino acid sequence LSGRSDQH (SEQ ID NO: 1165). In some embodiments, the CM comprises the amino acid sequence LSGRSDTH (SEQ ID NO: 1166). In some embodiments, the CM comprises the amino acid sequence LSGRSDYH (SEQ ID NO: 1169). In some embodiments, the CM comprises the amino acid sequence LSGRSDNP (SEQ ID NO: 1520). In some embodiments, the CM comprises the amino acid sequence LSGRSANP (SEQ ID NO: 1695). In some embodiments, the CM comprises the amino acid sequence LSGRSANI (SEQ ID NO: 1696). In some embodiments, the CM comprises the amino acid sequence LSGRSDNI (SEQ ID NO: 1697).

[000372] In some embodiments, the CM is a substrate for an MMP and includes the sequence ISSGLSS (SEQ ID NO: 334); QNQALRMA (SEQ ID NO: 305); AQNLLGMV (SEQ ID NO: 304); STFPFGMF (SEQ ID NO: 307); PVGYTSSL (SEQ ID NO: 335); DWLYWPPI (SEQ ID NO: 336); ISSGLSS (SEQ ID NO: 308); LKAAPRWA (SEQ ID NO: 337); GPSHLVLT (SEQ ID NO: 338); LPGGLSPW (SEQ ID NO: 339); MGLFSEAG (SEQ ID NO: 340); SPLPLRVP (SEQ ID NO: 341); RMHLRSLG (SEQ ID NO: 342); LAAPLGLL (SEQ ID NO: 306); AVGLLAPP (SEQ ID NO: 303); LLAPSHRA (SEQ ID NO: 343); PAGLWLDP (SEQ ID NO: 309); MIAPVAYR (SEQ ID NO: 1698); RPSPMWAY (SEQ ID NO: 1699); WATPRPMR (SEQ ID NO: 1700); FRLLDWQW (SEQ ID NO: 1701); ISSGL (SEQ ID NO: 1702); ISSGLS (SEQ ID NO: 1703); and/or ISSGLL (SEQ ID NO: 1704).

[000373] In some embodiments, the CM comprises the amino acid sequence ISSGLSS (SEQ ID NO: 334). In some embodiments, the CM comprises the amino acid sequence QNQALRMA (SEQ ID NO: 305). In some embodiments, the CM comprises the amino acid sequence AQNLLGMV (SEQ ID NO: 304). In some embodiments, the CM comprises the amino acid sequence STFPFGMF (SEQ ID NO: 307). In some embodiments, the CM comprises the amino acid sequence PVGYTSSL (SEQ ID NO: 335). In some embodiments, the CM comprises the amino acid sequence DWLYWPPI (SEQ ID NO: 336). In some embodiments, the CM comprises the amino acid sequence ISSGLSS (SEQ ID NO: 308). In some embodiments, the CM comprises the amino acid sequence LKAAPRWA (SEQ ID NO: 337). In some embodiments, the CM comprises the amino acid sequence GPSHLVLT (SEQ ID NO: 338). In some embodiments, the CM comprises the amino acid sequence LPGGLSPW (SEQ ID NO: 339). In some embodiments, the CM comprises the amino acid sequence MGLFSEAG (SEQ ID NO: 340). In some embodiments, the CM comprises the amino acid sequence

SPLPLRVP (SEQ ID NO: 341). In some embodiments, the CM comprises the amino acid sequence RMHLRSLG (SEQ ID NO: 342). In some embodiments, the CM comprises the amino acid sequence LAAPLGLL (SEQ ID NO: 306). In some embodiments, the CM comprises the amino acid sequence AVGLLAPP (SEQ ID NO: 303). In some embodiments, the CM comprises the amino acid sequence LLAPSHRA (SEQ ID NO: 343). In some embodiments, the CM comprises the amino acid sequence PAGLWLDP (SEQ ID NO: 309). In some embodiments, the CM comprises the amino acid sequence MIAPVAYR (SEQ ID NO: 1698). In some embodiments, the CM comprises the amino acid sequence RPSPMWAY (SEQ ID NO: 1699). In some embodiments, the CM comprises the amino acid sequence WATPRPMR (SEQ ID NO: 1700). In some embodiments, the CM comprises the amino acid sequence FRLLDWQW (SEQ ID NO: 1701). In some embodiments, the CM comprises the amino acid sequence ISSGL (SEQ ID NO: 1702). In some embodiments, the CM comprises the amino acid sequence ISSGLLS (SEQ ID NO: 1703). In some embodiments, the CM comprises the amino acid sequence ISSGLL (SEQ ID NO: 1704).

[000374] In some embodiments, the CM is a substrate for thrombin. In some embodiments, the CM is a substrate for thrombin and includes the sequence GPRSFGL (SEQ ID NO: 344) or GPRSFG (SEQ ID NO: 345). In some embodiments, the CM comprises the amino acid sequence GPRSFGL (SEQ ID NO: 344). In some embodiments, the CM comprises the amino acid sequence GPRSFG (SEQ ID NO: 345).

[000375] In some embodiments, the CM comprises an amino acid sequence selected from the group consisting of NTLSGRSENHSG (SEQ ID NO: 298); NTLSGRSGNHGS (SEQ ID NO: 299); TSTSGRSANPRG (SEQ ID NO: 300); TSGRSANP (SEQ ID NO: 301); VAGRSMRP (SEQ ID NO: 310); VVPEGRRS (SEQ ID NO: 311); ILPRSPAF (SEQ ID NO: 312); MVLGRSLL (SEQ ID NO: 313); QGRAITFI (SEQ ID NO: 314); SPRSIMLA (SEQ ID NO: 315); and SMLRSMPL (SEQ ID NO: 316).

[000376] In some embodiments, the CM comprises the amino acid sequence NTLSGRSENHSG (SEQ ID NO: 298). In some embodiments, the CM comprises the amino acid sequence NTLSGRSGNHGS (SEQ ID NO: 299). In some embodiments, the CM comprises the amino acid sequence TSTSGRSANPRG (SEQ ID NO: 300). In some embodiments, the CM comprises the amino acid sequence TSGRSANP (SEQ ID NO: 301). In some embodiments, the CM comprises the amino acid sequence VAGRSMRP (SEQ ID NO: 310). In some

embodiments, the CM comprises the amino acid sequence VVPEGRRS (SEQ ID NO: 311). In some embodiments, the CM comprises the amino acid sequence ILPRSPAF (SEQ ID NO: 312). In some embodiments, the CM comprises the amino acid sequence MVLGRSLL (SEQ ID NO: 313). In some embodiments, the CM comprises the amino acid sequence QGRAITFI (SEQ ID NO: 314). In some embodiments, the CM comprises the amino acid sequence SPRSIMLA (SEQ ID NO: 315). In some embodiments, the CM comprises the amino acid sequence SMLRSMPL (SEQ ID NO: 316).

[000377] In some embodiments, the CM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294-361, 1092-1112, 1157, 1158, 1161, 1162, 1165, 1166, 1169, 1520, and 1695-1704. In some embodiments, the CM comprises an amino acid sequence selected from the group consisting of SEQ ID NOs: 214, 294, 300, 302, 303, 305, 308, 318, 347, 361, 1092-1102, 1111, and 1157.

[000378] In some embodiments, the CM is a substrate for a neutrophil elastase. In some embodiments, the CM is a substrate for a serine protease. In some embodiments, the CM is a substrate for uPA. In some embodiments, the CM is a substrate for legumain. In some embodiments, the CM is a substrate for matriptase. In some embodiments, the CM is a substrate for a cysteine protease. In some embodiments, the CM is a substrate for a cysteine protease, such as a cathepsin.

[000379] In some embodiments, the CM is a CM1-CM2 substrate and includes the sequence ISSGLLSGRSDNH (SEQ ID NO: 214); ISSGLLSSGGSGGSLSGRSDNH (SEQ ID NO: 346); AVGLLAPPGGTSTSGRSANPRG (SEQ ID NO: 347); TSTSGRSANPRGGGAVGLLAPP (SEQ ID NO: 348); VHMPLGFLGPGGTSTSGRSANPRG (SEQ ID NO: 349); TSTSGRSANPRGGGVHMPLGFLGP (SEQ ID NO: 350); AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318); LSGRSDNHGGAVGLLAPP (SEQ ID NO: 351); VHMPLGFLGPGGLSGRSDNH (SEQ ID NO: 352); LSGRSDNHGGVHMPLGFLGP (SEQ ID NO: 353); LSGRSDNHGGSGGSISSGLLSS (SEQ ID NO: 354); LSGRSGNHGGSGGSISSGLLSS (SEQ ID NO: 355); ISSGLLSSGGSGGSLSGRSGNH (SEQ ID NO: 356); LSGRSDNHGGSGGSGQNQALRMA (SEQ ID NO: 357); QNQALRMAGGSGGSLSGRSDNH (SEQ ID NO: 358); LSGRSGNHGGSGGSGQNQALRMA (SEQ ID NO: 359); QNQALRMAGGSGGSLSGRSGNH (SEQ ID NO: 360); ISSGLLSGRSGNH (SEQ ID NO: 361); ISSGLLSGRSANPRG (SEQ ID

NO: 1092); AVGLLAPPTSGRSANPRG (SEQ ID NO: 1093); AVGLLAPPSGRSANPRG (SEQ ID NO: 1094); ISSGLLSGRSDDH (SEQ ID NO: 1095); ISSGLLSGRSDIH (SEQ ID NO: 1096); ISSGLLSGRSDQH (SEQ ID NO: 1097); ISSGLLSGRSDTH (SEQ ID NO: 1098); ISSGLLSGRSDYH (SEQ ID NO: 1099); ISSGLLSGRSDNP (SEQ ID NO: 1100); ISSGLLSGRSANP (SEQ ID NO: 1101); ISSGLLSGRSANI (SEQ ID NO: 1102); AVGLLAPPGGLSGRSDDH (SEQ ID NO: 1103); AVGLLAPPGGLSGRSDIH (SEQ ID NO: 1104); AVGLLAPPGGLSGRSDQH (SEQ ID NO: 1105); AVGLLAPPGGLSGRSDTH (SEQ ID NO: 1106); AVGLLAPPGGLSGRSDYH (SEQ ID NO: 1107); AVGLLAPPGGLSGRSDNP (SEQ ID NO: 1108); AVGLLAPPGGLSGRSANP (SEQ ID NO: 1109); AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110); ISSGLLSGRSDNI (SEQ ID NO: 1111); and/or AVGLLAPPGGLSGRSDNI (SEQ ID NO: 1112).

[000380] In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDNH (SEQ ID NO: 214), which is also referred to herein as substrate 2001. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSSGGSGGSLSGRSDNH (SEQ ID NO: 346), which is also referred to herein as substrate 1001/LP'/0001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGTSTSGRSANPRG (SEQ ID NO: 347), which is also referred to herein as substrate 1004/LP'/0003, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence TSTSGRSANPRGGGAVGLLAPP (SEQ ID NO: 348), which is also referred to herein as substrate 0003/LP'/1004, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence VHMPGLGFLGPGGTSTSGRSANPRG (SEQ ID NO: 349), which is also referred to herein as substrate 1003/LP'/0003, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence TSTSGRSANPRGGGVHMPGLGFLGP (SEQ ID NO: 350), which is also referred to herein as substrate 0003/LP'/1003, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318), which is also referred to herein as substrate 1004/LP'/0001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In

some embodiments, the CM1-CM2 substrate includes the sequence LSGRSDNHGGAVGLLAPP (SEQ ID NO: 351), which is also referred to herein as substrate 0001/LP'/1004, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence VHMPLGFLGPGGLSGRSDNH (SEQ ID NO: 352), which is also referred to herein as substrate 1003/LP'/0001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence LSGRSDNHGGVHMPLGFLGP (SEQ ID NO: 353), which is also referred to herein as substrate 0001/LP'/1003, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence LSGRSDNHGGSGGSISSGLLSS (SEQ ID NO: 354), which is also referred to herein as substrate 0001/LP'/1001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence LSGRSGNHGGSGGSISSGLLSS (SEQ ID NO: 355), which is also referred to herein as substrate 0002/LP'/1001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSSGGSGGSLSGRSGNH (SEQ ID NO: 356), which is also referred to herein as substrate 1001/LP'/0002, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence LSGRSDNHGGSGGSQNQALRMA (SEQ ID NO: 357), which is also referred to herein as substrate 0001/LP'/1002, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence QNQALRMAGGSGGSLSGRSDNH (SEQ ID NO: 358), which is also referred to herein as substrate 1002/LP'/0001, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence LSGRSGNHGGSGGSQNQALRMA (SEQ ID NO: 359), which is also referred to herein as substrate 0002/LP'/1002, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence QNQALRMAGGSGGSLSGRSGNH (SEQ ID NO: 360), which is also referred to herein as substrate 1002/LP'/0002, where LP' as used in this CM1-CM2 substrate is

the amino acid sequence GGSGGS (SEQ ID NO: 1519). In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSGNH (SEQ ID NO: 361), which is also referred to herein as substrate 2002. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSANPRG (SEQ ID NO: 1092), which is also referred to herein as substrate 2003. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPTSGRSANPRG (SEQ ID NO: 1093), which is also referred to herein as substrate 2004. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPSGRSANPRG (SEQ ID NO: 1094), which is also referred to herein as substrate 2005. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDDH (SEQ ID NO: 1095), which is also referred to herein as substrate 2006. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDIH (SEQ ID NO: 1096), which is also referred to herein as substrate 2007. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDQH (SEQ ID NO: 1097), which is also referred to herein as substrate 2008. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDTH (SEQ ID NO: 1098), which is also referred to herein as substrate 2009. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDYH (SEQ ID NO: 1099), which is also referred to herein as substrate 2010. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDNP (SEQ ID NO: 1100), which is also referred to herein as substrate 2011. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSANP (SEQ ID NO: 1101), which is also referred to herein as substrate 2012. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSANI (SEQ ID NO: 1102), which is also referred to herein as substrate 2013. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDDH (SEQ ID NO: 1103), which is also referred to herein as substrate 3006. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDIH (SEQ ID NO: 1104), which is also referred to herein as substrate 3007. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDQH (SEQ ID NO: 1105), which is also referred to herein as substrate 3008. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDTH (SEQ ID NO: 1106), which is also referred to herein as substrate 3009. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDYH (SEQ ID NO: 1107), which is also referred to herein as substrate

3010. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSDNP (SEQ ID NO: 1108), which is also referred to herein as substrate 3011. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSANP (SEQ ID NO: 1109), which is also referred to herein as substrate 3012. In some embodiments, the CM1-CM2 substrate includes the sequence AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110), which is also referred to herein as substrate 3013. In some embodiments, the CM1-CM2 substrate includes the sequence ISSGLLSGRSDNI (SEQ ID NO: 1111), which is also referred to herein as substrate 2014. In some embodiments, the CM1-CM2 substrate includes the sequence and/or AVGLLAPPGGLSGRSDNI (SEQ ID NO: 1112), which is also referred to herein as substrate 3014. In some embodiments, the CM1-CM2 substrate includes the sequence GLSGRSDNHGGAVGLLAPP (SEQ ID NO: 1970), which is also referred to herein as substrate 0001/LP'/1004, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG. In some embodiments, the CM1-CM2 substrate includes the sequence GLSGRSDNHGGVHMPLGFLGP (SEQ ID NO: 1971), which is also referred to herein as substrate 0001/LP'/1003, where LP' as used in this CM1-CM2 substrate is the amino acid sequence GG.

[000381] In some embodiments, the CM is a substrate for at least two proteases. In some embodiments, each protease is selected from the group consisting of those shown in Table 3. In some embodiments, the CM is a substrate for at least two proteases, wherein one of the proteases is selected from the group consisting of a MMP, thrombin, a neutrophil elastase, a cysteine protease, uPA, legumain and matriptase and the other protease is selected from the group consisting of those shown in Table 3. In some embodiments, the CM is a substrate for at least two proteases selected from the group consisting of a MMP, thrombin, a neutrophil elastase, a cysteine protease, uPA, legumain and matriptase.

[000382] In some embodiments, the activatable antibody includes at least a first CM and a second CM. In some embodiments, the first CM and the second CM are each polypeptides of no more than 15 amino acids long. In some embodiments, the first CM and the second CM in the activatable antibody in the uncleaved state have the structural arrangement from N-terminus to C-terminus as follows: MM-CM1-CM2-AB or AB-CM2-CM1-MM. In some embodiments, at least one of the first CM and the second CM is a polypeptide that functions as a substrate for a protease selected from the group consisting of a MMP, thrombin, a neutrophil elastase, a

cysteine protease, uPA, legumain, and matriptase. In some embodiments, the first CM is cleaved by a first cleaving agent selected from the group consisting of a MMP, thrombin, a neutrophil elastase, a cysteine protease, uPA, legumain, and matriptase in a target tissue and the second CM is cleaved by a second cleaving agent in a target tissue. In some embodiments, the other protease is selected from the group consisting of those shown in Table 3. In some embodiments, the first cleaving agent and the second cleaving agent are the same protease selected from the group consisting of a MMP, thrombin, a neutrophil elastase, a cysteine protease, uPA, legumain, and matriptase, and the first CM and the second CM are different substrates for the enzyme. In some embodiments, the first cleaving agent and the second cleaving agent are the same protease selected from the group consisting of those shown in Table 3. In some embodiments, the first cleaving agent and the second cleaving agent are different proteases. In some embodiments, the first cleaving agent and the second cleaving agent are co-localized in the target tissue. In some embodiments, the first CM and the second CM are cleaved by at least one cleaving agent in the target tissue.

[000383] In some embodiments, the activatable antibody is exposed to and cleaved by a protease such that, in the activated or cleaved state, the activated antibody includes a light chain amino acid sequence that includes at least a portion of LP2 and/or CM sequence after the protease has cleaved the CM.

[000384] In some embodiments, the activatable antibody also includes an agent conjugated to the AB. In some embodiments, the agent conjugated to the AB or the AB of an activatable antibody is a therapeutic agent. In some embodiments, the agent is conjugated to the AB via a cleavable linker. In some embodiments, the agent is conjugated to the AB via a linker that includes at least one CM1-CM2 substrate sequence. In some embodiments, the agent is conjugated to the AB via a noncleavable linker.

[000385] In some embodiments, the agent is an anti-inflammatory agent.

[000386] In some embodiments, the activatable antibody also includes a detectable moiety. In some embodiments, the detectable moiety is a diagnostic agent.

[000387] In some embodiments, the activatable antibody also includes a signal peptide. In some embodiments, the signal peptide is conjugated to the activatable antibody via a spacer. In some embodiments, the spacer is conjugated to the activatable antibody in the absence of a signal peptide. In some embodiments, the spacer is joined directly to the MM of the activatable

antibody. In some embodiments, the spacer is joined directly to the MM of the activatable antibody in the structural arrangement from N-terminus to C-terminus of spacer-MM-CM-AB. An example of a spacer joined directly to the N-terminus of MM of the activatable antibody is selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG (SEQ ID NO: 2045), QG (SEQ ID NO: 2046), and G. In some embodiments, the spacer includes at least the amino acid sequence QGQSGQG (SEQ ID NO: 362). In some embodiments, the spacer includes at least the amino acid sequence QGQSGQ (SEQ ID NO: 913). In some embodiments, the spacer includes at least the amino acid sequence QGQSG (SEQ ID NO: 914). In some embodiments, the spacer includes at least the amino acid sequence QGQS (SEQ ID NO: 915). In some embodiments, the spacer includes at least the amino acid sequence QGQ (SEQ ID NO: 916). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 917). In some embodiments, the spacer includes at least the amino acid residue Q. In some embodiments, the spacer includes at least the amino acid sequence GQSGQG (SEQ ID NO: 2042). In some embodiments, the spacer includes at least the amino acid sequence QSGQG (SEQ ID NO: 2043). In some embodiments, the spacer includes at least the amino acid sequence SGQG (SEQ ID NO: 2044). In some embodiments, the spacer includes at least the amino acid sequence GQG (SEQ ID NO: 2045). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 2046). In some embodiments, the spacer includes at least the amino acid sequence G. In some embodiments, the activatable antibody does not include a spacer sequence.

[000388] In some embodiments, the AB of the activatable antibody naturally contains one or more disulfide bonds. In some embodiments, the AB can be engineered to include one or more disulfide bonds.

[000389] In some embodiments, the serum half-life of the activatable antibody is longer than that of the corresponding antibody; e.g., the pK of the activatable antibody is longer than that of the corresponding antibody. In some embodiments, the serum half-life of the activatable antibody is similar to that of the corresponding antibody. In some embodiments, the serum half-life of the activatable antibody is at least 15 days when administered to an organism. In some

embodiments, the serum half-life of the activatable antibody is at least 12 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 11 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 10 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 9 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 8 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 7 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 6 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 5 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 4 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 3 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 2 days when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 24 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 20 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 18 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 16 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 14 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 12 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 10 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 8 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 6 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 4 hours when administered to an organism. In some embodiments, the serum half-life of the activatable antibody is at least 3 hours when administered to an organism.

[000390] In some embodiments, the activatable anti-PD-1 antibody is monospecific.

[000391] In some embodiments, the activatable anti-PD-1 antibody is multispecific, *e.g.*, by way of non-limiting example, bispecific or trifunctional. In some embodiments, the activatable anti-PD-1 antibody is formulated as part of a pro-Bispecific T Cell Engager (BITE) molecule, *i.e.*, the BITE includes a masking moiety and a cleavable moiety. In some embodiments, the activatable anti-PD-1 antibody is formulated as part of a pro-Chimeric Antigen Receptor (CAR) modified T cell, modified NK cell, or other modified immune effector cell. In some embodiments, an activatable anti-PD-1 antibody is formulated as part of another engineered receptor on an immune effector cell; *i.e.*, the pro-CAR or other pro-engineered receptor includes a masking moiety and a cleavable moiety.

[000392] In some embodiments, the activatable antibody or antigen-binding fragment thereof is incorporated in a multispecific activatable antibody or antigen-binding fragment thereof, where at least one arm of the multispecific activatable antibody specifically binds PD-1. In some embodiments, the activatable antibody or antigen-binding fragment thereof is incorporated in a bispecific activatable antibody or antigen-binding fragment thereof, where at least one arm of the bispecific activatable antibody specifically binds PD-1.

[000393] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a heavy chain amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[000394] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000395] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, a heavy chain amino acid sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid sequence selected from the group consisting

of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, a heavy chain amino acid sequence comprising an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45 or SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 45. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from amino acid sequence SEQ ID NO: 21 and a light chain that comprises or is derived from amino acid sequence SEQ ID NO: 47.

[000396] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 21. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or of SEQ ID NO: 47. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45 or 47. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a

bispecific activatable antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 45. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 21, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 47.

[000397] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody includes: (a) a variable heavy chain complementarity determining region 1 (VH CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 653-657; (b) a variable heavy chain complementarity determining region 2 (VH CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 658-663; (c) a variable heavy chain complementarity determining region 3 (VH CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 664-668; (d) a variable light chain complementarity determining region 1 (VL CDR1) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 669-677; (e) a variable light chain complementarity determining region 2 (VL CDR2) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 678-682; and (f) variable light chain complementarity determining region 3 (VL CDR3) comprising an amino acid sequence selected from the group consisting of SEQ ID NOs: 683-687

[000398] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664).

[000399] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein

as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[000400] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675) or RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDSYGISFMN (SEQ ID NO: 675); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3

sequence comprises QQSKDVPWT (SEQ ID NO: 683). In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GFTFSGYAMS (SEQ ID NO: 653); the VH CDR2 sequence comprises YISNSGGNAH (SEQ ID NO: 658); and the VH CDR3 sequence comprises EDYGTSPFVY (SEQ ID NO: 664); the VL CDR1 sequence comprises RASESVDAYGISFMN (SEQ ID NO: 676); the VL CDR2 sequence comprises AASNQGS (SEQ ID NO: 678); and the VL CDR3 sequence comprises QQSKDVPWT (SEQ ID NO: 683).

[000401] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1346, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 626.

[000402] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a heavy chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 1514, and a light chain that comprises or is derived from the amino acid sequence of SEQ ID NO: 638.

[000403] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1346, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 626.

[000404] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence, a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH

CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); and the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707).

[000405] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[000406] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GITFSNSG (SEQ ID NO: 1705); the VH CDR2 sequence comprises IWYDGSKR (SEQ ID NO: 1706); the VH CDR3 sequence comprises TNDDY (SEQ ID NO: 1707); the VL CDR1 sequence comprises QSVSSY (SEQ ID NO: 1708); the VL CDR2 sequence comprises DAS (SEQ ID NO: 1709); and the VL CDR3 sequence comprises QQSSNWPRT (SEQ ID NO: 1710).

[000407] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the complementarity determining region (CDR) sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes the combination of the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of the CDR sequences of the heavy chain amino acid sequence of SEQ ID NO: 1514, and the CDR sequences of the light chain amino acid sequence of SEQ ID NO: 638.

[000408] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence,

a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, and a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); and the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713).

[000409] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[000410] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, includes a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein the VH CDR1 sequence comprises GYTFTNYY (SEQ ID NO: 1711); the VH CDR2 sequence comprises INPSNGGT (SEQ ID NO: 1712); the VH CDR3 sequence comprises RRDYRFDMGFDY (SEQ ID NO: 1713); the VL CDR1 sequence comprises KGVSTSGYSY (SEQ ID NO: 1714); the VL CDR2 sequence comprises LAS (SEQ ID NO: 1715); and the VL CDR3 sequence comprises QHSRDLPLT (SEQ ID NO: 1716).

[000411] In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37. In some embodiments, at least one arm of the multispecific activatable antibody, *e.g.*, a bispecific activatable antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37.

[000412] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000413] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59.

[000414] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1346, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 626.

[000415] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, includes a heavy chain amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 1514, and a light chain amino acid that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98% or 99% identical to the amino acid sequence of SEQ ID NO: 638.

[000416] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a combination of a variable heavy chain complementarity determining region 1 (VH CDR1, also referred to herein as CDRH1) sequence,

a variable heavy chain complementarity determining region 2 (VH CDR2, also referred to herein as CDRH2) sequence, a variable heavy chain complementarity determining region 3 (VH CDR3, also referred to herein as CDRH3) sequence, a variable light chain complementarity determining region 1 (VL CDR1, also referred to herein as CDRL1) sequence, a variable light chain complementarity determining region 2 (VL CDR2, also referred to herein as CDRL2) sequence, and a variable light chain complementarity determining region 3 (VL CDR3, also referred to herein as CDRL3) sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence shown in Table 8.

[000417] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence, a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence, wherein at least one CDR sequence is selected from the group consisting of a VH CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR1 sequence shown in Table 8; a VH CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR2 sequence shown in Table 8; a VH CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VH CDR3 sequence shown in Table 8; a VL CDR1 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR1 sequence shown in Table 8; a VL CDR2 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR2 sequence shown in Table 8; and a VL CDR3 sequence that includes a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to a VL CDR3 sequence shown in Table 8.

[000418] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, and a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8.

[000419] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein the combination is a combination of the three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000420] In some embodiments, at least one arm of the multispecific activatable antibody, e.g., a bispecific activatable antibody, comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, a VH CDR3 sequence shown in a single row in Table 8, and a VL CDR1 sequence, a VL CDR2 sequence, and a VL CDR3 sequence shown in a single row in Table 8, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding VH CDR sequence shown in a single row in Table 8 and the corresponding VL CDR sequence shown in a single row in Table 8.

[000421] In some embodiments, the antibody comprises a heavy chain that comprises a combination of a VH CDR1 sequence, a VH CDR2 sequence, and a VH CDR3 sequence, wherein each CDR sequence in the combination comprises a sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the corresponding CDR sequence in a combination of three heavy chain CDR sequences (VH CDR1, VH CDR2, VH CDR3) shown in a single row in Table 8.

[000422] In some embodiments, the anti-PD-1 antibodies and/or activatable anti-PD-1 antibodies described herein are used in conjunction with one or more additional agents or a combination of additional agents. Suitable additional agents include other experimental anti-cancer agents in development for specific applications, current pharmaceutical and/or surgical therapies for an intended application, such as, for example, cancer. For example, the anti-PD-1 antibodies and/or activatable anti-PD-1 antibodies can be used in conjunction with an additional chemotherapeutic or anti-neoplastic agent.

[000423] In some embodiments, the additional agent includes at least one experimental anti-cancer agent in development for specific applications. In some embodiments, the additional agent includes at least one agent that is already approved for one indication but is being tested in another. In some embodiments, the additional agent includes at least one agent that is not currently approved in any indication but is being tested in one or more indications for the purposes of obtaining regulatory approval. In some embodiments, the additional agent includes

at least one agent that is a pharmaceutical therapy for an intended application. In some embodiments, the additional agent includes at least one agent that is a surgical therapy for cancer. In some embodiments, the additional agent includes at least one agent that is a pharmaceutical therapy for an intended application. In some embodiments, the additional agent includes at least one agent that is a surgical therapy for cancer. In some embodiments, the additional agent is a chemotherapeutic agent. In some embodiments, the additional agent is an anti-neoplastic agent. In some embodiments, the additional agents are a combination of any two or more of these agents.

[000424] In some embodiments, the additional agent(s) is a chemotherapeutic agent, such as a chemotherapeutic agent selected from the group consisting of docetaxel, paclitaxel, abraxane (i.e., albumin-conjugated paclitaxel), doxorubicin, oxaliplatin, carboplatin, cisplatin, irinotecan, and gemcitabine.

[000425] In some embodiments, the additional agent(s) is a checkpoint inhibitor, a kinase inhibitor, an agent targeting inhibitors in the tumor microenvironment, and/or a T cell or NK agonist. In some embodiments, the additional agent(s) is radiation therapy, alone or in combination with another additional agent(s) such as a chemotherapeutic or anti-neoplastic agent. In some embodiments, the additional agent(s) is a vaccine, an oncovirus, and/or a DC-activating agent such as, by way of non-limiting example, a toll-like receptor (TLR) agonist and/or α -CD40. In some embodiments, the additional agent(s) is a tumor-targeted antibody designed to kill the tumor via ADCC or via direct conjugation to a toxin (e.g., an antibody drug conjugate (ADC)).

[000426] In some embodiments, the checkpoint inhibitor is an inhibitor of a target selected from the group consisting of CTLA-4, LAG-3, PD-1, PD-1, TIGIT, TIM-3, B7H4, BTLA, and Vista. In some embodiments, the kinase inhibitor is selected from the group consisting of B-RAFi, MEKi, and Btk inhibitors, such as ibrutinib. In some embodiments, the kinase inhibitor is crizotinib. In some embodiments, the B-RAF inhibitor is vemurafenib. In some embodiments, the tumor microenvironment inhibitor is selected from the group consisting of an IDO inhibitor, an α -CSF1R inhibitor, an α -CCR4 inhibitor, a TGF-beta, a myeloid-derived suppressor cell, or a T-regulatory cell. In some embodiments, the agonist is selected from the group consisting of Ox40, GITR, CD137, ICOS, CD27, and HVEM.

[000427] In some embodiments, the inhibitor is a CTLA-4 inhibitor. In some embodiments, the inhibitor is a LAG-3 inhibitor. In some embodiments, the inhibitor is a PD-1 inhibitor. In some embodiments, the inhibitor is a PD-1 inhibitor. In some embodiments, the inhibitor is a TIGIT inhibitor. In some embodiments, the inhibitor is a TIM-3 inhibitor. In some embodiments, the inhibitor is a B7H4 inhibitor. In some embodiments, the inhibitor is a Vista inhibitor. In some embodiments, the inhibitor is a B-RAFi inhibitor. In some embodiments, the inhibitor is a MEKi inhibitor. In some embodiments, the inhibitor is a Btk inhibitor. In some embodiments, the inhibitor is ibrutinib. In some embodiments, the inhibitor is crizotinib. In some embodiments, the inhibitor is an IDO inhibitor. In some embodiments, the inhibitor is an α -CSF1R inhibitor. In some embodiments, the inhibitor is an α -CCR4 inhibitor. In some embodiments, the inhibitor is a TGF-beta. In some embodiments, the inhibitor is a myeloid-derived suppressor cell. In some embodiments, the inhibitor is a T-regulatory cell.

[000428] In some embodiments, the agonist is Ox40. In some embodiments, the agonist is GITR. In some embodiments, the agonist is CD137. In some embodiments, the agonist is ICOS. In some embodiments, the agonist is CD27. In some embodiments, the agonist is HVEM.

[000429] In some embodiments, the anti-PD-1 antibody and/or activatable antibody is administered during and/or after treatment in combination with one or more additional agents such as, for example, a chemotherapeutic agent, an anti-inflammatory agent, and/or an immunosuppressive agent. In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent are formulated into a single therapeutic composition, and the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and additional agent are administered simultaneously. Alternatively, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and additional agent are separate from each other, *e.g.*, each is formulated into a separate therapeutic composition, and the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent are administered simultaneously, or the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent are administered at different times during a treatment regimen. For example, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody is administered prior to the administration of the additional agent, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody is administered subsequent to the administration of the additional agent, or the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent are administered in an alternating fashion. As described herein, the anti-PD-1

antibody and/or activatable anti-PD-1 antibody and additional agent are administered in single doses or in multiple doses.

[000430] In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) are administered simultaneously. For example, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) can be formulated in a single composition or administered as two or more separate compositions. In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) are administered sequentially, or the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent are administered at different times during a treatment regimen.

[000431] In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody is administered during and/or after treatment in combination with one or more additional agents such as, by way of non-limiting example, a chemotherapeutic agent, an anti-inflammatory agent, and/or an immunosuppressive agent, such as an alkylating agent, an anti-metabolite, an anti-microtubule agent, a topoisomerase inhibitor, a cytotoxic antibiotic, and/or any other nucleic acid damaging agent. In some embodiments, the additional agent is a taxane, such as paclitaxel (e.g., Abraxane®). In some embodiments, the additional agent is an anti-metabolite, such as gemcitabine. In some embodiments, the additional agent is an alkylating agent, such as platinum-based chemotherapy, such as carboplatin or cisplatin. In some embodiments, the additional agent is a targeted agent, such as a kinase inhibitor, e.g., sorafenib or erlotinib. In some embodiments, the additional agent is a targeted agent, such as another antibody, e.g., a monoclonal antibody (e.g., bevacizumab), a bispecific antibody, or a multispecific antibody. In some embodiments, the additional agent is a proteasome inhibitor, such as bortezomib or carfilzomib. In some embodiments, the additional agent is an immune modulating agent, such as lenolidomide or IL-2. In some embodiments, the additional agent is radiation. In some embodiments, the additional agent is an agent considered standard of care by those skilled in the art. In some embodiments, the additional agent is a chemotherapeutic agent well known to those skilled in the art.

[000432] In some embodiments, the additional agent is another antibody or antigen-binding fragment thereof and/or another activatable antibody or antigen-binding fragment thereof. In some embodiments the additional agent is another antibody or antigen-binding fragment thereof and/or another activatable antibody or antigen-binding fragment thereof against the same target

as the first antibody or antigen-binding fragment thereof and/or an activatable antibody or antigen-binding fragment thereof, e.g., against PD-1. In some embodiments the additional agent is another antibody or antigen-binding fragment thereof and/or another activatable antibody or antigen-binding fragment thereof against a target different than the target of the first antibody or antigen-binding fragment thereof, the first conjugated antibody or antigen-binding fragment thereof and/or activatable antibody or antigen-binding fragment thereof.

[000433] As a non-limiting example, the antibody or antigen-binding fragment and/or the AB of an activatable antibody is a binding partner for any target listed in Table 1.

Table 1: Exemplary Targets

1-92-LFA-3	CD52	DL44	HVEM	LIF-R	STEAP1
Alpha-4 integrin	CD56	DLK1	Hyaluronidase	Lewis X	STEAP2
Alpha-V integrin	CD64	DLL4	ICOS	LIGHT	TAG-72
alpha4beta1 integrin	CD70	DPP-4	IFNalpha	LRP4	TAPA1
alpha4beta7 integrin	CD71	DSG1	IFNbeta	LRRC26	TGFbeta
AGR2	CD74	EGFR	IFNgamma	MCSP	TIGIT
Anti-Lewis-Y		EGFRviii	IgE	Mesothelin	TIM-3
Apelin J receptor	CD80	Endothelin B receptor (ETBR)	IgE Receptor (FcεRI)	MRP4	TLR2
APRIL	CD81	ENPP3	IGF	MUC1	TLR4
B7-H4	CD86	EpCAM	IGF1R	Mucin-16 (MUC16, CA-125)	TLR6
BAFF	CD95	EPHA2	IL1B	Na/K ATPase	TLR7
BTLA	CD117	EPHB2	IL1R	Neutrophil elastase	TLR8
C5 complement	CD125	ERBB3	IL2	NGF	TLR9
C-242	CD132 (IL-2RG)	F protein of RSV	IL11	Nicastrin	TMEM31
CA9	CD133	FAP	IL12	Notch Receptors	TNFalpha
CA19-9 (Lewis a)	CD137	FGF-2	IL12p40	Notch 1	TNFR
Carbonic anhydrase 9	CD138	FGF8	IL-12R, IL-12Rbeta1	Notch 2	TNFRS12 A
CD2	CD166	FGFR1	IL13	Notch 3	TRAIL-R1

CD3	CD172A	FGFR2	IL13R	Notch 4	TRAIL-R2
CD6	CD248	FGFR3	IL15	NOV	Transferrin
CD9	CDH6	FGFR4	IL17	OSM-R	Transferrin receptor
CD11a	CEACAM5 (CEA)	Folate receptor	IL18	OX-40	TRK-A
CD19	CEACAM6 (NCA-90)	GAL3ST1	IL21	PAR2	TRK-B
CD20	CLAUDIN-3	G-CSF	IL23	PDGF-AA	uPAR
CD22	CLAUDIN-4	G-CSFR	IL23R	PDGF-BB	VAP1
CD24	cMet	GD2	IL27/IL27R (wsx1)	PDGFRalpha	VCAM-1
CD25	Collagen	GITR	IL29	PDGFRbeta	VEGF
CD27	Cripto	GLUT1	IL-31R	PD-1	VEGF-A
CD28	CSFR	GLUT4	IL31/IL31R	PD-L1	VEGF-B
CD30	CSFR-1	GM-CSF	IL2R	PD-L2	VEGF-C
CD33	CTLA-4	GM-CSFR	IL4	Phosphatidyl-serine	VEGF-D
CD38	CTGF	GP IIb/IIIa receptors	IL4R	P1GF	VEGFR1
CD40	CXCL10	Gp130	IL6, IL6R	PSCA	VEGFR2
CD40L	CXCL13	GPIIB/IIIA	Insulin Receptor	PSMA	VEGFR3
CD41	CXCR1	GPNMB	Jagged Ligands	RAAG12	VISTA
CD44	CXCR2	GRP78	Jagged 1	RAGE	WISP-1
CD44v6		HER2/neu	Jagged 2	SLC44A4	WISP-2
CD47	CXCR4	HGF	LAG-3	Sphingosine 1 Phosphate	WISP-3
CD51	CYR61	hGH			

[000434] As a non-limiting example, the antibody or antigen-binding fragment and/or the AB of an activatable antibody is or is derived from an antibody listed in Table 2.

Table 2: Exemplary sources for Abs

Antibody Trade Name (antibody name)	Target
Avastin™ (bevacizumab)	VEGF
Lucentis™ (ranibizumab)	VEGF
Erbix™ (cetuximab)	EGFR
Vectibix™ (panitumumab)	EGFR
Remicade™ (infliximab)	TNF α
Humira™ (adalimumab)	TNF α
Tysabri™ (natalizumab)	Integrin α 4
Simulect™ (basiliximab)	IL2R
Soliris™ (eculizumab)	Complement C5

Raptiva™ (efalizumab)	CD11a
Bexxar™ (tositumomab)	CD20
Zevalin™ (ibritumomab tiuxetan)	CD20
Rituxan™ (rituximab)	CD20
Ocrelizumab	CD20
Arzerra™ (ofatumumab)	CD20
Gazyva™ (Obinutuzumab)	CD20
Zenapax™ (daclizumab)	CD25
Adcetris™ (brentuximab vedotin)	CD30
Myelotarg™ (gemtuzumab)	CD33
Mylotarg™ (gemtuzumab ozogamicin)	CD33
Campath™ (alemtuzumab)	CD52
ReoPro™ (abiximab)	Glycoprotein receptor IIb/IIIa
Xolair™ (omalizumab)	IgE
Herceptin™ (trastuzumab)	Her2
Kadcyla™ (trastuzumab emtansine)	Her2
Synagis™ (palivizumab)	F protein of RSV
(ipilimumab)	CTLA-4
(tremelimumab)	CTLA-4
Hu5c8	CD40L
(pertuzumab)	Her2-neu
(ertumaxomab)	CD3/Her2-neu
Orencia™ (abatacept)	CTLA-4
(tanezumab)	NGF
(bavituximab)	Phosphatidylserine
(zalutumumab)	EGFR
(mapatumumab)	EGFR
(matuzumab)	EGFR
(nimotuzumab)	EGFR
ICR62	EGFR
mAb 528	EGFR
CH806	EGFR
MDX-447	EGFR/CD64
(edrecolomab)	EpCAM
RAV12	RAAG12
huJ591	PSMA
Enbrel™ (etanercept)	TNF-R
Amevive™ (alefacept)	1-92-LFA-3
Antril™, Kineret™ (ankinra)	IL-1Ra
GC1008	TGFbeta
	Notch, e.g., Notch 1
	Jagged 1 or Jagged 2
(adecatumumab)	EpCAM
(figitumumab)	IGF1R
(tocilizumab)	IL-6 receptor

Stelara™ (ustekinumab)	IL-12/IL-23
Prolia™ (denosumab)	RANKL

[000435] In some embodiments, the additional antibody or antigen binding fragment thereof and/or activatable antibody or antigen binding fragment thereof is a monoclonal antibody, domain antibody, single chain, Fab fragment, a F(ab')₂ fragment, a scFv, a scAb, a dAb, a single domain heavy chain antibody, or a single domain light chain antibody. In some embodiments, the additional antibody or antigen binding fragment thereof and/or activatable antibody or antigen binding fragment thereof is a mouse, other rodent, chimeric, humanized or fully human monoclonal antibody.

[000436] The disclosure also provides methods of producing an anti-PD-1 antibody and/or activatable anti-PD-1 antibody polypeptide by culturing a cell under conditions that lead to expression of the polypeptide, wherein the cell comprises an isolated nucleic acid molecule encoding an antibody and/or an activatable antibody described herein, and/or vectors that include these isolated nucleic acid sequences. The disclosure provides methods of producing an antibody and/or activatable antibody by culturing a cell under conditions that lead to expression of the antibody and/or activatable antibody, wherein the cell comprises an isolated nucleic acid molecule encoding an antibody and/or an activatable antibody described herein, and/or vectors that include these isolated nucleic acid sequences.

[000437] The invention also provides a method of manufacturing activatable antibodies that in an activated state binds PD-1 by (a) culturing a cell comprising a nucleic acid construct that encodes the activatable antibody under conditions that lead to expression of the activatable antibody, wherein the activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM), and an antibody or an antigen binding fragment thereof (AB) that specifically binds PD-1, (i) wherein the CM is a polypeptide that functions as a substrate for a protease; and (ii) wherein the CM is positioned in the activatable antibody such that, when the activatable antibody is in an uncleaved state, the MM interferes with specific binding of the AB to PD-1 and in a cleaved state the MM does not interfere or compete with specific binding of the AB to PD-1; and (b) recovering the activatable antibody. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

[000438] In some embodiments, the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-

MM. In some embodiments, the activatable antibody comprises a linking peptide between the MM and the CM. In some embodiments, the activatable antibody comprises a linking peptide between the CM and the AB. In some embodiments, the activatable antibody comprises a first linking peptide (LP1) and a second linking peptide (LP2), and wherein the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-LP1-CM-LP2-AB or AB-LP2-CM-LP1-MM. In some embodiments, the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: spacer-MM-LP1-CM-LP2-AB or AB-LP2-CM-LP1-MM-spacer.

[000439] In some embodiments, the two linking peptides need not be identical to each other.

[000440] In some embodiments, at least one of LP1 or LP2 comprises an amino acid sequence selected from the group consisting of (GS)_n, (GGS)_n, (GSGGS)_n (SEQ ID NO: 363) and (GGGS)_n (SEQ ID NO: 364), where n is an integer of at least one.

[000441] In some embodiments, at least one of LP1 or LP2 comprises an amino acid sequence selected from the group consisting of GGSG (SEQ ID NO: 365), GGSGG (SEQ ID NO: 366), GSGSG (SEQ ID NO: 367), GSGGG (SEQ ID NO: 368), GGGSG (SEQ ID NO: 369), and GSSSG (SEQ ID NO: 370).

[000442] In some embodiments, LP1 comprises the amino acid sequence GSSGGSGGSGGSG (SEQ ID NO: 371), GSSGGSGGSGG (SEQ ID NO: 372), GSSGGSGGSGGS (SEQ ID NO: 373), GSSGGSGGSGGSGGS (SEQ ID NO: 374), GSSGGSGGSG (SEQ ID NO: 375), GSSGGSGGSGS (SEQ ID NO: 376), GGGSSGGS (SEQ ID NO: 65), or GGGSSGG (SEQ ID NO: 1040).

[000443] In some embodiments, LP2 comprises the amino acid sequence GSS, GGS, GGGS (SEQ ID NO: 377), GSSGT (SEQ ID NO: 378) or GSSG (SEQ ID NO: 379).

[000444] The invention provides methods of preventing, delaying the progression of, treating, alleviating a symptom of, or otherwise ameliorating a disorder or disease in a subject by administering to a subject in need thereof a therapeutically effective amount of an anti-PD-1 antibody, conjugated anti-PD-1 antibody, activatable anti-PD-1 antibody and/or conjugated activatable anti-PD-1 antibody described herein.

[000445] The invention also provides a method of reducing immune suppression comprising administering to a subject in need thereof a therapeutically effective amount of an

anti-PD-1 antibody and/or activatable anti-PD-1 antibody described herein. In some embodiments, the immune suppression is suppression of T cell activity. In some embodiments, the immune suppression is mediated by engagement of PD-1 on T cells to PD-L1 or PD-L2 on tumor cells or other immune cells. In some embodiments, the invention provides a method to reduce or inhibit binding of PD-L1 (also referred to herein as PDL1) and/or PD-L2 (also referred to herein as PDL2), to PD-1 on T cells. The ligands PD-L1 and/or PD-L2 are often found on tumor cells or other immune cells.

[000446] The invention also provides methods of preventing, delaying the progression of, treating, alleviating a symptom of, or otherwise ameliorating cancer in a subject by administering a therapeutically effective amount of an anti-PD-1 antibody and/or activatable anti-PD-1 antibody described herein to a subject in need thereof. PD-1 is known to be expressed on immune cells, such as T cells, in a variety of cancers, such as, by way of non-limiting example, melanoma, non-small cell lung cancer, nasopharyngeal cancer, glioblastoma/mixed glioma, colon adenocarcinoma, hepatocellular carcinoma, urothelial cancer, multiple myeloma, ovarian cancer, gastric carcinoma, esophageal cancer, pancreatic cancer, renal cell carcinoma (RCC), breast cancer, lymphomas, such as Hodgkin's lymphoma, and leukemias. (See e.g., Chen et al., "Molecular Pathways: Next-Generation Immunotherapy – Inhibiting Programmed Death-Ligand 1 and Programmed Death-1," Clin. Can. Res., vol. 18: 6580-6587 (2012), the contents of which are hereby incorporated by reference in their entirety).

[000447] In some embodiments, the cancer is a bladder cancer, a bone cancer, a breast cancer, a carcinoid, a cervical cancer, a cholangiocarcinoma, a colon cancer, an endometrial cancer, a glioma, a head and neck cancer, a liver cancer, a lung cancer, a lymphoma, such as Hodgkin's lymphoma, a melanoma, an ovarian cancer, a pancreatic cancer, a prostate cancer, a renal cancer, a sarcoma, a skin cancer, a stomach cancer, a testis cancer, a thymus cancer, a thyroid cancer, a urogenital cancer, and/or a urothelial cancer.

[000448] In some embodiments, the cancer is selected from the group consisting of melanoma (MEL), renal cell carcinoma (RCC), squamous non-small cell lung cancer (NSCLC), non-squamous NSCLC, colorectal cancer (CRC), castration-resistant prostate cancer (CRPC), hepatocellular carcinoma (HCC), squamous cell carcinoma of the head and neck, thymoma, carcinomas of the esophagus, ovary, gastrointestinal tract and breast, or a hematologic

malignancy such as multiple myeloma, B-cell lymphoma, T-cell lymphoma, Hodgkin's lymphoma, primary mediastinal B-cell lymphoma, and chronic myelogenous leukemia.

[000449] The invention also provides methods of treating cancer patients with an autoimmune or inflammatory disease by administering a therapeutically effective amount of an anti-PD-1 antibody and/or activatable anti-PD-1 antibody described herein to a subject in need thereof. In some embodiments, the autoimmune disease is colitis, RA, pancreatitis, diabetes, or pneumonitis.

[000450] An anti-PD-1 antibody and/or an activatable anti-PD-1 antibody used in any of the embodiments of these methods and uses can be administered at any stage of the disease. For example, such an anti-PD-1 antibody and/or activatable anti-PD-1 antibody can be administered to a patient suffering cancer of any stage, from early to metastatic. The terms subject and patient are used interchangeably herein.

[000451] In some embodiments, the subject is a mammal, such as a human, non-human primate, companion animal (*e.g.*, cat, dog, horse), farm animal, work animal, or zoo animal. In some embodiments, the subject is a human. In some embodiments, the subject is a companion animal. In some embodiments, the subject is an animal in the care of a veterinarian.

[000452] The anti-PD-1 antibody and/or activatable anti-PD-1 antibody and therapeutic formulations thereof are administered to a subject suffering from or susceptible to a disease or disorder associated with immune suppression, such as immune suppression mediated by engagement of PD-1 on T cells by PD-L1 or PD-L2 on tumor cells or other immune cells. In some embodiments, the immune suppression is suppression of T cell activity. A subject suffering from or susceptible to a disease or disorder associated with such immune suppression is identified using any of a variety of methods known in the art. For example, subjects suffering from cancer or other neoplastic condition are identified using any of a variety of clinical and/or laboratory tests such as, physical examination and blood, urine and/or stool analysis to evaluate health status. For example, subjects suffering from inflammation and/or an inflammatory disorder are identified using any of a variety of clinical and/or laboratory tests such as physical examination and/or bodily fluid analysis, *e.g.*, blood, urine and/or stool analysis, to evaluate health status.

[000453] Administration of an anti-PD-1 antibody and/or activatable anti-PD-1 antibody to a patient suffering from a disease or disorder associated with immune suppression, such as

immune suppression mediated by engagement of PD-1 on T cells by PD-L1 or PD-L2 on tumor cells or other immune cells, is considered successful if any of a variety of laboratory or clinical objectives is achieved. For example, administration of an anti-PD-1 antibody, and/or activatable anti-PD-1 antibody to a patient suffering from a disease or disorder associated with such immune suppression is considered successful if one or more of the symptoms associated with the disease or disorder is alleviated, reduced, inhibited or does not progress to a further, *i.e.*, worse, state. Administration of an anti-PD-1 antibody and/or conjugated activatable anti-PD-1 antibody to a patient suffering from a disease or disorder associated with immune suppression, such as immune suppression mediated by engagement of PD-1 on T cells to PD-L1 or PD-L2 on tumor cells or other immune cells, is considered successful if the disease or disorder enters remission or does not progress to a further, *i.e.*, worse, state.

[000454] In some embodiments, the anti-PD-1 antibody and/or conjugated activatable anti-PD-1 antibody is administered during and/or after treatment in combination with one or more additional agents such as, for example, a chemotherapeutic agent, an anti-inflammatory agent, and/or an immunosuppressive agent. In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) are administered simultaneously. For example, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) can be formulated in a single composition or administered as two or more separate compositions. In some embodiments, the anti-PD-1 antibody and/or activatable anti-PD-1 antibody and the additional agent(s) are administered sequentially.

[000455] The invention also provides methods and kits for using the activatable anti-PD-1 antibodies in a variety of diagnostic and/or prophylactic indications. For example, the invention provides methods and kits for detecting the presence or absence of a cleaving agent and a target of interest in a subject or a sample by (i) contacting a subject or sample with an anti-PD-1 activatable antibody, wherein the anti-PD-1 activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, and an antigen binding domain or fragment thereof (AB) that specifically binds the target of interest, wherein the anti-PD-1 activatable antibody in an uncleaved, non-activated state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to PD-1, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB and is not a

modified form of a natural binding partner of the AB; and (b) wherein, when the AB is in an uncleaved, non-activated state, the MM interferes with specific binding of the AB to PD-1, and when the AB is in a cleaved, activated state the MM does not interfere or compete with specific binding of the AB to PD-1; and (ii) measuring a level of activated anti-PD-1 activatable antibody in the subject or sample, wherein a detectable level of activated anti-PD-1 activatable antibody in the subject or sample indicates that the cleaving agent and PD-1 are present in the subject or sample and wherein no detectable level of activated anti-PD-1 activatable antibody in the subject or sample indicates that the cleaving agent, PD-1 or both the cleaving agent and PD-1 are absent in the subject or sample.

[000456] In some embodiments, the activatable anti-PD-1 antibody is an activatable anti-PD-1 antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable anti-PD-1 antibody is not conjugated to an agent. In some embodiments, the activatable anti-PD-1 antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable anti-PD-1 antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000457] In some embodiments of these methods and kits, the activatable anti-PD-1 antibody includes a detectable label. In some embodiments of these methods and kits, the detectable label includes an imaging agent, a contrasting agent, an enzyme, a fluorescent label, a chromophore, a dye, one or more metal ions, or a ligand-based label. In some embodiments of these methods and kits, the imaging agent comprises a radioisotope. In some embodiments of these methods and kits, the radioisotope is indium or technetium. In some embodiments of these methods and kits, the contrasting agent comprises iodine, gadolinium or iron oxide. In some embodiments of these methods and kits, the enzyme comprises horseradish peroxidase, alkaline phosphatase, or β -galactosidase. In some embodiments of these methods and kits, the fluorescent label comprises yellow fluorescent protein (YFP), cyan fluorescent protein (CFP), green fluorescent protein (GFP), modified red fluorescent protein (mRFP), red fluorescent protein tdimer2 (RFP tdimer2), HCRED, or a europium derivative. In some embodiments of these methods and kits, the luminescent label comprises an N- methylacrydium derivative. In some embodiments of these methods, the label comprises an Alexa Fluor[®] label, such as Alex Fluor[®]

680 or Alexa Fluor[®] 750. In some embodiments of these methods and kits, the ligand-based label comprises biotin, avidin, streptavidin or one or more haptens.

[000458] In some embodiments of these methods and kits, the subject is a mammal. In some embodiments of these methods, the subject is a human. In some embodiments, the subject is a non-human mammal, such as a non-human primate, companion animal (e.g., cat, dog, horse), farm animal, work animal, or zoo animal. In some embodiments, the subject is a rodent.

[000459] In some embodiments of these methods and kits, the method is an *in vivo* method. In some embodiments of these methods, the method is an *in situ* method. In some embodiments of these methods, the method is an *ex vivo* method. In some embodiments of these methods, the method is an *in vitro* method.

[000460] In some embodiments of the methods and kits, the method is used to identify or otherwise refine a patient population suitable for treatment with an anti-PD-1 activatable antibody of the disclosure, followed by treatment by administering that activatable anti-PD-1 antibody to a subject in need thereof. For example, patients that test positive for both the target (e.g., PD-1) and a protease that cleaves the substrate in the cleavable moiety (CM) of the anti-PD-1 activatable antibody being tested in these methods are identified as suitable candidates for treatment with such an anti-PD-1 activatable antibody comprising such a CM, and the patient is then administered a therapeutically effective amount of the activatable anti-PD-1 antibody that was tested. Likewise, patients that test negative for either or both of the target (e.g., PD-1) and the protease that cleaves the substrate in the CM in the activatable antibody being tested using these methods might be identified as suitable candidates for another form of therapy. In some embodiments, such patients can be tested with other anti-PD-1 activatable antibodies until a suitable anti-PD-1 activatable antibody for treatment is identified (e.g., an anti-PD-1 activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, the patient is then administered a therapeutically effective amount of the activatable anti-PD-1 antibody for which the patient tested positive. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

[000461] Pharmaceutical compositions according to the invention can include an antibody of the invention and a carrier. These pharmaceutical compositions can be included in kits, such as, for example, diagnostic kits.

Brief Description of the Drawings

[000462] Figure 1 is graph depicting the results of an ELISA binding assay using various murine antibodies that bind human PD-1 (i.e., anti-hPD-1 antibodies) referred to herein as 245-M3, also referred to herein as M3 (VH: SEQ ID NO: 9; VL: SEQ ID NO: 11), 245-M5, also referred to herein as M5 (VH: SEQ ID NO: 13; VL: SEQ ID NO: 15), 136-M13, also referred to herein as M13 (VH: SEQ ID NO: 1; VL: SEQ ID NO: 3), 136-M19, also referred to herein as M19 (VH: SEQ ID NO: 5; VL: SEQ ID NO: 7), and 136-M14, also referred to herein as M14 (VH: SEQ ID NO: 17; VL: SEQ ID NO: 19). The anti-PD-1 antibodies nivolumab (also referred to herein as “nivo” or “NV1”) and pembrolizumab (also referred to herein as “pembro” or “PM1” or “PM1 AB”) were used as positive controls.

[000463] Figure 2 is a series of graphs demonstrating that various murine anti-hPD-1 antibodies block binding of the ligands human PD-L1 (hPD-L1) and human PD-L2 (hPD-L2) to human PD-1.

[000464] Figure 3 is a series of graphs demonstrating that various murine anti-hPD-1 antibodies block binding of the ligands human PD-L1 (hPD-L1) and human PD-L2 (hPD-L2) to human PD-1.

[000465] Figure 4 is a graph depicting binding to human PD-1 of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.0 (VH: SEQ ID NO: 21; VL: SEQ ID NO: 39), A1.2 (VH: SEQ ID NO: 21; VL: SEQ ID NO: 43), and A1.4 (VH: SEQ ID NO: 21; VL: SEQ ID NO: 45) as determined by ELISA. The anti-PD-1 antibody pembrolizumab (“PM1”) was used as positive control.

[000466] Figure 5 is a graph depicting binding to human PD-1 of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.5 (VH: SEQ ID NO: 21; VL: SEQ ID NO: 47), A1.6 (VH: SEQ ID NO: 21; VL: SEQ ID NO: 49), Ba2 (VH: SEQ ID NO: 29; VL: SEQ ID NO: 57), Bb2 (VH: SEQ ID NO: 31; VL: SEQ ID NO: 57), C1.1 (VH: SEQ ID NO: 33; VL: SEQ ID NO: 41), and D4 (VH: SEQ ID NO: 37; VL: SEQ ID NO: 59) as determined by ELISA. The anti-PD-1 antibodies nivolumab (“NV1”) and pembrolizumab (“PM1”) were used as positive controls.

[000467] Figure 6 is a series of graphs depicting the ability of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.0, A1.2, and A1.4 to inhibit binding between human PD-1 (hPD-1) and both human PDL-1 (hPDL-1) and human PDL-2 (hPDL-2) as

determined by ELISA. The anti-PD-1 antibody pembrolizumab (“PM1”) was used as positive control.

[000468] Figure 7 is a series of graphs depicting the ability of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.5 and C1.1 to inhibit binding between hPD-1 and both hPDL-1 and hPDL-2 as determined by ELISA. The anti-PD-1 antibodies nivolumab (“NV1”) and pembrolizumab (“PM1”) were used as positive controls.

[000469] Figure 8 is a graph depicting the ability of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.5, Ba2, Bb2, C1.1, and D4 to inhibit binding between hPD-1 and hPDL-1 as determined by ELISA. The anti-PD-1 antibodies nivolumab (“NV1”) and pembrolizumab (“PM1”) were used as positive controls.

[000470] Figure 9 is a graph depicting the ability of various anti-hPD-1 antibodies of the disclosure referred to herein as A1.5, Ba2, Bb2, C1.1, and D4 to inhibit binding between hPD-1 and hPDL-2 as determined by ELISA. The anti-PD-1 antibodies nivolumab (“NV1”) and pembrolizumab (“PM1”) were used as positive controls.

[000471] Figure 10 is a graph depicting that the anti-PD-1 antibody referred to herein as A1.5 specifically binds hPD-1 as determined by ELISA. The A1.5 anti-PD-1 antibody was tested against a panel of human and mouse proteins.

[000472] Figures 11A and 11B are a series of graphs depicting the ability of the anti-PD-1 antibody A1.5 to block Fab fragments of the anti-PD-1 antibodies nivolumab (“NV1”) and pembrolizumab (“PM1”) from binding to PD-1. These graphs also demonstrate that the anti-PD-1 antibody A1.5 binds a distinct epitope from either NV1 and/or PM1.

[000473] Figure 12 is a graph demonstrating that the anti-PD-1 antibody A1.5 has a potency similar to that of nivolumab (“NV1”) and pembrolizumab (“PM1”) measured in a T-cell restimulation assay. In Figure 12, PMBCs were treated as follows: PBMCs: CMV+ HemaCare Donor C were plated at 2×10^5 cells/well; stimulated with 5 µg/mL CMV lysate +/- anti-PD-1 or isotype control; IFN-γ ELISA assay performed on day 4 supernatant.

[000474] Figure 13 is a series of graphs demonstrating that the anti-PD-1 antibodies A1.5 and Ba2 bind monomeric human PD-1 (hPD1) with similar or higher affinity than the anti-PD-1 antibodies nivolumab (“Nivo”) and pembrolizumab (“Pembro”).

[000475] Figure 14 is a series of graphs depicting the binding to human PD-1 by anti-PD-1 antibody A1.4 and by various activatable anti-PD-1 antibodies that include the various masking

moieties referred to herein as PD01 (SEQ ID NO: 66), PD02 (SEQ ID NO: 67), PD03 (SEQ ID NO: 68), PD08 (SEQ ID NO: 73), PD09 (SEQ ID NO: 74), and PD10 (SEQ ID NO: 75) as determined by ELISA.

[000476] Figure 15 is a series of graphs depicting the binding to hPD-1 by anti-PD-1 antibody A1.5 and by various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody A1.5 and the masking moieties referred to herein as PD01 (SEQ ID NO: 66), PD02 (SEQ ID NO: 67), PD03 (SEQ ID NO: 68), PD05 (SEQ ID NO: 70), PD06 (SEQ ID NO: 71) (Plate 1); PD08 (SEQ ID NO: 73), PD09 (SEQ ID NO: 74), PD10 (SEQ ID NO: 75), PD11 (SEQ ID NO: 76), PD12 (SEQ ID NO: 77) (Plate 2); PD13 (SEQ ID NO: 78), PD14 (SEQ ID NO: 79), PD15 (SEQ ID NO: 80), PD16 (SEQ ID NO: 81), PD17 (SEQ ID NO: 82) (Plate 3); PD18 (SEQ ID NO: 83), PD19 (SEQ ID NO: 84), PD20 (SEQ ID NO: 85), PD21 (SEQ ID NO: 86), PD22 (SEQ ID NO: 87) (Plate 4); and PD23 (SEQ ID NO: 88), PD24 (SEQ ID NO: 89) (Plate 5).

[000477] Figure 16 is a graph depicting the binding to hPD-1 by anti-PD-1 antibody A1.5 and various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody A1.5, the cleavable moiety referred to herein as 3001, which includes the sequence AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318), and the masking moieties referred to herein as PD02, PD12, and PD16.

[000478] Figure 17 is a series of graphs depicting the binding to hPD-1 by the anti-PD-1 antibody A1.5 and by various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody A1.5, the cleavable moiety referred to herein as 2001, which includes the sequence ISSGLLSGRSDNH (SEQ ID NO: 214), and the masking moieties referred to herein as PD05 (SEQ ID NO: 70), PD06 (SEQ ID NO: 71), PD09 (SEQ ID NO: 74) (upper left panel); PD12 (SEQ ID NO: 77), PD23 (SEQ ID NO: 88), PD25 (SEQ ID NO: 90) (upper right panel); PD26 (SEQ ID NO: 91), PD27 (SEQ ID NO: 92), PD28 (SEQ ID NO: 93) (lower left panel); and PD30 (SEQ ID NO: 95), PD33 (SEQ ID NO: 98), PD35 (SEQ ID NO: 100) (lower right panel).

[000479] Figure 18 is a series of graphs depicting the binding to hPD-1 by the anti-PD-1 antibody A1.5 and by various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody A1.5 and the masking moieties referred to herein as PD34 (SEQ ID NO: 99), PD06 (SEQ ID NO: 71), PD09 (SEQ ID NO: 74) (upper left panel); PD34 (SEQ ID NO: 99), PD12 (SEQ ID NO: 77), PD17 (SEQ ID NO: 82) (upper right panel); PD34 (SEQ ID NO: 99), PD19

(SEQ ID NO: 84), PD25 (SEQ ID NO: 90) (lower left panel); and PD34 (SEQ ID NO: 99), PD26 (SEQ ID NO: 91), PD28 (SEQ ID NO: 93) (lower right panel).

[000480] Figure 19 is a graph depicting the ability of anti-PD1 A1.5 activatable antibodies of the disclosure to increase CMV-stimulated IFN-gamma secretion as compared to control hIgG4, but with decreased potency relative to anti-PD1 A1.5 parental antibody. The tested activatable antibodies include the anti-PD-1 antibody A1.5, the cleavable moiety referred to herein as 2001, and the masking moieties referred to herein as PD06, PD05, PD11, PD12, PD14, and PD19. In Figure 19, PMBCs were treated as follows: PBMCs: CMV+ HemaCare Donor C were plated at 2×10^5 cells/well; stimulated with 5 μ g/mL CMV lysate +/- anti-PD-1 or isotype control; IFN- γ ELISA assay performed on day 4 supernatant.

[000481] Figures 20A and 20B are a series of graphs depicting the ability of the anti-PD-1 antibody referred to herein as J43v2 (Heavy Chain (HC): SEQ ID NO: 546; Light Chain (LC): SEQ ID NO: 543) to block murine PD-L1 (mPD-L1) and murine PD-L2 (mPD-L2) from binding to cells expressing murine PD-1 (mPD-1) with single digit nM EC₅₀ values.

[000482] Figure 21 is a graph depicting the dose-dependent induction of diabetes in NOD mice by the antibody referred to herein as J43v2 m2a (HC: SEQ ID NO: 546; LC: SEQ ID NO: 543).

[000483] Figure 22 is a series of graphs depicting the binding to hPD-1 by anti-PD-1 antibody J43v2 m2a and by various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody J43v2 m2a and the masking moieties referred to herein as MP01 (SEQ ID NO: 384), MP02 (SEQ ID NO: 385), MP03 (SEQ ID NO: 386), MP04 (SEQ ID NO: 387), MP05 (SEQ ID NO: 388) (upper left panel); MP06 (SEQ ID NO: 389), MP07 (SEQ ID NO: 390), MP08 (SEQ ID NO: 391), MP09 (SEQ ID NO: 392), MP10, also referred to herein as MP010 (SEQ ID NO: 393) (upper right panel); MP11, also referred to herein as MP011 (SEQ ID NO: 394), MP12, also referred to herein as MP012 (SEQ ID NO: 395), MP13, also referred to herein as MP013 (SEQ ID NO: 396), MP14, also referred to herein as MP014 (SEQ ID NO: 397) (bottom panel).

[000484] Figure 23 is a graph depicting the binding to hPD-1 by the anti-PD-1 antibody J43v2 m2a and by various activatable anti-PD-1 antibodies that include the anti-PD-1 antibody J43v2 m2a, the cleavable moiety referred to herein as 2001, and the masking moieties referred to herein as MP5-2 (SEQ ID NO: 565), MP7-1 (SEQ ID NO: 558), MP7-5 (SEQ ID NO: 562),

MP8-2 (SEQ ID NO: 549) (upper panel); and MP7-1 (SEQ ID NO: 558), MP8-9 (SEQ ID NO: 556), MP8-8 (SEQ ID NO: 555) (lower panel).

[000485] Figure 24 is a graph depicting that the anti-PD-1 J43v2 antibody induced diabetes in NOD mice with increased frequency as dosage increased and that anti-PD-1 J43v2 activatable antibodies exhibited reduced diabetes compared to antibodies at similar doses.

[000486] Figures 25A and 25B are a series of graphs depicting the binding to hPD-1 by the anti-PD-1 antibody A1.5 and by of various activatable antibodies of the disclosure comprising the anti-PD1 antibody A1.5, the masking moiety referred to herein as PD34 (SEQ ID NO: 99), and for Figure 25A, the substrates referred to herein as 2006 (SEQ ID NO: 1095), 2007 (SEQ ID NO: 1096), 2008 (SEQ ID NO: 1097), 2009 (SEQ ID NO: 1098), and 2001 (SEQ ID NO: 214), and for Figure 25B, the substrates referred to herein as 2001 (SEQ ID NO: 214), 2008 (SEQ ID NO: 1097), 2012 (SEQ ID NO: 1101), 2011 (SEQ ID NO: 1100), and 2003 (SEQ ID NO: 1092).

[000487] Figures 26A, 26B, and 26C are a series of graphs depicting the binding to hPD-1 by the anti-PD-1 antibody nivolumab (NV1) and by of various activatable antibodies of the disclosure comprising the anti-PD1 antibody NV1, the substrate referred to herein as 2001 (SEQ ID NO: 214), and the masking moieties referred to herein NV01 (SEQ ID NO: 1206), NV02 (SEQ ID NO: 1207), NV03 (SEQ ID NO: 1208), NV04 (SEQ ID NO: 1209), NV05 (SEQ ID NO: 1210), NV06 (SEQ ID NO: 1211), NV07 (SEQ ID NO: 1212), NV08 (SEQ ID NO: 1213), NV09 (SEQ ID NO: 1214), NV10 (SEQ ID NO: 1215), NV11 (SEQ ID NO: 1216), and NV12 (SEQ ID NO: 1217).

[000488] Figures 27A, 27B, and 27C are a series of graphs depicting the binding to hPD-1 by the anti-PD-1 antibody pembrolizumab (PM1) and by various activatable antibodies of the disclosure comprising the anti-PD1 antibody PM1, the substrate referred to herein as 2001 (SEQ ID NO: 214), and the masking moieties referred to herein PM01 (SEQ ID NO: 1351), PM02 (SEQ ID NO: 1352), PM03 (SEQ ID NO: 1353), PM04 (SEQ ID NO: 1354), PM05 (SEQ ID NO: 1355), PM06 (SEQ ID NO: 1356), PM07 (SEQ ID NO: 1357), PM08 (SEQ ID NO: 1358), PM09 (SEQ ID NO: 1359), PM10 (SEQ ID NO: 1360), PM11 (SEQ ID NO: 1361), and PM12 (SEQ ID NO: 1362).

[000489] Figure 28 is a graph depicting the masking moiety specificity for various activatable antibodies of the disclosure that comprise the anti-PD-1 A1.5 (PD) antibody, the NV1 (NV) antibody, and the PM1 (PM) antibody and a variety of mask and substrate combinations.

[000490] Figures 29A and 29B are a series of graphs depicting the binding to PD1 by the IgG2a effector negative (EN) anti-PD1 antibody J43 (J43 m2a EN) and by various activatable antibodies of the disclosure comprising the IgG2a effector negative (EN) anti-PD1 antibody J43 (J43 m2a EN) and a variety of mask and substrate combinations.

[000491] Figure 30 is a graph depicting that the anti-PD-1 J43m2a EN antibody induced diabetes in NOD mice with increased frequency as dosage increased and that anti-PD-1 J43m2a EN activatable antibodies exhibited reduced diabetes compared to antibodies at similar doses.

[000492] Figures 31A and 31B are a series of graphs depicting that anti-PD1 activatable antibodies of the disclosure MP7-1 2001 m2a EN and MP8-2 2001 m2a EN inhibit the growth of MC38 syngeneic tumors in a manner similar to positive control anti-PD1 antibody J43 m2a EN both as single agents (Figure 31A) and in combination with the commercially available anti-CTLA4 antibody 9D9 mIgG2b (Figure 31B).

[000493] Figures 32A, 32B, 32C, 32D, and 32E are a series of graphs depicting that the anti-human PD1 antibody referred to herein as A1.5 Ab (i.e., VH of SEQ ID NO: 21 and VL of SEQ ID NO: 47) blocks PD-L1/PD-L2 binding to PD1 and potently activates T cells in an antigen recall assay. Figure 31A shows the binding of the anti-human PD-1 A1.5 Ab to immobilized human PD1 as detected by standard plate ELISA, and Figure 31B shows the binding of A1.5 Ab to cynomolgus PD1 as detected by ELISA. Figure 32C depicts the inhibition of biotinylated human PD-L1 to immobilized PD1 by A1.5 Ab as determined by ELISA, and Figure 32D depicts the inhibition of biotinylated human PD-L2 to immobilized PD1 as determined by ELISA. Figure 32E demonstrates that the A1.5 Ab enhances IFN- γ production in a CMV T cell restimulation assay.

[000494] Figures 33A and 33B are a series of graphs depicting that the activatable anti-PD1 antibody referred to herein as A1.5 PD34 2001 (i.e., VH of SEQ ID NO: 21, VL of SEQ ID NO: 47, masking moiety of SEQ ID NO: 99, and cleavable moiety of SEQ ID NO: 214) binds human PD1 with decreased affinity relative to the parental PD-1 Ab, i.e., A1.5 Ab (Figure 33A), and the A1.5 PD34 2001 activatable antibody shows functional masking in a CMV T cell antigen recall assay (Figure 33B).

[000495] Figure 34 is a graph and a table depicting the results of pharmacokinetic (PK) analysis of plasma samples from cynomolgus monkeys after dosing with a single IV bolus dose

of either the A1.5 antibody or the activatable antibody referred to herein as A1.5 PD34 2001 at 1 mg/kg or at 5 mg/kg. Mean PK parameters shown for both A1.5 and A1.5 PD34 2001.

[000496] Figure 35 is a graph depicting that the combination of 10 mg/kg anti-PD-1 J43 antibody plus 10 mg/kg anti-CTLA-4 antibody dosed on days 0, 4 and 7 induced diabetes in 50% of NOD mice by day eleven while the same dosing schedule of activatable anti-PD-1 J43 antibodies with CTLA-4 antibody resulted in no induction of diabetes to day eighteen.

[000497] Figure 36 is a graph depicting that the anti-PD1 activatable antibodies MP8-2 2012 m2a EN and MP8-2 2011 m2a EN inhibit the growth of MC38 syngeneic tumors similar to positive control anti-PD1 antibody J43 m2a EN, both as single agents and in combination with anti-CTLA4 antibody 9D9 mIgG2b.

[000498] Figure 37 is a series of graphs depicting that combination CTLA4 antibody and J43 activatable antibody treatment protects mice against rechallenge with MC38 tumor cells.

[000499] Figure 38 is a graph depicting that an anti-PD-1 antibody administered alone or in combination with an anti-CTLA4 antibody induced diabetes in NOD mice, whereas an anti-PD-1 activatable antibody as a single agent or in combination with an anti-CTLA4 antibody resulted in no induction of diabetes to day fifteen.

[000500] Figure 39 is a graph demonstrating that anti-PD-1 activatable A1.5 and activatable nivolumab antibodies of the embodiments enhance IFN-gamma production in a CMV T cell restimulation assay as compared to a control antibody, but not to the extent effected by anti-PD-1 antibody A1.5 or nivolumab.

[000501] Figure 40 is a graph demonstrating that an anti-PD-1 activatable pembrolizumab antibody of the embodiments enhances IFN-gamma production in a CMV T cell restimulation assay as compared to a control antibody, but not to the extent effected by anti-PD-1 antibody A1.5 or pembrolizumab.

Detailed Description of the Invention

[000502] The present invention provides monoclonal antibodies (mAbs), activatable antibodies, and antigen-binding fragments thereof that specifically bind programmed cell death protein 1 (PD-1), also known as CD279. The use of the term “PD-1” is intended to cover any variation thereof, such as, by way of non-limiting example, PD1 and/or PD 1, all variations are used herein interchangeably. Aberrant expression and/or activity of PD-1 and PD-1-related signaling has been implicated in the pathogenesis of many diseases and disorders, such as cancer.

[000503] PD-1, a cell surface receptor that belongs to the immunoglobulin superfamily, is expressed on T cells and pro-B cells. PD-1 is known to bind two ligands, PD-L1 and PD-L2. PD-1 is expressed on the surface of activated T cells, and the interaction between PD-1 and PD-L1 and/or PD-L2 functions as an immune checkpoint, as the binding of PD-L1 or PD-L2 to PD-1 inactivates the T cell. Thus, PD-1 plays a role in down-regulating the immune system by preventing the activation of T-cells, which, in turn, reduces autoimmunity and promotes self-tolerance.

[000504] The anti-PD-1 monoclonal antibodies and activated activatable anti-PD-1 antibodies of the disclosure bind and neutralize or otherwise inhibit the ability of PD-1 to bind or otherwise interact with PD-L1 and/or PD-L2. The anti-PD-1 monoclonal antibodies and activated activatable anti-PD-1 antibodies of the disclosure bind and neutralize or otherwise inhibit at least one biological activity of PD-1. For example, the anti-PD-1 monoclonal antibodies and activated activatable anti-PD-1 antibodies of the disclosure bind PD-1 and block or otherwise inhibit ligand activation of PD-1 on activated T cells. In contrast to traditional chemotherapies and other targeted anti-cancer therapies, which exert their effects by direct cytotoxic or tumor growth inhibition, the monoclonal antibodies and activatable antibodies of the disclosure block a negative regulator of T-cell activation and response, thereby allowing the immune system to attack the tumor.

[000505] The activatable anti-PD-1 antibodies of the disclosure include an antibody or antigen-binding fragment thereof that specifically binds PD-1 coupled to a masking moiety (MM), such that coupling of the MM reduces the ability of the antibody or antigen-binding fragment thereof to bind PD-1. The MM is coupled via a cleavable moiety (CM) that includes a sequence that functions as a substrate for a protease.

[000506] The activatable anti-PD-1 antibodies of the disclosure are activated when the cleavable moiety is cleaved by a protease. For example, the protease is produced by a tumor that is in proximity to T cells that express PD-1. In some embodiments, the protease is produced by a tumor that is co-localized with T cells that express PD-1. In the activated, i.e., cleaved state, the activatable anti-PD-1 antibodies of the disclosure bind PD-1 expressed on a T cell surface.

[000507] The activatable anti-PD-1 antibodies are used in methods of treating, preventing, delaying the progression of, ameliorating and/or alleviating a symptom of a disease or disorder associated with binding of a ligand selected from the group consisting of PD-L1 and PD-L2 to

PD-1 on a T cell. Such ligands are often found on tumor cells and other immune cells. Binding of such ligands to PD-1 can lead to immune suppression, such as suppression of T cell activity, which is reduced by anti-PD1 antibodies and activatable anti-PD-1 antibodies of the disclosure. For example, the activatable anti-PD-1 antibodies are used in methods of treating, preventing, delaying the progression of, ameliorating and/or alleviating a symptom of a cancer or other neoplastic condition.

[000508] Exemplary activatable anti-PD-1 antibodies of the invention include, for example, activatable antibodies that include a heavy chain and a light chain that are, or are derived from, antibodies described in the Examples, for example in Examples 1, 2, 8, 9, 14, and 15.

[000509] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 1, 5, 9, 13, 17, 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 3, 7, 11, 15, 19, 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, 59, 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539, 541, 572, 574, 576, 578, 580, 582, 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, 997, 999, 1001, 1003, 1005, 1007, 1009, 1011, 1013, 1015, 1017, 1019, 1021, 1023, 1025, 1027, 1030, 1032, 1034, 1036, 1039, 1041-1090, 1113-1120, 1123, 1124, 1127, 1128, 1131, 1132, 1134, 1135, 1138, 1139, 1143, 1144, 1147, 1148, 1151, 1152, 1155, 1156, 1159, 1160, 1163, 1164, 1167, 1168, 1170, 1171, 1174, 1175, 1178, 1179, 1182, 1183, 1186, 1187, 1190, 1191, 1194, 1195, 1198, 1199, 1202, 1203, 2055, 2054, 2057, 2056, 2059, and 2058.

[000510] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, 59, 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539,

541, 572, 574, 576, 578, 580, 582, 584, 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, 997, 999, 1001, 1003, 1005, 1007, 1009, 1011, 1013, 1015, 1017, 1019, 1021, 1023, 1025, 1027, 1029, 1030, 1032, 1034, 1036, 1039, 1041-1090, 1113-1120, 1123, 1127, 1131, 1134, 1138, 1144, 1148, 1152, 1156, 1160, 1164, 1168, 1170, 1174, 1178, 1182, 1186, 1190, 1194, 1198, 1203, 2055, 2054, 2057, 2056, 2059, and 2058.

[000511] In some embodiments, the light chain comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 39, 41, 43, 45, 47, 49, 51, 53, 55, 57, and 59. In some embodiments, the light chain comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 584, 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, and 997.

[000512] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 243, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 572, 574, 576, 578, 580, 582, 1029, 1120, 1124, 1128, 1132, 1135, 1139, 1143, 1147, 1151, 1155, 1159, 1163, 1167, 1171, 1175, 1179, 1183, 1187, 1191, 1195, 1199, 1203, 2055, 2054, 2057, 2056, 2059, and 2058.

[000513] In some embodiments, the light chain comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539, 541, 572, 574, 576, 578, 580, 582, 999, 1001, 1003, 1005, 1007, 1009, 1011, 1013, 1015, 1017, 1019, 1021, 1023, 1025, 1027, 1029, 1030, 1032, 1034, 1036, 1039, 1041-1090, 1113-1120, 1123, 1127, 1131, 1134, 1138, 1144, 1148, 1152, 1156, 1160, 1164, 1168, 1170, 1174, 1178, 1182, 1186, 1190, 1194, 1198, 1203, 2055, 2054, 2057, 2056, 2059, and 2058.

[000514] In some embodiments, the activatable anti-PD-1 antibody includes a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of

SEQ ID NOs: 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 243, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539, 541, 572, 574, 576, 578, 580, 582, 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, 997, 1027, 1028, 1029, 1030, 1032, 1034, 1036, 1039, 1041-1076, 1113-1120, 1123, 1124, 1127, 1128, 1131, 1132, 1134, 1135, 1138, 1139, 1143, 1144, 1147, 1148, 1151, 1152, 1155, 1156, 1159, 1160, 1163, 1164, 1167, 1168, 1170, 1171, 1174, 1175, 1178, 1179, 1182, 1183, 1186, 1187, 1190, 1191, 1194, 1195, 1198, 1199, 1202, 12032055, 2054, 2057, 2056, 2059, and 2058.

[000515] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 21, 23, 25, 27, 29, 31, 33, 35, and 37, and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOs: 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 243, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, 997, 1028, 1029, 1041-1076, 1138, 1139, 1143, 1144, 1147, 1148, 1151, 1152, 1155, 1156, 1159, 1160, 1163, 1164, 1167, 1168, 1170, 1171, 1174, 1175, 1178, 1179, 1182, 1183, 1186, 1187, 1190, 1191, 1194, 1195, 1198, 1199, 1202, 1203, 2055, 2054, 2057, 2056, 2059, and 2058.

[000516] In some embodiments, the light chain comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOs: 919, 921, 923, 925, 927, 929, 931, 933, 935, 937, 939, 941, 943, 945, 947, 949, 951, 953, 955, 957, 959, 961, 963, 965, 967, 969, 971, 973, 975, 977, 979, 981, 983, 985, 987, 989, 991, 993, 995, 997, 1028, 1041-1076, 1138, 1144, 1148, 1152, 1156, 1160, 1164, 1168, 1170, 1174, 1178, 1182, 1186, 1190, 1194, 1198, 1202, 2055, 2057, and 2059.

[000517] In some embodiments, the light chain comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOs: 215, 217, 219, 221, 223, 225, 227, 229, 231, 233, 235, 237, 239, 241, 243, 245, 247, 249, 251, 253, 255, 257, 259, 261, 263, 265, 267, 269, 271, 273, 275, 277, 279, 281, 283, 285, 287, 289, 291, 293, 1029, 1139, 1143, 1147,

1151, 1155, 1159, 1163, 1167, 1171, 1175, 1179, 1183, 1187, 1191, 1195, 1199, 1203, 2054, 2056, and 2058.

[000518] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NO: 546 and a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOS: 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539, 541, 572, 574, 576, 578, 580, 582, 999, 1001, 1003, 1005, 1007, 1009, 1011, 1013, 1015, 1017, 1019, 1021, 1023, 1025, 1027, 1030, 1032, 1034, 1036, 1039, 1077-1090, 1113-1120, 1123, 1124, 1127, 1128, 1131, 1132, 1134, and 1135.

[000519] In some embodiments, the activatable anti-PD-1 antibody includes a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOS: 999, 1001, 1003, 1005, 1007, 1009, 1011, 1013, 1015, 1017, 1019, 1021, 1023, 1025, 1027, 1030, 1032, 1034, 1036, 1039, 1070-1090, 1119, 1123, 1127, 1131, and 1134.

[000520] In some embodiments, the activatable anti-PD-1 antibody includes a light chain that comprises or is derived from an amino acid sequence selected from the group consisting of SEQ ID NOS: 515, 517, 519, 521, 523, 525, 527, 529, 531, 533, 535, 537, 539, 541, 572, 574, 576, 578, 580, 582, 1120, 1124, 1128, 1132, and 1135.

[000521] In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from a heavy chain amino acid sequence shown in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a light chain that comprises or is derived from a light chain amino acid sequence shown in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a heavy chain that comprises or is derived from a heavy chain amino acid sequence shown in Table 7, and a light chain that comprises or is derived from a light chain amino acid sequence shown in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group A in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group B in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group C in Table 7. In some embodiments, the

activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group D in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group E in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group F in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group H in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group I in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group J in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group K in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group L in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group M in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain variable region sequence and the light chain variable region sequence as shown in Group N in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain variable region sequence and the light chain variable region sequence as shown in Group O in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group P in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of a heavy chain variable region sequence and a light chain variable region sequence from the combinations shown in Group Q in Table 7.

[000522] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the complementarity determining region (CDR) sequences of a heavy chain sequence from the heavy chain sequences shown in Group A in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group A in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group A in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group A in Table 7.

[000523] In some embodiments, the activatable anti-PD-1 antibody includes a combination of CDRs of a heavy chain sequence from the heavy chain sequences shown in Group B in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group B in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group B in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group B in Table 7.

[000524] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group C in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group C in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group C in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group C in Table 7.

[000525] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group D in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group D in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group D in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group D in Table 7.

[000526] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group E in

Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group E in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group E in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group E in Table 7.

[000527] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group F in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group F in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group F in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group F in Table 7.

[000528] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group G in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group G in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group G in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group G in Table 7.

[000529] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group H in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group H in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group H in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group H in Table 7.

[000530] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group I in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group I in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a

heavy chain sequence from the heavy chain sequences shown in Group I in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group I in Table 7.

[000531] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group J in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group J in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group J in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group J in Table 7.

[000532] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group K in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group K in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group K in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group K in Table 7.

[000533] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group L in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group L in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group L in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group L in Table 7.

[000534] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group M in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group M in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group M in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group M in Table 7.

[000535] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group N in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group N in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group N in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group N in Table 7.

[000536] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group O in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group O in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group O in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group O in Table 7.

[000537] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group P in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group P in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group P in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group P in Table 7.

[000538] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group Q in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group Q in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group Q in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group Q in Table 7.

[000539] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group R in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the

CDRs of a light chain sequence from the light chain sequences shown in Group R in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group R in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group R in Table 7.

[000540] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group S in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group S in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group S in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group S in Table 7.

[000541] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group T in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group T in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group T in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group T in Table 7.

[000542] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group U in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group U in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group U in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group U in Table 7.

[000543] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a

heavy chain sequence from the heavy chain sequences shown in Group V in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group V in Table 7.

[000544] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 619 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1856 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 619 and the light chain sequence of SEQ ID NO: 1856 as shown in Group V in Table 7.

[000545] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1846 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1858 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1846 and the light chain sequence of SEQ ID NO: 1858 as shown in Group V in Table 7.

[000546] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1843 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1859 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1843 and the light chain sequence of SEQ ID NO: 1859 as shown in Group V in Table 7.

[000547] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1847 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1847 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000548] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1848 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1860 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain

sequence of SEQ ID NO: 1848 and the light chain sequence of SEQ ID NO: 1860 as shown in Group V in Table 7.

[000549] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1844 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1861 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1844 and the light chain sequence of SEQ ID NO: 1861 as shown in Group V in Table 7.

[000550] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1841 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1841 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000551] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1842 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1842 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000552] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1845 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1845 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000553] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1835 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1857 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain

sequence of SEQ ID NO: 1835 and the light chain sequence of SEQ ID NO: 1857 as shown in Group V in Table 7.

[000554] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1836 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1862 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1836 and the light chain sequence of SEQ ID NO: 1862 as shown in Group V in Table 7.

[000555] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1837 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1863 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1837 and the light chain sequence of SEQ ID NO: 1863 as shown in Group V in Table 7.

[000556] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1838 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1864 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1838 and the light chain sequence of SEQ ID NO: 1864 as shown in Group V in Table 7.

[000557] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1838 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1865 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1838 and the light chain sequence of SEQ ID NO: 1865 as shown in Group V in Table 7.

[000558] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1839 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1864 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain

sequence of SEQ ID NO: 1839 and the light chain sequence of SEQ ID NO: 1864 as shown in Group V in Table 7.

[000559] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1839 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1865 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1839 and the light chain sequence of SEQ ID NO: 1865 as shown in Group V in Table 7.

[000560] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1840 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1866 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1840 and the light chain sequence of SEQ ID NO: 1866 as shown in Group V in Table 7.

[000561] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1841 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1867 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1841 and the light chain sequence of SEQ ID NO: 1867 as shown in Group V in Table 7.

[000562] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1841 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1868 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1841 and the light chain sequence of SEQ ID NO: 1868 as shown in Group V in Table 7.

[000563] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1849 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1858 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain

sequence of SEQ ID NO: 1849 and the light chain sequence of SEQ ID NO: 1858 as shown in Group V in Table 7.

[000564] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1853 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1859 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1853 and the light chain sequence of SEQ ID NO: 1859 as shown in Group V in Table 7.

[000565] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1869 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1869 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000566] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1850 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1860 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1850 and the light chain sequence of SEQ ID NO: 1860 as shown in Group V in Table 7.

[000567] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1851 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 1861 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1851 and the light chain sequence of SEQ ID NO: 1861 as shown in Group V in Table 7.

[000568] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1852 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain

sequence of SEQ ID NO: 1852 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000569] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1854 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1854 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000570] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of the heavy chain sequence of SEQ ID NO: 1855 and a combination of the CDRs of the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the heavy chain sequence of SEQ ID NO: 1855 and the light chain sequence of SEQ ID NO: 737 as shown in Group V in Table 7.

[000571] In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group W in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a light chain sequence from the light chain sequences shown in Group W in Table 7. In some embodiments, the activatable anti-PD-1 antibody includes a combination of the CDRs of a heavy chain sequence from the heavy chain sequences shown in Group W in Table 7 and the CDRs of a light chain sequence from the heavy chain sequences shown in Group W in Table 7.

Table 7. Variable Heavy Chain Region (VH) and Variable Light Chain Region (VL) Sequences for Activatable Antibodies that Bind PD-1

Group A	
VH	QVQLVESGGDVVQPGGSLRLSCAASGVAFSNGMHVVRQAPGKGLEWVAVIWDGSKNYYADSVKGRFTISR DNSKNMPLYLQMNSLRAEDTAMYYCARNDYWGQGTLLTVSS (SEQ ID NO: 615)
VH	QVQLVESGGDVVQPGSLRLSCAASGLTFTNYGFHWVRQAPGKGLEWVAVIWDGSKKYYADSVKGRFTISR DNSKNTLYLQMNNLRAEDTAVYYCATGDDYWGQGTLLTVSS (SEQ ID NO: 617)
VH	QVYLVESGGGVVQPGSLRLSCAASGFTFSNGMHVVRQAPGKGLEWVALIWDGSKNYYADSVKGRFTISR DNSKNTLYLQMTSLRVEDTAVYYCASNVDPHWGQGTLLTVSS (SEQ ID NO: 618)
VH	QVQLVESGGGVVQPGSLRLDCKASGITFSNSGMHWVRQAPGKGLEWVAVIWDGSKRYADSVKGRFTISR DNSKNTLFLQMNSLRAEDTAVYYCATNDYWGQGTLLTVSS (SEQ ID NO: 1346)
VH	QLQLQESGPGLVKPSSETLSLTCTVSGGSLRSSTFFWGWIRQPPGKGLEWIGSIYYSGSTYYNPSLKSRVTIS VDTSKNQFSLKLSSVTAADTAVYYCVRDYDILTGDDEYWGQGTLLTVSS (SEQ ID NO: 620)
VH	QVQLVESGGGVVQPGSLRLSCTTSGITFSSYGFHWVRQAPGKGLEWVAVIWDGSKKYYADSVKGRFTLSR DDSKNTLYLQMNSLRAEDTAVYYCVTGDDYWGQGTLLTVSS (SEQ ID NO: 621)
VH	QLQLQESGPGLVKPSSETLSLTCSVSGGSLRSSTFFWGWIRQPPGKGLEWIASIFYSGETYFNPSLKSRVTIS VDTSRNQFSLKLSSVTAADTAVYYCARDYDILTGDDEYWGQGTLLTVSS (SEQ ID NO: 623)
VL	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLIIYDASNRATGIPARFSGSGSGTDFT LTISSELPEDFAVYYCQQRSNWPLTFGGGTKVEIK (SEQ ID NO: 624)
VL	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLIIYDTSNRATGIPARFSGSGSGTDFT LTISSELPEDFAVYYCQQRSNWPLTFGGGTKVEIK (SEQ ID NO: 625)
VL	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLIIYDASNRATGIPARFSGSGSGTDFT LTISSELPEDFAVYYCQQSSNWPRTFGQGTKVEIK (SEQ ID NO: 626)
VL	DIQMTQSPSSLSASVGDVRSITCRASQGISSWLAWYQQKPEKAPKSLIYAASNLRSVPSRFSGSGSGTDFT LTISSLQPEDFATYYCQQYYSPRTFGQGTKVEIK (SEQ ID NO: 628)
VL	EIVLTQSPATLSLSPGERATLSCRASQSVSSYLAWYQQKPGQAPRLIIYDASNRATGIPARFSGSGSGTDFT LTISSELPEDFAVYYCQQRSNWPLTFGGGTKVEIK (SEQ ID NO: 629)
VL	DIQMTQSPSSLSASVGDRTITCRASQGISSWLAWYQQKPEKAPKSLIYAASSLQSGVPSRFSGSGSGTDFT LTISSLQPEDFATYYCQQYYSPRTFGQGTKVEIK (SEQ ID NO: 630)
Group B	
VH	QVQLQQPGAELVKPGASVKLSCKASGYTFTSYLYWMKQRPQGQLEWIGGVNPSNGGTNFSEKFKSKATLTV DKSSSTAYMQLSLTSSEDSAVYYCTRRDSNYDGGFDYWGQGTLLTVSSAK (SEQ ID NO: 631)
VH	QVQLQQPGAELVKPGTSVKLSCKASGYTFTNYYMYWVKQRPQGQLEWIGGINPSNGGTNFNEKFKNKATLTV DSSSSTTYMQLSLTSSEDSAVYYCTRRDYRFDMGFDYWGQGTLLTVSSAK (SEQ ID NO: 632)

VH	MDWTWSILFLVAAPTGAHSQVQLVQSGVEVKKPGASVKVSKASGYFTFTNYYMYWVRQAPQGQLEWMGGINP SNGGTNFNEKFKNRVTLTDSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSS (SEQ ID NO: 633)
VH	QVQLVQSGVEVKKPGASVKVSKASGYFTFTNYYMYWVRQAPQGQLEWMGGINPSNGGTNFNEKFKNRVTLT DSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSS (SEQ ID NO: 634)
VH	EVQLVLSGGGFVQPGGSLKLSCAASGFTFSSYAMSWVRQNPERRLVWVATITGGGRNTYYPDSVKGRFTISR DNAKNTLYLQMSSLRSEDAMYYCTRQGYDGYTWFAYWGQGTTLTVS (627)
VL	DIVLTQSPSTSLAVSLGQRATISCRASKSVSTSGFSYLHWYQQKPGQPPKLLIFLASNLESGVPARFSGSGS TDFTLNIHPVEEEDAATYYCQHSWELPLTFGAGTKLELK (SEQ ID NO: 635)
VL	DIVLTQSPASLAVSLGQRAAISCRASKGVSTSGYSYLHWYQQKPGQSPKLLIYLASYLESGVPARFSGSGS TDFTLNIHPVEEEDAATYYCQHSRDLPLTFGTGKLELK (SEQ ID NO: 636)
VL	MAPVQQLGLLVFLPAMRCEIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIY LASYLESGVPARFSGSGSGTDFTLTISSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 637)
VL	EIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLASYLESGVPARFSGSGS TDFTLTISSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 638)
VL	MAPVQQLGLLVFLPAMRCEIVLTQSPSLPVTGPGEPAISCRASKGVSTSGYSYLHWYQQKPGQSPQLLIY LASYLESGVPDRFSGSGSGTDFTLKISRVEAEDVGYYCQHSRDLPLTFGQGTKLEIK (SEQ ID NO: 639)
VL	EIVLTQSPSLPVTGPGEPAISCRASKGVSTSGYSYLHWYQQKPGQSPQLLIYLASYLESGVPDRFSGSGS TDFTLKISRVEAEDVGYYCQHSRDLPLTFGQGTKLEIK (SEQ ID NO: 640)
VL	MAPVQQLGLLVFLPAMRCDIVMTQTPLSLPVTGPGEPAISCRASKGVSTSGYSYLHWYQQKPGQSPQLLIY LASYLESGVPDRFSGSGSGTAFTLKISRVEAEDVGLYYCQHSRDLPLTFGQGTKLEIK (SEQ ID NO: 641)
VL	DIVMTQTPLSLPVTGPGEPAISCRASKGVSTSGYSYLHWYQQKPGQSPQLLIYLASYLESGVPDRFSGSGS TAFTLKISRVEAEDVGLYYCQHSRDLPLTFGQGTKLEIK (SEQ ID NO: 642)
VL	DIVLTQSPSTSLAVSLGQRATISCRASEVDNSGISFMNWFQKPGQPPKLLIYAASNPGSGVPARFSGSGS TDFSLNIHPMEEDDTAMYFCQQSKEVPWTFGGGTELEIKR (SEQ ID NO: 725)
HC	MAVLGLLFLCLVTFPSCVLSQVQLVQSGVEVKKPGASVKVSKASGYFTFTNYYMYWVRQAPQGQLEWMGGINP SNGGTNFNEKFKNRVTLTDSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSSASTKG PSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLG TKTYTCNVDHKPSNTKVDKRVESKYGPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQE DPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVTLHQLDNLNGKEYKCKVSNKGLPSSIEKTISKAKG QPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTPPVLDSDGSFFLYSRLTVD KSRWQEGNVFSCSVMEALHNHYTQKSLSLSLGK (SEQ ID NO: 643)

HC	QVQLVQSGVEVKKPGASVKVSCASGYTFTNYYMYWVRQAPGQGLEWMGGINPSNGGTNFNEKFKNRVTLT DSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSSASTKGPSVFPLAPCSRSTSESTAA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTKYTCNVDPKPSNTKVDK RVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVDSQEDPEVQFNWYVDGVEVHNAK TKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTK NQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEAL HNHYTQKSLSLSPGK (SEQ ID NO: 644)
HC	MAVLGLLFLCLVTFPSCVLSQVQLVQSGVEVKKPGASVKVSCASGYTFTNYYMYWVRQAPGQGLEWMGGINP SNGGTNFNEKFKNRVTLTDSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSSASTKG PSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLG TQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDV SHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISK AKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKL TVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPGK (SEQ ID NO: 645)
HC	QVQLVQSGVEVKKPGASVKVSCASGYTFTNYYMYWVRQAPGQGLEWMGGINPSNGGTNFNEKFKNRVTLT DSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFDYWGQGTTVTVSSASTKGPSVFPLAPSSKSTSGGTA LGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGTQTYICNVNHKPSNTKVDK KVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVH NAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRDE LTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMH EALHNHYTQKSLSLSPGK (SEQ ID NO: 646)
LC	MAPVQLLGLLVFLPAMRCEIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIY LASYLESGVPARFSGSGSDFTLTITSSLEPEDFAVYYCQHSRDLPLTFGGGTKEIKRTVAAPSVFIFPPS DEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYA CEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 647)
LC	EIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLASYLESGVPARFSGSGS GSDFTLTITSSLEPEDFAVYYCQHSRDLPLTFGGGTKEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFY PREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRG EC (SEQ ID NO: 648)
LC	MAPVQLLGLLVFLPAMRCEIVLTQSPSLPVTGPGEPAISCRASKGVSTSGYSYLHWYLLQKPGQSPQLLIY LASYLESGVPDRFSGSGSDFTLTKISRVEAEDGVYYCQHSRDLPLTFGGGTKEIKRTVAAPSVFIFPPS DEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYA CEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 649)
LC	EIVLTQSPSLPVTGPGEPAISCRASKGVSTSGYSYLHWYLLQKPGQSPQLLIYLASYLESGVPDRFSGSGS GSDFTLTKISRVEAEDGVYYCQHSRDLPLTFGGGTKEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFY PREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRG EC (SEQ ID NO: 650)

LC	MAPVQLLGLLVLFPLPAMRCDIVMTQTPLSLPVTGPGEPAISCRASKGVSTSGYSYLHWYLQKPGQSPQLLIYLASYLESQVDFRFSGSGSGTAFTLKISRVEAEDVGLYYCQHSRDLPLTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 651)
LC	DIVMTQTPLSLPVTGPGEPAISCRASKGVSTSGYSYLHWYLQKPGQSPQLLIYLASYLESQVDFRFSGSGSGTAFTLKISRVEAEDVGLYYCQHSRDLPLTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 652)
Group C	
VH	DVQLQESGPGLVKPSQSLSLTCTVTGYSITSDYAWNWIQFPGNKLEWMGYINYSGSTSYNPSLKSRSITRDTSKNQFFLQLNSVTTEDTATYYCARWIGSSAWYFDVWGAGTTVT (SEQ ID NO: 726)
VL	DVLMQTPLSLPVS LGDQASISCRSGQNI VHSNGNTYLEWYLQKPGQSPKLLIYKVS NRFFGVPDRISGSGSGTDFTLKISRVEAEDLGVYFCFQGSHPFTFGSGTKLEIK (SEQ ID NO: 727)
VH	QVQLQQPGAELVKPGASVKLSCKASGYTFTTYYLYWVRQRPQGQLEWIGGINPSNGGTNFNEKFKSKATLTVDKSSSTAYMQINSLTSEDSAVYYCTRRDYRDRGFDYWGQGTSTVTV (SEQ ID NO: 728)
VL	DIVLTQSPASLAVSLGQRATISCRASKSVSTSGFNIIHWYQQKPGQPPKLLIYLASNLESQVDFRFSGSGSGTDFTLNIHPVEDEDAATYYCQHSRELPLTFGAGTKLEIK (SEQ ID NO: 729)
VH	QVQLQQSGAELVKPGASVKMSCKAFGYTFTTYPPIEWMKQNHGKSLEWIGNFHPYNDTKYNEKFKGKAKLTV EKSSSTTVYLELSRLTSDDSAVYYCARENYGSHGGFVYWGQGTSLTVTV (SEQ ID NO: 730)
VL	ENVLTQSPAIMASAPGEKVTMTCRASSSVISSYLHWYQQKSGASPKLWIYSTSNLASQVDFRFSGSGSGTSLTISSVEAEDAATYYCQQYNGYPLTFGAGTKLEIK (SEQ ID NO: 731)
Group D	
	LFTVTVPKELYIIIEHGSNVTLECNFDTGSHVNLGAITASLQKVENDTSPHRERATLLEEQLPLGKASFHIPQVQVRDEGQYQCIIYIGVAWDYKYLTLKVKASYRKINTHILKVPETDEVELTCQATGYPLAEVSWPNVSVPAN TSHSRTPEGLYQVTSVLRLKPPPGRNFSVFWNTHVRELTLASIDLQSQMEPRTHPTWEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSRDELTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSQSVMEALHNHYTQKSLSLSPGK (SEQ ID NO: 732)
Group E	
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFTSYGISWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTTDTSTNTAYMELRSLRSDDTAVYYCARDADYSSGSGYWGQGTSLTVTVSS (SEQ ID NO: 733)
VH	QVQLVQSGAEVKKPGASVRVSCKASGYTLTSYIIHWVRQAPGQGLEWMGIINPRGATISYAQKFQGRVTMTRDTSTSTVYMELRNLKSEDTALYYCATAGIYGFDYWGRTSLTVTVSS (SEQ ID NO: 734)
VH	QVQLQESGPGLVKPSQTLSTCTVSGGSISSGAYYWSWIRQHPGKGLEWIGYIYYNGNTYYNPSLRSLVTISVDASKNQFSLKLSSVTAADTAVYYCARASDYVWGGYRYMDAFDIWGRGTLITVSS (SEQ ID NO: 735)

VH	GAHSEVQLVQSGGGVVQPGRSLRLSCAASGFTFSSYWCDRMSWVRQAPGKGLEWVSAISGSGGSTYYADSVK GRFTISRDN SKNTLYLQMNSLRAEDTAVYYCAKENWGSYFDLWGQGT TTVTVSS (SEQ ID NO: 736)
VL	SYELTQPPSVSVSPGQTARITCSGDALPKQYAYWYQQKPGQAPVMVIYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDEADYYCQSADNSITYRVFGGGTKVTVL (SEQ ID NO: 737)
VL	QSALTQPASVSGSPGQSITISCTGTSNDVGGYNYVSWYQHHPGKAPKLIIYDVTNRPSGVSDRFGSGSKSGNT ASLTISGLLAEDEGDYYCSSLTYTIVTNFEVLFGGGTKLTV (SEQ ID NO: 738)
VL	QSVLTQPPSASGTPGQRTISCSGSNSNIGSNSNVNWWYQQLPGTAPKLLIYGNNQRPSGVPDRFGSGSKSGTSA SLAISGLQSENEADYYCAAWDDSLNGPVFGRGTVTLGE (SEQ ID NO: 739)
VL	GVHSDIVMTQSPSTLSASVGDRVITITCRASQGISSWLAWYQQKPGRAPKVLIYKASTLESQVPSRFSGSGSG TDFTLTITSSLPEDFATYYCQQSYSTPWTFTGQGTKLEIKR (SEQ ID NO: 740)
Group F	
VH	QVQLVQSGHEVKQPGASVKMSCKASGYSTSSWIHWVKQAPGQGLEWIGYIYPSTGFTEYNQKFKDRATLTA DKSTSTAYMELSSSLRSED TAVYYCARWRDSSGYHAMDYWGQGT LTVTVSS (SEQ ID NO: 741)
VH	QVQLVQSGHEVKQPGASVKMSCKASGYSTSSWIHWVRQAPGQGLEWIGYIYPSTGFTEYNQKFKDRATLTA DKSTSTAYMELSSSLRSED TAVYYCARWRDSSGYHAMDYWGQGT LTVTVSS (SEQ ID NO: 742)
VL	EIVLTQSPATLSLSPGQRLTISCRASQSVSTSGYSYMHWWYQQKPDQSPKLLIKFGSNLESGIPARFSGSGSG TDFTLTITSSLEPEDFATYYCQHSWEIPYTFGQGTKLEIK (SEQ ID NO: 743)
VL	DIVLTQSPATLSLSPGQRLTISCRASQSVSTSGYSYMHWWYQQKPDQSPKLLIKFGSNLESGIPARFSGSGSG TDFTLTITSSLEPEDFAVYYCQHSWEIPYTFGQGTKLEIK (SEQ ID NO: 744)
Group H	
VH	QGQLVQSGAEVKKPGASVKVSCASGYTFTDYEMHWVRQAPIHGLEWIGVIESETGGTAYNQKFKGRVTITA DKSTSTAYMELSSSLRSED TAVYYCAREGITTVATYYWYFDVWGQGT TTVTVSS (SEQ ID NO: 745)
VL	DVVMQTQSPLSLPVTLGQPASISCRSSQSIVHSNGNTYLEWYLQKPGQSPQLLIYKVSNRFGVPDRFGSGSGS GTDFTLKISRVEAEDVGVIYCFQGSVPLTFGQGTKLEIK (SEQ ID NO: 746)
Group I	
VH	MGLGLQWVFFVALLKGVHCEVRLLLESGLLVKPEGSLKLSCVASGFTFSDYFMSWVRQAPGKGLEWVAHIYT KSYNYATYYSGSVKGRFTISRDDSRSMVYLQMNLRTEDTATYYCTRDGSGYPSLDFWGQGTQVTVSSATTT APSVYPLAPACDSTTKS (SEQ ID NO: 747)
VL	YELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLT IRDVRAEDEGDYYCFSGYVDSDSLKYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDYFP GSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAECCL (SEQ ID NO: 748)

Bs	METDTLLLWVLLLWVPGSTGDAAQPARRARRTKLGTELGSPGLQEFVRLLESGGGLVKPEGSLKLSCVASG FTFSDYFMSWVRQAPGKGLEWVAHIYTKSYNYATYYSGSVKGRFTISRDDSRSMVYLQMNLRTEDTATYYC TRDGSYGPSLDFWGGQTQVTVSSGGGSDIQMTQSPSSLPASLGDRVTINCQASQDISNYLNWYQQKPGKAP KLLIYYTNKLADGVPSRFSGSGSGRDSSTFISSLESEDIGSYCQQYYNYPWTFGPGTKLEIKGGGSGGGG
Ab	SGGGGSEVQLVESGGGLVQPGKSLKLSCEASGFTFSGYGMHWVRQAPGRGLESVAYITSSSINIKYADAVKG RFTVSRDNAKNLLFLQMNILKSEDTAMYYCARFDWDKNYWGQGTMTVTVSSGGGGSYELTQPPSASVNVGETV KITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSG YVDSDSKLYVFGSGTQLTVLGPRGGPEQKLISEEDLNSAVDHHHHHH (SEQ ID NO: 749)
Group J	
VH	QLVQSGAEVKKPGASVKVSCKASGYTFTSYGISWVRQAPGQGLEWMGWISAYNGNTNYAQLQGRVTMTTDT STSTATMELRSLRSDDTAVYYCARGRGYSYGIDAFDIWGQGTMTV (SEQ ID NO: 750)
VH	LSYVLTQPPSVSVSPGQTARITCSGDALPKQYAYWYQQKPGQAPVLVIYKDSERPSGIPERFSGSSSGTTVT LTISGVQAEDEADYYCQSADSSGTYVVFGGGTKLTVLGQP (SEQ ID NO: 751)
Group K	
VH	QVQLQQSGPGLVRPSQTLNLSLSDISGDSVSSNSATWNWIRQSPSRGLEWLGRFTFYRSKWYHDYALSVKSRIT INPDTSKNQFSLQLNSVSPGDTAVYFCVREDIDGRLDYWGQGTTLTVTVSS (SEQ ID NO: 752)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKIGNPTYAQGFTGRFVFSL DTSVSTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVTVSS (SEQ ID NO: 753)
VH	MAEVQLLESGLAEVKKPGASVKVSCKASGYTFTSHYMHWVRQAPGQGLEWMGVINPSGGSTSYAQKFQGRVTM TRDTSTSTVYMDLSSSLRSEDVAVYYCARRSEAYYHGMDVWGQGTTLTVTVSS (SEQ ID NO: 754)
VH	EVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKIGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVTVSS (SEQ ID NO: 755)
VH	QVQLVESGGGLVQPGGSLRLSCEATGFTFSRYWMHWVRQAPGKGLEWVSSISSSSSYIYADSVKGRFTISR DNAKNSLYLQMNLSRAEDTAVYYCARDTLEYYGSGILENAMGYGMDVWGQGTTLTVTVSS (SEQ ID NO: 756)
VH	EVQLVESGGGLVRPGGSLRLACAASGFSFSDYYMTWIRQAPGRGLEWIAIYISDSGQTVHYADSVKGRFTISR DNTKNSLFLQVNTLRAEDTAVYYCAREDLLGYLQSWGQGTTLTVTVSS (SEQ ID NO: 757)
VH	EVQLVESGGGVVQPGSLRLSCLAASGFTFSSYWMWVRQAPGKGLEWVANIKQDGSEKYYVDSVKGRFTISR DNAKNSLYLQMNLSRAEDTAVYYCAREGEHDAFDIWGQGTMTVTVSS (SEQ ID NO: 758)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKIGNPTYAQGFTGRFVFSL DTSVSTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVTVSS (SEQ ID NO: 759)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKIGNPTYAQGFTGRFVFSL DTSVSTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVTVSS (SEQ ID NO: 760)
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTRYTMHWVRQAPQRLEWMGWINAGNGNTKYSQKFQGRVTITR DTSASTAYMELSSSLRSEDVAVYYCAKVSAGTESWFDPWGQGTTLTVTVSS (SEQ ID NO: 761)

VH	QVQLVQSGAEVKKPGASVKVSCSKASGYTFTSYGISWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTSTAYMELRSLRSDDTAVYYCARGLYGDEDYWGQGTTLTVVSS (SEQ ID NO: 762)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKIGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 763)
VH	QMQLVQSGAEVKKPGESLKI SCKSGSYFTSYWIGWVRQMPGKGLEWMGIIYPGDS DTRYSPSFQGVVTISA DKSISTAYLQWSSLKASDTAMYYCAGSVTRKRYSSSWPPFDYWGQGTTLTVVSS (SEQ ID NO: 764)
VH	QVQLQQWGAGLLKSSETLSLSCAVYGGTFRDDHWSWIRQPPGKGLEWIGESHHTGRTIYNPSLRSRVTMSID TSKNEFSLILRSVTAADTATYFCARGNNYVWGNQEDFWGQGTTLTVVSS (SEQ ID NO: 765)
VH	QVQLQQSGPGLVRPSQTLSLSCDISGDSVSSNSATWNWIRQSPSRGLEWLGRTFYRSKWYHDYALSVKSRIT INPDTSKNQFSLQLNSVSPGDTAVYFCVREDIDGRLDYWGQGTTLTVVSS (SEQ ID NO: 766)
VH	EVQLVESGGALVQPGGSLRLSCAVSGFTFSDHYMDWVRQAPGKGLEWVARSRNKGNSYTTEYAASVVRGRTI SRDDSKNSLYLQMNLSLKTEDTAVYYCVRVGVVPALDGMVWGQGTTLTVVSS (SEQ ID NO: 767)
VH	EVQLVESGGGLVQPGGSLRLSCAASGFTFSSYWMHWVRQAPGKGLEWVSSISSSSSYIYADSVKGRFTISR DNAKNSLYLQMNLSLRAEDTAVYYCARDTLEYYGSGILENAMGYYGMDVWGQGTTLTVVSS (SEQ ID NO: 768)
VH	EVQLLESGGGVVQTGRSLRLSCSDSGSTFRSQAMHWVRQTPGKGLEWLAVTSHDGSKTYADSVKGRFTISR DNSKNTLYLQMNLSLRGEDTAVYYCARGGRGYTYDHSFFDYWGQGTTLTVVSS (SEQ ID NO: 769)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKDGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 770)
VH	QVQLVQSGSELKKPGASVKISCKASGYIFSDNGVNWVRQAPGQGLEWMGWINTKTGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 771)
VH	QVQLVQSGSELKKPGASVKISCKASGYTFS DNGVNWVRQAPGQGLEWMGWINTKDGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 772)
VH	QVQLVQSGSELKKPGASVKISCKASGYTFS DNGVNWVRQAPGQGLEWMGWINTKTGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVS (SEQ ID NO: 773)
VH	QVQLVQSGSELKKPGASVKISCKASGYKFS DNGVNWVRQAPGQGLEWMGWINTKSGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 774)
VH	QVQLVQSGSELKKPGASVKISCKASGYTFS DNGVNWVRQAPGQGLEWMGWINTKSGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 775)
VH	QVQLVQSGSELKKPGASVKISCKASGYKFS DNGVNWVRQAPGQGLEWMGWINTKDGNPTYAQGFTGRFVFSL DTSISTTYLQISSLQAGDTAVYYCAREHDYYYGMDVWGQGTTLTVVSS (SEQ ID NO: 776)
VL	LPVLTQPASVSGSPGQSITISCTGTSSDVGGYNYVSWYQQHPGKAPKLMIEVSKRPSGVPDRFSGSKSGNT ASLTVSGLQAEDEADYYCSAWDDSLNADVFGGGTKLTVL (SEQ ID NO: 777)
VL	QPVLTQPPSASGTPGQRTISCSGSSSNIGTNTVNWYQQVPGTAPKLLIHGNDQRPSGVPDRFSGSKSDTSA SLAITGLQSDDDADYYCSAWDDSLNADVFGGGTKLTVL (SEQ ID NO: 778)
VL	QAVLTQPPSASATPGQRTISCSGSDSNIGTNYVYWYQQFPGTAPQPLIYRDNQRPSGVPDRFSGSKSGTSA SLAISGLRSEDEATYFCSTWDDSLNGWVFGGGTKLTVL (SEQ ID NO: 779)

VL	QPVLTQPRSVSGSPGQSITTSCTGTSSDVGGYNYVSWYQQHPGKAPKLMIEVSNRPSGVSNRFGSGSKSGNT ASLTISGLQAEDEADYYCSSYTSSSTLEVFGTGKVTVL (SEQ ID NO: 780)
VL	YELMQPPSVSGAPGQRTISCTGSSSNIGAAVDVHWYQQLPKGAPKLMFANSNRPSGVPDRFGSGSKSGTSA SLAITGLQAEDEADYYCQSYDISLRAVFGTGKLTVL (SEQ ID NO: 781)
VL	SYELMQPPSASGTPGQRTISCSGSSSNIGTNTVNWYQHLPGTAPKLLIYSNNQRPSGVPDRFGSGSKSGTSA SLAISGLQSEDEADYYCATWDDSPNGWVFGGGTKLTVL (SEQ ID NO: 782)
VL	QAVLTQPPSVSAAPGQRTISCSGSSNIGADTYVSWYQQLPGTAPRLLIYDNDQRPSGIPDRFGSGSKSGTSA TLGITGLQTGDEADYYCGTWDDSSLSGVFGTGKVTVL (SEQ ID NO: 783)
VL	QSVLTQPASVSGSPGQSVTISCTGSSSDVGAYNFVSWYRQYPGKAPKLLIYEVNKRPSDVPDRFGSGSKFGNT ASLTVSGLQADDEADYYCSSYAGSTDVFGTGKVTVL (SEQ ID NO: 784)
VL	LPVLTQPPSVSGTPGQRTISCSGSSSNIGSNTVNWYQQLPGTAPKLLIYTNNQRPSGVPDRFGSGSKSGTSA SLAISGLQSEDEADYYCAAWDESNGDVFGTGKVTVL (SEQ ID NO: 785)
VL	AIRMTQSPSFLSASVGDRVTITCRTSQNIYNYLNWYQQKPGKAPELLI FVASSLQSGVPSRFGSGSGTDFT LTISSLQPEDFATYFCLQDHSYPYTFGQGTKVEIK (SEQ ID NO: 786)
VL	LPVLTQPPSVSEVPGQRTISCSGGISNIGSNAVNWYQHFGKAPKLLIYYNDLLPSGVSDRFSASKSGTSA SLAISGLRSEDEADYYCAAWDDNLSAYVFATGKVTVL (SEQ ID NO: 787)
VL	QSALTQPPSASGSPGQSVTISCTGTSSDVGGYNYVSWYQHHPGKAPKLMIEVSKRPSGVPDRFGSGSKSAIT ASLTISGLLTEDEADYYCSAWDDSLNADVFGGGTKVTVL (SEQ ID NO: 788)
VL	DIQMTQSPSSLSASVGDRVTITCRASQSISTYLNWYQQKPGKAPKVLITDASSLETGVPSRFGSGSGTDFT FTISSLQPEDTATYFCQQYDDLPTFGQGTKLEIK (SEQ ID NO: 789)
VL	QAGLTQPRSVSGSPGQSVTISCTGTSSDVGGYNYVSWYQQHPGKAPKLMIDVTKRPSGVSNRFGSGSKSGNT ASLTISGLQAEDEADYYCSSYTSSSTYVFGTGKVTVL (SEQ ID NO: 790)
VL	QPVLTQPASVSGSPGQSITISCTGTSSDVGGYNYVSWYQQHPGKAPKLMIDVSNRPSGVSNRFGSGSKSGNT ASLAITGLQSDDDADYYCSAWDDSLNADVFGGGTKLTVL (SEQ ID NO: 791)
VL	QAGLTQPPSVSKGLRQTATLTCTGNSNNIGDQGAAWLQQHQGHPPRLLSYRNNNRPSGISERLSASRSGNIA SLTITGLQPEDEADYYCSAWDDSSLSVWVFGGGTKLTVL (SEQ ID NO: 792)
VL	QAVLTQPPSASGTPGQRTISCSGSSSNIGSNTVNWYQQLPGTAPKLLIYSNNQRPSGVPDRFGSGSKSGTSA SLAISGLQSEDEADYYCAAWDDSLNGYVFGTGKLTVL (SEQ ID NO: 793)
VL	AIRMTQSPSTLSASVGDRVTITCRASENIRNLLAWYQQKPGKAPELLIHGASTLGTGVPSRFGSGSGTEFT LTISSLQPDDEFATYYCQQYESYFNTFGQGTKVEIK (SEQ ID NO: 794)
Group L	
VH	QVQLVESGPGVKKPGSSSLKLSCTVSGFTFSSYDYYMHVVRQAPNGLEWMAVIWYSGSNTYYNDLSKSRFSI TRDNSKNTAYMQNLNLAEDTAVYYCARAYFGVDVWGQGLTVTVSS (SEQ ID NO: 795)
VL	DIVMTQSPASLSVSGDRATISCRASQGIGNTLAWYQQKPGQAPKRLLIYRASQGIGNTLGVPARFSGDGD GTDFTLTIDDLPEPEDFATYYCQQYDHPVPLTFGQGTKLEIK (SEQ ID NO: 796)
Group M	

VH	EVQLVQSGAEVKKPGASVKVSCKAFGYTFTTYPIEWMRQAHGQGLEWIGNFHPYNDDTKYNEKFKGRVTMTV DKSTTTVYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 797)
VH	EVQLVQSGAEVKKPGASVKVSCKAFGYTFTTYPIEWMRQAHGQGLEWIGNFHPYNDDTKYNEKFKGRVTMTR DTSTSTVYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 798)
VH	EVQLVQSGAEVKKPGASVKVSCKAFGYTFTTYPIEWVRQAPGQGLEWMGNFHPYNDDTKYNEKFKGRVTMTR DTSTSTVYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 799)
VH	EVQLVQSGAEVKKPGSSVKVSCKASGYTFTTYPIEWVRQAPGQGLEWMGNFHPYNDDTKYNEKFKGRVTITA DKSTSTAYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 800)
VH	EVQLVQSGAEVKKPGSSVKVSCKAFGYTFTTYPIEWMRQAHGQGLEWIGNFHPYNDDTKYNEKFKGRVTITV DKSTTTVYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 801)
VH	EVQLVQSGAEVKKPGSSVKVSCKAFGYTFTTYPIEWMRQAHGQGLEWIGNFHPYNDDTKYNEKFKGRVTITA DKSTSTAYMELSSLRSEDTAVYYCARENYGSHGGFVYWGQGTLLTVS (SEQ ID NO: 802)
VL	ENVLTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLWIYSTSNLASGVPDRFSGSGSGTSY TLTISSLQPEDFATYYCQQYNGYPLTFGGGTKVEIK (SEQ ID NO: 803)
VL	ENVLTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLFIYSTSNLASGVPSRFSGSGSGTDY TLTISSLQPEDFATYYCQQYNGYPLTFGGGTKVEIK (SEQ ID NO: 804)
VL	ENVLTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLWIYSTSNLASGVPDRFSGSGSGTSY TLTISSLQPEDFATYYCQQYNSYPLTFGGGTKVEIK (SEQ ID NO: 805)
VL	ENVMTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLFIYSTSNLASGVPSRFSGSGSGTDY TLTISSLQPEDFATYYCQQYNGYPLTFGGGTKVEIK (SEQ ID NO: 806)
VL	ENVMTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLFIYSTSNLASGVPSRFSGSGSGTDY TLTISSLQPEDFATYYCQQYNSYPLTFGGGTKVEIK (SEQ ID NO: 807)
VL	ENVLTQSPFSLASVGDRTITCRASSSVISSYLHWYQQKPAKAPKLFIYSTSNLASGVPSRFSGSGSGTDY TLTISSLQPEDFATYYCQQYNSYPLTFGGGTKVEIK (SEQ ID NO: 808)
VL	ENVLTQSPGTLSSLSPGERATLSCRASSSVISSYLHWYQQKPGQAPRLWIYSTSNLASGVPDRFSGSGSGTSY TLTISRLEPEDFATYYCQQYNGYPLTFGGGTKVEIK (SEQ ID NO: 809)
VL	ENVLTQSPGTLSSLSPGERATLSCRASSSVISSYLHWYQQKPGQAPRLWIYSTSNLASGVPDRFSGSGSGTSY TLTISRLEPEDFATYYCQQYNSYPLTFGGGTKVEIK (SEQ ID NO: 810)
Group N	
VH	QVQLQQPGAELVKPGASVKLSCKASGYTFTSYLYWMKQRPQGQGLEWIGGVNPSNGGTNFSEKFKSKATLTV DKSSSTAYMQLSLSTSEDSAVYYCTRRDSNYDGGFDYWGQGTLLTVSSAK (SEQ ID NO: 631)
VL	DIVLTQSPSTSLAVSLGQRATISCRASKSVSTSGFSYLHWYQQKPGQPPKLLIFLASNLESGVPPARFSGSGSG TDFTLNHPVEEEDAATYYCQHSWELPLTFGAGTKLELK (SEQ ID NO: 635)
Group O	
VH	EVQLVESGGGLVQPGGSLRLSCAASGFTFSDYGMHWVRQAPGKGLEWVAYISSGSYTIYSADSVKGRFTISR DNAKNTLYLQMSSLRAEDTAVYYCARRGYGSFYEYFDYWGQGTLLTVS (SEQ ID NO: 1521)

VL	QIVLTQSPATLSLSPGERATLSCSASSSVSYMYWYQQKPGQAPRLLIYLTSNRATGIPARFSGSGSGTDYTL TISSLEPEDFAVYYCQQWSSNPFTFGQGTKLEIK (SEQ ID NO: 1522)
Group P	
VH	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWVRQATGQGLEWMGNIYPGTGGSNFDEKFKNRVTITA DKSTSTAYMELSSLRSED TAVYYCTRWTTGTGAYWGQGT TTVTVSS (SEQ ID NO: 1523)
VH	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWIRQSPSRGLEWLGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTTGTGAYWGQGT TTVTVSS (SEQ ID NO: 1524)
VH	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWVRQAPGQGLEWMGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTTGTGAYWGQGT TTVTVSS (SEQ ID NO: 1525)
VL	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLT WYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTEFTLTITSSLPDDFATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1526)
VL	DIQMTQSPSSLSASVGDRVTITCKSSQSLDSDGNQKNFLT WYQQKPGQAPRLLIYWASTRESGIPPRFSGSG YGTDFTLTINNIESEDAAYYFCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1527)
VL	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLT WYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLPEDIATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1528)
VL	DIVMTQTPLSLPVTPGEPASISCKSSQSLDSDGNQKNFLT WYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1529)
VL	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLT WYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1530)
VL	EIVLTQSPDFQSVTPKEKVTITCKSSQSLDSDGNQKNFLT WYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1531)
VL	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLT WYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1532)
VL	DIQMTQSPSSLSASVGDRVTITCKSSQSLDSDGNQKNFLT WYQQKPGQSPQLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1533)
VL	DVVMTQSPPLSLPVTLGQPASISCKSSQSLDSDGNQKNFLT WYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIK (SEQ ID NO: 1534)
HC	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWVRQATGQGLEWMGNIYPGTGGSNFDEKFKNRVTITA DKSTSTAYMELSSLRSED TAVYYCTRWTTGTGAYWGQGT TTVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDKRV SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVDSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVSVLT VLVHQLDNLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFIYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 1535)

HC	QVQLVQSGAEVKKPGASVKVSCKASGYTFTTYWMHWIRQSPSRGLEWLGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTGTGAYWGQGTITVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDRVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 1536)
HC	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWIRQSPSRGLEWLGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTGTGAYWGQGTITVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDRVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 1537)
HC	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWVRQAPGQGLEWMGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTGTGAYWGQGTITVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDRVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 1538)
HC	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWVRQATGQGLEWMGNIYPGTGGSNFDEKFKNRVTITA DKSTSTAYMELSSLRSEDVAVYYCTRWTGTGAYWGQGTITVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDRVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLG (SEQ ID NO: 1539)
HC	EVQLVQSGAEVKKPGESLRISCKGSGYTFTTYWMHWIRQSPSRGLEWLGNIYPGTGGSNFDEKFKNRFTISR DNSKNTLYLQMNSLRAEDTAVYYCTRWTGTGAYWGQGTITVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDHKPSNTKVDRVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLG (SEQ ID NO: 1540)

LC	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLTQYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTEFTLTITSSLPDDFATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1541)
LC	DIQMTQSPSSLSASVGDRVTITCKSSQSLDSDGNQKNFLTQYQQKPGQAPRLLIYWASTRESGIPPRFSGSG YGTDFTLTINNIESEDAAYYFCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1542)
LC	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLTQYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLPEDIAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1543)
LC	DIVMTQTPLSLPVTPGEPASISCKSSQSLDSDGNQKNFLTQYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1544)
LC	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLTQYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1545)
LC	EIVLTQSPDFQSVTPKEKVTITCKSSQSLDSDGNQKNFLTQYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1546)
LC	DIQMTQSPSSLSASVGDRVTITCKSSQSLDSDGNQKNFLTQYQQKPGQSPQLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1547)
LC	DVVMQTSPSLSPVTLGQPASISCKSSQSLDSDGNQKNFLTQYQQKPGKAPKLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1548)
LC	EIVLTQSPATLSLSPGERATLSCKSSQSLDSDGNQKNFLTQYQQKPGQAPRLLIYWASTRESGVPSRFSGSG SGTDFTFITSSLEAEDAATYYCQNDYSYPYTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFN RGEC (SEQ ID NO: 1834)
Group Q	

VH	EVQLLESGGGLVQPGGSLRLSCSAS <u>GF</u> TFSSYTMNWVRQAPGKGLEWVSGISDTGGNTYYTDSVKGRFTVSR DNSKNTLSLQMNSLRAEDTAVYYC <u>AKDQ</u> GGSYPPYFHYWGQGSGLTVTVSS (SEQ ID NO: 1549)
VH	EVQLVESGGGLVQPGGSLRLSCAAS <u>GF</u> TVSNNYMSWVRQAPGKGLEWVSVIYSGGFTYYTDSVKGRFTISR NSKNTLYLQMNSLRAEDTAVYYCARYYYDTSYWTFFDYWGQGTSLTVTVSS (SEQ ID NO: 1550)
VH	QVQLVESGGGVVQSGRSLRLSCAAS <u>GF</u> TFSSYGMHWVRQAPGKGLEWVAVIWYDGSNIYYSDSVKGRFTISR ANSKNTLYLQMNSLRAEDTAVYYC <u>CARPGH</u> WNYFFEYWGQGTSLTVTVSS (SEQ ID NO: 1551)
VH	EVQLVESGGGLVQPGGSLRLSCGAS <u>GF</u> TFRNYDMHWVRQITGKGLEWVSAIGSAGDTYYPDSVKGRFTISRE NAKNSLYLQMNSLRVGD TAVYYC <u>TRDIHCS</u> STRCYGMDVWGQGTSLTVTVSS (SEQ ID NO: 1552)
VH	EVQLVESGGGLVKPGGSLRLSCAAS <u>GF</u> KFSNEWMSWVRQAPGKGLEWVGRIKSKTDGGTTDYAAPVKGRFTI SRDDSKNTLYLQMNSLKTEDTAVYYC <u>TTDQDF</u> WSGYTGADYYGMDVWGQGTMTVTVSS (SEQ ID NO: 1553)
VH	QMQLQQWGAGLLKPSETLSLTCVVYGGSLNGYYWSWIRQSPGKGLEWIGEIDHSGSTNYNPSLKNRVTMSVD TSKIQFSLKLTSTVTADTAVYYC <u>CAREGL</u> LPFDYWGQGTSLTVTVSS (SEQ ID NO: 1554)
VH	QLQLQESGPDLVKPSDTLSLTCVSDDSISSTYYWAWIRQPPGKGLEWIGSMSYNGNNYYNPSLKSRVAIS AGTSQKQFSLKLTSTVTAADTAVYHC <u>CARHLG</u> YNGNWYPFDYWGQGTSLTVTVSS (SEQ ID NO: 1555)
VH	EVQVLESGGGLVEPGRSLRLSCKAS <u>GF</u> TFDDYAMHWVRQTPGKALEWVSGINWSGNNIGYADSVKGRFTISK DDAKNSLYLQMNSLRPEDTALYYCTKDISITGTLDADFVWGQGTMTVTVSS (SEQ ID NO: 1556)
VH	QVQLVESGGGVVQPGRSLRLSCAAS <u>GF</u> TFSSYGMHWVRQAPGKGLEWVAIIWSDGDSEYNLDSVKGRFTISR DNSKNTLYLQMNSLRVEDSAVYYC <u>CARDR</u> LEDIYWQGTMTVTVSS (SEQ ID NO: 1557)
VH	EVQLLESGGVLVQPGGSLRLSCAAS <u>GF</u> TFSNFGMTWVRQAPGKGLEWVSGISGGGRDITYFADSVKGRFTISR DNSKNTLYLQMNSLKGEDTAVYYC <u>CVK</u> WGNIFYFDYWGQGTSLTVTVSS (SEQ ID NO: 1558)
VH	EVQLVESGGGVVRPGGSLRLSCAAS <u>GF</u> TFDDYGMWVRQVPGKGLEWVSGISWNDGKTVYAESVKGRFIISR DNAKNSLYLEMNSLRAEDTALYYC <u>CARDWQ</u> YLIERYFDYWGQGTSLTVTVSS (SEQ ID NO: 1559)
VH	EVQLVESGGGVVRPGGSLRLSCTAS <u>GF</u> TFDDYGMWVRQAPGKGLEWISGIGWTGGRSSYADSVRGRFTISR DNAKNSLYLQMNSLGAEDTALYYC <u>CARDRQ</u> WLQWYFDYWGQGTSLTVTVSS (SEQ ID NO: 1560)
VH	EVQLVESGGRVVRPGGSLRLSCAAS <u>GF</u> TFDDYGMWVRQLPGKGLEWVAGISWNDGKTVYAESVKGRFIISR DNAKNSLHLEMNSLRAEDTALYYC <u>CARDWQ</u> YLIDRYFDFWQGTSLTVTVSS (SEQ ID NO: 1561)
VH	EVQLVESGGGLVQPGGSLRLSCAAS <u>GF</u> TFDDYAMHWVRQAPGKGLEWVSGIGWSSGSIGYADSVKGRFTISR DNAKNSLYLQMDSLRPEDSALYYC <u>AKAYT</u> FMITLYFDYWGQGTSLTVTVSS (SEQ ID NO: 1562)
VH	EVQLVESGGGLVQPGRSLRLSCAAS <u>GF</u> TFDDYDMHWVRQAPGKGLEWVSGSGWNRGSLGYADSVKGRFTISR DNAKNSLYLQMNSVRVEDTALYYC <u>AKGFV</u> VSAAYFDYWGQGTSLTVTVSS (SEQ ID NO: 1563)
VH	QVQLVQSGAEVKRPGSSVKVSKVSGVTFRNFAIIWVRQAPGQGLEWMGGIIPFFSAANYAQSFGQGRVTITP DESTSTAFMELASLRSED TAVYYC <u>CAREGER</u> GHTYGFYWGQGTSLTVTVSS (SEQ ID NO: 1564)
VH	EVQLVESGGGLVQSGRSLRLSCAAS <u>GF</u> TFDDYAMHWVRQPPGKGLEWVSGINWNRGRTGYADSVKGRFTISR DNAKNSLYLQMNDLRVEDTALYYC <u>AKAEQ</u> WLDEGYFDYWGQGTSLTVTVSS (SEQ ID NO: 1565)
VH	EVQLVESGGGLVQRGGSLRLSCAASGFSSSYAMNWVRQAPGKGLEWVSTISDSGGSTYYADSVKGRFTISR DNSKNTLSLQMNSLRAEDTAVYYC <u>AKDQ</u> GGSYPPYFHYWGQGTSLTVTVSS (SEQ ID NO: 1566)

VH	EVQLVESGGGLVQPGRSLRLSCAAS <u>GFTTFEDY</u> AMHWVRQAPGKGLEWVSGIGWSNVKIGYADSVKGRFTISR DNVRNSLYLQMNSLRTEDTAFYYC <u>VKAYT</u> SMLTLYFDYWGQGTMTVTVSS (SEQ ID NO: 1567)
VH	QVQLVQSGAEVKRPGASVKVSCKASGYTFTSFYMYWVRQAPGQGLEWMGI <u>INPSDGST</u> SNAQKFQGRVTMTR DTSTSTVYMELSSLRSEDTAVYYCARRVAGDIFDIWGQGTMTVTVSS (SEQ ID NO: 1568)
VH	QVQLQESGPGLVKPSSETLSLTCTVSGGSISSYHWNWIRQSPGKGLEWIGYIYYIGSTDYNPSLESRTISVD TSKNQFSLKLLSVTAADTAVYYC <u>CARVPVGATGASDV</u> WGQGTMTVTVSS (SEQ ID NO: 1569)
VH	EVQLVESGGGSVVRPGGSLRLSCVVS <u>GFTTFEDY</u> GLSWVRQIPGKGLEWVSGISWTGGNTGYADSVKGRFTISR DNAKNSLYLQMNSLRAEDTALYHCTRD <u>RQWL</u> MQWYFDYWGQGTMTVTVSS (SEQ ID NO: 1570)
VH	QVQLVESGGGVVQPGRSLRLSCSAS <u>GFTTF</u> SAYAMHWVRQAPGKGLEWVA <u>AI</u> SYGGSDKYYADSVKGRFTISR DNSKNTLYLQMNSLRDDEDTAVYYC <u>AKSAHWN</u> FFFDYWGQGTMTVTVSS (SEQ ID NO: 1571)
VH	EVQLVESGGGLVQPGRSLRLSCVAS <u>G</u> FALHDYAMHWVRQVPKGKLEWVSSISWNSGVIYADSLKGRFTISR DNAKNSLYLQMNSLRAEDTALYYC <u>AKGSGSY</u> YVSWFDPWGQGTMTVTVSS (SEQ ID NO: 1572)
VH	QLQLQESGPGLVQPSETLSLTCTVSGDSISSTAYHWDWIRQPPGKGLEWIGTITYNGNTYFNPSLKSRVTIS VDTSKNQFSLKLLSMTAAETAUFYCARHLGYN <u>SDFFPFDF</u> WGQGTMTVTVSS (SEQ ID NO: 1573)
VH	EVQLVESGGGLVVRPGGSLRLSCAAS <u>GFTTF</u> STYAMAWVRQTPGKGLEGVSAIGGSGDSTYYVDSVKGRFTISR DNSKSTLFLQMNSLRAEDTAVYYC <u>VKVRNYDGSF</u> DIWGQGTMTVTVSS (SEQ ID NO: 1574)
VL	DIQMTQSPSSLSASVGDRVTITCRAS <u>Q</u> SISSYLIWYQQKPGTAPKFLIYAASSLQSGVPSRFRSGSGSGTDFT LTISSLQPEDFATYYC <u>QQSYSTP</u> PITFGQGTGLEIK (SEQ ID NO: 1575)
VL	EIVMTQSPATLSVSPGERATLSCRAS <u>QSVSSN</u> LAWYQQKPGQAPRLLIYGASTRATGIPARFSGSGSGTEFT LTISSLQSGDFAVYYC <u>QQYNNWPLT</u> FGGGTKVEIN (SEQ ID NO: 1576)
VL	EIVMTQSPATLSVSPGERATLSCRAS <u>QSVSSN</u> LAWYQQKPGQAPRLLIYGASTRATGIPARFSGSGSGTEFT LTISSLQSGDFAVYYC <u>QQYNNWPLT</u> FGGGTKVEIN (SEQ ID NO: 1577)
VL	DIQMTQSPSSLSASVGDRVTITCRAS <u>Q</u> SINNYLNWYQQKPGKAPKLLIYTASSLQSGVPSRFRSGSGSGTDFT LTISSLQPEDFATYYC <u>QQSYSTP</u> PLTFGQGTGLEIK (SEQ ID NO: 1578)
VL	DIQMTQSPSSLSASVGDRVTITCRAS <u>Q</u> SISNYLNWYQQKPGKAPKLLIYAASSLQSGVPSRFRSGSGSGTDFT LTISSLQPEDFATYYC <u>QQSYSTP</u> PITFGQGTGLEIK (SEQ ID NO: 1579)
VL	DIQMTQSPSSLSASVGDRVTITCRAS <u>Q</u> SISSYLNWYQQKPGKAPKLLIYAASSLQSGVPSRFRSGSGSGTDFT LTISSLQPEDFATYYC <u>QQSYSTP</u> PITFGQGTGLEIK (SEQ ID NO: 1580)
VL	EIVLTQSPGTLSLSPGERATLSCRAS <u>QSVSY</u> SNYLAWYQQNPGQAPRLLIYAASNRTGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYC <u>CHQYATSPWT</u> FGQGTGLEIK (SEQ ID NO: 1581)
VL	EIVLTQSPGTLSLSPGERATLSCRAS <u>QSVSSSY</u> LAWYQQKPGQAPRLLIYGASSRTTGIPDRFSGSGSGTDF TLTISRLEPEDFAVYYC <u>QQYGSSPWT</u> FGQGTGLEIK (SEQ ID NO: 1582)
VL	DIQMTQSPISVSASVGDRVTITCRAS <u>Q</u> GISNWLAWYQQKPGIAPKLLIYSASSLQSGVPSRFRSGSGSGTDFT LTIGSLQPEDFATYYC <u>QQAHSFPLT</u> FGGGTKVEIK (SEQ ID NO: 1583)
VL	DIQMTQSPSSLSASVGDRVTITCRAS <u>Q</u> GIRNDLGWYQQKPGKAPKRLIYAASNLSLQSGVPSRFRSGSGSGTEFT LTISSLQPEDFATYYC <u>LQHNSYPLT</u> FGGGTKVEIK (SEQ ID NO: 1584)

VL	DIQMTQSPSSLSASVGDSTITITCRAS <u>LSINTFLN</u> WYQQKPGKAPNLLIYAASSLHGGVPSRFSGSGSGTDFTLTIRTLQPEDFATYYC <u>QQSSNTPFT</u> FGPGTVVDFR (SEQ ID NO: 1585)
VL	EIVLTQSPGTLISLSPGERATLSCRAS <u>QSVSSSYLAWY</u> QQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYC <u>QQYGSSPWT</u> FGQGTKVEIK (SEQ ID NO: 1586)
VL	DIQMTQSPSSLSASVGDRTITITCRAS <u>QSISSYL</u> NWYQQKPGKAPKLLIYAASSLQSGVPSRFSGSGSGTDFTLTISLQPEDFATYYC <u>QQSYSTPPIT</u> FGQGTTRLEIK (SEQ ID NO: 1587)
HC	EVQLLESGGVLVQPGGSLRLSCAAS <u>GFTFSNFGMTWVRQ</u> APGKGLEWVSGISGGGRDITYFADSVKGRFTISRDN SKNTLYLQMNLSLKGEDTAVYYC <u>CVKWGNIYFDYWGQ</u> GLTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTI SKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSV MHEALHNHYTQKSLSLGLK (SEQ ID NO: 1588)
HC	EVQVVESGGGLVEPGRSLRLSCKAS <u>GFTFDDYAMHWVRQ</u> TPGKALEWVSGISWSGNNIGYADSVKGRFTISKDDAKNSLYLQMNSLRPEDTALYYCTKDISITGTLDADFVWGQGTMTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTI SKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSV MHEALHNHYTQKSLSLGLK (SEQ ID NO: 1589)
HC	EVQLVESGGGLVLRPGGSLRLSCAAS <u>GFTFSTYAMAWVRQ</u> TPGKLEGVSAIGGSGDSTYYVDSVKGRFTISRDN SKSTLFLQMNSLRAEDTAVYYC <u>CVKVRNYDGSFDI</u> WGQGTMTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTI SKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSV MHEALHNHYTQKSLSLGLK (SEQ ID NO: 1590)
HC	QVQLVQSGAEVKRPGSSVKVSGVTFRNFAI I <u>WVRQAPGQGLEWMGGIIPFFSA</u> ANYAQSFGGRVTITPDESTSTA FMELASLRSEDTAVYYC <u>CAREGERGHTYGF</u> DYWGQGTMTVTVSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTI SKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSV MHEALHNHYTQKSLSLGLK (SEQ ID NO: 1591)

LC	DIQMTQSPSSLSASVGDSTITITCRAS <u>LSINTFLN</u> WYQQKPGKAPNLLIYAASSLHGGVPSRFRSGSGSGTDFT LTIRTLQPEDFATYYC <u>QQSSNTPFT</u> FGPGTVVDFRRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREA KVQWKVDNALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 1592)
LC	DIQMTQSPISVSASVGDRTITITCRAS <u>QGISNWL</u> AWYQQKPGIAPKLLIYSASSLQSGVPSRFRSGSGSGTDFT LTIGSLQPEDFATYYC <u>QQAHSFPLT</u> FGGGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREA KVQWKVDNALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 1593)
LC	EIVLTQSPGTLSLSPGERATLSCRAS <u>QSVSSSYL</u> AWYQQKPGQAPRLLIYGASSRATGIPDRFSGSGSGTDFT TLTISRLEPEDFAVYYCQQYGSSPWTFGQGTKVEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 1594)
LC	DIQMTQSPSSLSASVGDRTITITCRAS <u>QSISSYL</u> NWYQQKPGKAPKLLIYAASSLQSGVPSRFRSGSGSGTDFT LTISSLQPEDFATYYC <u>QQSYSTPPIT</u> FGQGTTRLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPRE AKVQWKVDNALQSGNSQESVTEQDSKDYSLSSLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 1595)
Group R	
VH	EVKLIVESGGGLVKPGGSLKLSCAASGFTFSSYGM <u>SWLRQT</u> PEKRLEWVATMSGGGRDIYYPDSMKGRFTISR DNAKNNLYLQMSSLRSEDALYYCAR <u>QYYDDWFAY</u> WGQGTTLTVSA (SEQ ID NO: 1719)
VH	QVQLKQSGPGLVQP <small>SQ</small> NLSVTCTVSGFSLT <u>TYGVHW</u> RQSPGKGLEWLGVIWGGSTDYNAAFISRLTISKD NARSQVFFKMNSLQVNDTAMYYCAREKS <u>SVYGNYVG</u> AMDYWGQGTSTTVSS (SEQ ID NO: 1720)
VH	EVKLIVESGGGLVKPGGSLKLSGASGFTFSSYGM <u>SWVRQT</u> PEKRLEWVATISGGGRDIYYPDSVKGRLTISR DNAKNNLYLQMSSLRSEDALYYCVR <u>QYYDDWFAY</u> WGQGTTLTVSA (SEQ ID NO: 1721)
VH	DVQLQESGPGLVKPSQSLSLTCTVTGYSITSDYAWNWIRQFPGNQLEWMAYISYSGYTSYNPSLKSRIISITR DTSKNQFFLQLNSVTTEDTATYYCARSLDYDYGTMDYWGQGTSTTVSS (SEQ ID NO: 1722)
VH	EVKLIVESGGGLVKPGGSLKLSCAASGFAFRSYDMSWVRQTPEKILEWVATISGGGSYTYQDSVKGRFTISR DNARNTLYLQMSSLRSEDALYYCASPYGPYFDYWGQGTTLTVSS (SEQ ID NO: 1723)
VH	DVQLQESGPGLVKPSQSLSLTCTVTGYSITSDYAWNWIRQFPGNQLEWMAYISYSGYTSYNPSLKSRIISITR DTSRNQFFLQLNSVTTEDTATYYCARSLDYDYGTMDYWGQGTSTTVSS (SEQ ID NO: 1724)
VH	EVKLIVESGGGLVKPGGSLKLSASGFSFSYDMSWVRQTPEKGLEWVATISGGGRNTYFIDSVKGRFTISR DNVKNLYLLMSSLRSEDALYYCASPYEGAVDFWGQGTSTTVSS (SEQ ID NO: 1725)
VH	EVKLIVESGGGLVKPGGSLKLSCAASGFTFSSYGM <u>SWVRQT</u> PEKRLEWATISGGGRDITYYLDVSKGRFTISRD NAKNNLYLQMSSLRSEDALYYCVR <u>QYYDDWFAY</u> WGQGTTLVNSA (SEQ ID NO: 1726)
VH	QVQLQQSGDELVRPGTSVKMSCKAAGYFTNNWIGWVKQRPGHGLEWIGDFYPGGGYTNYNEKFKGKATLTA DTSSSTAYMQLSLTSEDYAIYYCARGYGTNYWYFDVWGAGTTTVTVSS (SEQ ID NO: 1727)
VH	QIHLVQSGPELKKPGETVKISKASGYFTNFNMWVKQAPGKGLKWMGWISGYTREPTYAADFKGRFAISL ETSASTAYLQINDLKNEDMATYFCARDVFDYWGQGTTLTVSS (SEQ ID NO: 1728)

VH	QIQLVQSGSELKKPGASVKVSCKASGYTFTNFGMNWVRQAPGQGLKWMGWISGYTREPTYAADFKGRFVISL DTSVSTAYLQISSSLKAEDTAVYYCARDVFDYWGQGTLLTVSS (SEQ ID NO: 1729)
VH	QVQLQESGPGILVKPSQTLSTCTVSGYSISSDYAWNWRQPPGKLEWMAYISYSGYTSYNPSLKSRTISR DTSKNQFSLKLSSVTAADTAVYYCARSLDYDYGTMQYWGQGTLLTVSS (SEQ ID NO: 1730)
VL	DIVLTQTPATLSVTPGDSVSLSCRASQSISNNLHWYQQKSHESPRLLIKYASQSISGIPSRFSGSGSGTDFT LNINSVETEDFGMYFCQQSNSWPLTFGAGTKLELKR (SEQ ID NO: 1731)
VL	SIVMTQTPKFLLVSAAGDRVTITCKASQSVSDDVAWYQQKPGQSPKLLIYYAFNRYTGVPDRFTGSGYGTDF FTISTVQSEDLAVYFCQQDYRSPWTFGGGKLEIKR (SEQ ID NO: 1732)
VL	DIVLTQSPATLSVTPGDSVSLSCRASQSISNDLHWYQQKSHESPRLLIKYVSQSISGIPSRFSGSGSGTDFT LSINSVETEDFGMYFCQQSDSWPLTFGAGTKLELKR (SEQ ID NO: 1733)
VL	QIVLSQSPAILLSASPGEKVTMTCRANSSVSSMHWYQQKPGSSPEPWIYAIISNLAFGVPTRFSGSGSGTSYSL TISRVEAEDAATYFCQQWSSRPPTFGGGKLEIKR (SEQ ID NO: 1734)
VL	DIQMNQSPSSLSASLGDTITITCHASQSVNVLWYQQKPGNIPKLLIYRASNLHTGVPSRFSGSGSGTGFT LTISLQPDIDATYFCQQGQSYPWTFGGGKLEIKR (SEQ ID NO: 1735)
VL	QIVLSQSPAILLSASPGEKVTMTCRANSSVSSMHWYQQKPGSSPEPWIYAIISNLAFGVPARFSGSGSGTSYSL TISRVEAEDAATYFCQQWNSRPPTFGGGKLEIKR (SEQ ID NO: 1736)
VL	DIVMTQSHKVMSTSVGDRVSITCKASQDVNDVAWYQQNPGQSPKLLIKWASTRHHGVPDRFTGSGSGTDFT LTISTVQSEDLADFFCQQYSTFPYTFGGGKLEIKR (SEQ ID NO: 1737)
VL	DIVLTQTPATLSVTPGDSVSLSCRASQSLSNNLHWYQQKSHESPRLLIKYASQSISGIPSRFSGSGSGTDFT LSINSVETEDFGMYFCQQSNSWPLTFGAGTKLEMKR (SEQ ID NO: 1738)
VL	NIVMTQTPKILFISAGDRVTITCKASQSVSNDVAWYQQKPGQSPKLLIYYAFTRYIGVPDRFTGSGYGTDF FTISTVQAEDLAVYFCQQDYSSPYTFGGGKLEIKR (SEQ ID NO: 1739)
VL	DIVLTQSPASLAVSLGQRATISCRASESDNYGYSFMNWFQKPGQPPKLLIYRASNLESGIPARFSGSGSR TNFTLTINPVEADDVATYFCQQSNADPTFGGGTNLEIKR (SEQ ID NO: 1740)
VL	DIVLTQSPASLAVSPGQRATITCRASESDNYGYSFMNWFQKPGQPPKLLIYRASNLESGVPARFSGSGSR TDFTLTINPVEANDTANYCQQSNADPTFGQGTKLEIK (SEQ ID NO: 1741)
VL	EIVLTQSPATLSLSPGERATLSCRANSSVSSMHWYQQKPGQSPPEPWIYAIISNLAFGVPARFSGSGSGTDYTL TISSLEPEDFAVYYCQQWSSRPPTFGQGTKLEIK (SEQ ID NO: 1742)
Group S	
VH	QVQLVQSGSEVKKSGSSVKVSCKTSGGTFSTITNYAINWVRQAPGQGLEWMGGILPIFGAAKYAQKFQDRVTI TADESTNTAYLELSSLTSEDATAMYYCARGKRWLQSDLQYWGQGTLLTVSS (SEQ ID NO: 1743)
VL	QPVLTQPASVSGSPGQSITISCTGSSSDVGSYDLVSWYQQSPGKVPKLLIYEGVKRPSGVSNRFGSGSGNT ASLTISGLQAEDADYCYSSYAGTRNFVFGGGTQLTVL (SEQ ID NO: 1744)
Group T	
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTNYYMYWVKQAPGQGLEWIGGINPSNGGTNYNEKFKNKATLTA DKSTSTAYMELSSLRSEDATAVYYCTRRDYYRDMGFDYWGQGTLLTVSS (SEQ ID NO: 1745)

VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYMYWVRQAPGQGLEWMGGVNPSNGGTNFNEKFKSRVTITA DKSTSTAYMELSSLRSED TAVYYCARRDYRYDMGFDYWGQGT TVTVSS (SEQ ID NO: 1746)
VH	EVQLQQSGPELVKPGASVKISCKASGYTFTNYYMYWVKQSHGKSLEWIGGINPSNGGTNYNEKFKNKATLTV DKSSSTAYMELNSLTSEDSAVYYCARRDYRYDMGFDYWGAGT TVTVSS (SEQ ID NO: 1747)
VH	EVQLQQSGPVLVKPGASVKMSCKASGYTFTSYMYWVKQSHGKSLEWIGGVNPSNGGTNFNEKFKSKATLTV DKSSSTAYMELNSLTSEDSAVYYCARRDYRYDMGFDYWGQGT TTLTVSS (SEQ ID NO: 1748)
VL	EIVLTQSPATLSLSPGERATISCRASKGVSTSGYSYLHWYQQKPGQAPRLLIY LASYLESGV PARFSGSGS TDFTLTISSELEPEDFATYYCQHSRELPLTFGTG TKVEIK (SEQ ID NO: 1749)
VL	QIVLTQSPA IM SASPGEKVTMT CR ASKGVSTSGYSYLHWYQQKPGSSPRLLIY LASYLESGV PVRFSGSGS TSYSLTISRMEAE DAATYYCQHSRELPLTFGT TRLEIK (SEQ ID NO: 1750)
Group U	
VH	QVQLQQSGAELMKPGASVKMSCKTTGYIFSSYIWGWVKQRP GH GLEWIGKIFPGSGSADYNENFKGKATFTV DTSSNTAYMQ LSSLTSEDSAVYYCARGYGNLYFDV WGAGT TVTVSS (SEQ ID NO: 619)
VH	DVQLQESGPGLVKPSQSLSLTCTVTGHSITSDYAWN WIRQ FPGNKLEWMGYISYSGRTSYNPSLT SRISITR DTSKNQFFLQLNSVT TEDTATYYCARGYALDYWGQGT SVTVSS (SEQ ID NO: 1835)
VH	EVKLVESGGGLVSPGGSLKLSCAASGFTFSTFGMSWVRQTPEKRLEWVATISGGGSDTYYPDSVQGRFIISR YNAKNNLYLQMN SLRPEDTALYYCARQGYDVYSWFAYWGQGT LTVVSA (SEQ ID NO: 1836)
VH	EVKLVESGGGLVKPGGSLKLSCAASGFTFSTYGM SWVRQTPEKRLQW VATISGGGSNTYYSDSVKGRFTISR DNAKNNLYLQMS SLRSED TALYYCARQ RDSAWFASWGQGT LTVVSA (SEQ ID NO: 1837)
VH	EVQLVESGGGLVKPGGSLRLS CAASGFTFSTFGMSWVRQAPGK GLEWVSTISGGGSDTYYPDSVQGRFTISR DNAKNSLYLQMN SLRAEDTAVYYCARQGYDVYSWFAYWGQGT LTVVSS (SEQ ID NO: 1838)
VH	EVQLVESGGGLVQPGGSLRLS CAASGFTFSTFGMSWVRQAPGK GLEWVATISGGGSDTYYPDSVQGRFTISR DNAKNSLYLQMN SLRAEDTAVYYCARQGYDVYSWFAYWGQGT LTVVSS (SEQ ID NO: 1839)
VH	EVQLVESGGGLVKPGGSLRLS CAASGFTFSTYGM SWVRQAPGKLEWVATISGGGSNTYYSDSVKGRFTISR DDSKNTLYLQMN SLKTEDTAVYYCARQDSAWFASWGQGT LTVVSS (SEQ ID NO: 1840)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFPHYGISWVRQAPGQGLEWMGWI SAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDVDYGTGSGYWGQGT LTVVSS (SEQ ID NO: 1841)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFTRQGISWVRQAPGQGLEWMGWI SAYNGNTKYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDVDYGS SGSYWGQGT LTVVSS (SEQ ID NO: 1842)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFSTFGISWVRQAPGQGLEWMGWI SAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDVDYSSGS SGSYWGQGT LTVVSS (SEQ ID NO: 1843)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFTRYGISWVRQAPGQGLEWMGW VSAHNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDADYGS SGSYWGQGT LTVVSS (SEQ ID NO: 1844)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFPHYGISWVRQAPGQGLEWMGWI SAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDAEYGS SGSYWGQGT LTVVSS (SEQ ID NO: 1845)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFTWYGISWVRQAPGQGLEWMGWI SAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSD DTAVYYCARDSEYSSGS SGSYWGQGT LTVVSS (SEQ ID NO: 1846)

VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FETYGI</u> SWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARD <u>AEYSLGSGYWGQ</u> GLTVTVSS (SEQ ID NO: 1847)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FRQYGI</u> SWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARD <u>AEYSGSGYWGQ</u> GLTVTVSS (SEQ ID NO: 1848)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFTWYGISWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARDSEYRSGSGYWGQGLTVTVSS (SEQ ID NO: 1849)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FRQYGI</u> SWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARD <u>AEYRSGSGYWGQ</u> GLTVTVSS (SEQ ID NO: 1850)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FTRYGI</u> SWVRQAPGQGLEWMGWVSAHNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARDADYRSGSGYWGQGLTVTVSS (SEQ ID NO: 1851)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFPYGISWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARDVDYRTGSGYWGQGLTVTVSS (SEQ ID NO: 1852)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FSTFGI</u> SWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARDVDYRSGSGYWGQGLTVTVSS (SEQ ID NO: 1853)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FRQGIS</u> SWVRQAPGQGLEWMGWISAYNGNTKYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARDVDYRSGSGYWGQGLTVTVSS (SEQ ID NO: 1854)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYRFPYGISWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARD <u>AEYRSGSGYWGQ</u> GLTVTVSS (SEQ ID NO: 1855)
VH	EVQLVQSGAEVKKPGASVKVSCKASGYR <u>FETYGI</u> SWVRQAPGQGLEWMGWISAYNGNTNYAQKLQGRVTMTT DTSTNTAYMELRSLRSDDTAVYYCARD <u>AEYRLGSGYWGQ</u> GLTVTVSS (SEQ ID NO: 1869)
VL	NIVMTQTPKFLIVSAGDRITITCKASQSVSDDVAWYQQKPGQSPKLLISYAFKRYIGVPDRFTGSGYGTDF FTISTVQAEDLAVYFCQQNYNSPYTFGGGTKLELKR (SEQ ID NO: 1856)
VL	QIVLSQSPAILSASPGEKVTMTCTRTSSSVNYMHWFQQKPGSSPKPIYATSKLASGVPARFSGSGSGTSYSL TISRVEAEDAATYFCQQWISDPWTFGGGTKLEIK (SEQ ID NO: 1857)
VL	SYELTQPPSVSVSPGQTARITCSGDALTTQYAYWYQQKPGQAPVMVIYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDADYCYCQADNSITYRVFGGGTKVTVL (SEQ ID NO: 1858)
VL	SYELTQPPSVSVSPGQTARITCSGDALSEQYAYWYQQKPGQAPVMVIYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDADYCYCQADNSITYRVFGGGTKVTVL (SEQ ID NO: 1859)
VL	SYELTQPPSVSVSPGQTARITCSGDALPKQYAYWYQQKPGQAPVMVIYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDADYCYCQADNSITYRVFGGGTKVTVL (SEQ ID NO: 737)
VL	SYELTQPPSVSVSPGQTARITCSGDALPKQYAYWYQQKPGQAPVMVLYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDADYCYCQSDNSITYRVFGGGTKVTVL (SEQ ID NO: 1860)
VL	SYELTQPPSVSVSPGQTARITCSGDALPMQYGYWYQQKPGQAPVMVIYKDTERPSGIPERFSGSSSGTKVTL TISGVQAEDADYCYCQADNSITYRVFGGGTKVTVL (SEQ ID NO: 1861)
VL	DIILTQSPASLAVSLGQRAAISCRASESDNSGISFMSWFQQKPGQPPKLLIYTASNQGSQGVPARFSGSGSG TEFSLNIHPMEEDDTAMYFCQQSKEVPWTFGGGTKLEIR (SEQ ID NO: 1862)

VL	DIVLTQSPASLAVSLGQRATISCRASENVDDYGVSFMNWFQQKPGQPPKLLIYPASNQSGVPPARFSGSGG TDFSLNIHPMEEDDTAMYFCQQSKEVPWTFGGGTKLEIK (SEQ ID NO: 1863)
VL	DIQLTQSPSFLSASVGRVTITCRASESDNSGISFMSWYQQKPGKAPKLLIYTASNQSGVPSRFSGSGG TEFTLTITISLQPEDFATYYCQQSKEVPWTFGQGTKVEIK (SEQ ID NO: 1864)
VL	EIVLTQSPATLSLSPGERATLSCRAESVDNSGISFMSWYQQKPGQAPRLLIYTASNQSGIPARFSGSGG TDFTLTISLQPEDFAVYYCQQSKEVPWTFGQGTKVEIK (SEQ ID NO: 1865)
VL	EIVLTQSPATLSLSPGERATLSCRASENVDDYGVSFMNWYQQKPGQAPRLLIYPASNQSGIPARFSGSGG TDFTLTISLQPEDFAVYYCQQSKEVPWTFGQGTKVEIK (SEQ ID NO: 1866)
VL	EIVLTQSPGTLSLSPGERATLSCRASENVDDYGVSFMNWYQQKPGQAPRLLIYPASNQSGIPDRFSGSGG TDFTLTISRLEPEDFAVYYCQQSKEVPWTFGQGTKVEIK (SEQ ID NO: 1867)
VL	DIVMTQSPDSLAVSLGERATINCRASENVDDYGVSFMNWYQQKPGQPPKLLIYPASNQSGVPDRFSGSGG TDFTLTISLQAEADVAVYYCQQSKEVPWTFGGGTKLEIK (SEQ ID NO: 1868)
Group V	
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIYPGSSLTNYNEKFKNRVTMTR DTSTSTVYMEISSLRSEDVAVYYCARLLTGTFAYWGQGLTVTVSS (SEQ ID NO: 1870)
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIYPGSSLTNYNEKFKNRVTMTR DTSTSTVYMEISSLRSEDVAVYYCARLSTGTFAYWGQGLTVTVSS (SEQ ID NO: 1871)
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIYPGSSITNYNEKFKNRVTMTR DTSTSTVYMEISSLRSEDVAVYYCARLTGTFAYWGQGLTVTVSS (SEQ ID NO: 1872)
VH	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIWPGSSLTNYNEKFKNRVTMTR DTSTSTVYMEISSLRSEDVAVYYCARLLTGTFAYWGQGLTVTVSS (SEQ ID NO: 1873)
VH	DIVMTQSPDSLAVSLGERATINCKSSQSLWDSGNQKNFLTWYQQKPGQPPKLLIYWTSTRESGVPDRFSGG SGTDFTLTISLQAEADVAVYYCQNDYFYPLTFGGGTKVEIK (SEQ ID NO: 1874)
VL	DIVMTQSPDSLAVSLGERATINCKSSQSLWDSGNQKNFLTWYQQKPGQPPKLLIYWTSTRESGVPDRFSGG SGTDFTLTISLQAEADVAVYYCQNDYFYPLTFGGGTKVEIK (SEQ ID NO: 1875)
VL	DIVMTQSPDSLAVSLGERATINCKSSQSLWDSGNQKNFLTWYQQKPGQPPKLLIYWTSTRESGVPDRFSGG SGTDFTLTISLQAEADVAVYYCQNDYFYPHTFGGGTKVEIK (SEQ ID NO: 1876)
VL	DIVMTQSPDSLAVSLGERATINCKSSQSLWDSGNQKNFLTWYQQKPGQPPKLLIYWTSTRESGVPDRFSGG SGTDFTLTISLQAEADVAVYYCQNDYFYPLTFGGGTKVEIK (SEQ ID NO: 1877)
HC	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIYPGSSLTNYNEKFKNRVTMTR DTSTSTVYMEISSLRSEDVAVYYCARLSTGTFAYWGQGLTVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNVDPKPSNTKVDKRV SKYGPCCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVDSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVSVLTVHLQDNLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFIYPSDIAVEWESNGQPENNYKTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLGK (SEQ ID NO: 1878)

HC	QVQLVQSGAEVKKPGASVKVSCKASGYTFTSYWINWVRQAPGQGLEWMGNIYPGSSLTNYNEKFKNRVTMTR DTSTSTVYMELSSLRSED TAVYYCAR <u>LSTGTFA</u> WGQGLTVTVSSASTKGPSVFPLAPCSRSTSESTAALGC LVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVTVPSSSLGKTYTCNV DKPSNTKVDK RVE SKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVDSQEDPEVQFNWYVDGVEVHNAKTKP REEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQV SLTCLVKGFYPSDIAVEWESNGQPENNYKTTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNH YTQKSLSLSLG (SEQ ID NO: 1879)
LC	DIVMTQSPDSLAVSLGERATINCKSSQSLWDSGNQKNFLT WYQQKPGQPPKLLIYWTSY RESGVPDRFSGSG SGTDFTLTITSSQLAEDVAVYYCQNDYFYPH T FGGGTKVEIKRGTVAAPSVFI FPPSDEQLKSGTASV VCLLN NFYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSSLTTLTKADY EKH KVYACEVTHQGLSSPVTKSF NRGEC (SEQ ID NO: 1880)
Group W	
VH	EVQLQESGP ELVR PGASVKMSCKASGYTFTSYWMHWVKQRPQG LEW IGMIDPSNSETSLNQKFKDKATLNV DKSTNTAYMQLSLTS EDSAVYYCAR SRGNAYEMDYWGQGTSVTVSS (SEQ ID NO: 1972)
VH	EVQLQESGP ELVR PGASVKMSCKASGYTFTSYWMHWVKQRPQG LEW IGMIEPSSSETSLNQKFKDKATLNV DKSSNTAYMQLSLTS EDSAVYYCAR SRGNAYEMDYWGQGTSVTVSS (SEQ ID NO: 1973)
VH	EVQLQESGP ELVR PGASVKMSCKASGYTFTSYWMHWVKQRPQG LEW IGMIDPYSSETSLNQKFKDKATLNV DKISNTAYMQLSLTS EDSAVYFCAR SRGNAYDMDYWGQGTSVTVSS (SEQ ID NO: 1974)
VH	EVQLQESGP ELVR PGASVKMSCKASGYTFTSYWMHWVKQRPQG LEW IGMIDPSNSETSLNQKFKDKATLNV DKSSKTAYMQLSLTS EDSAVYYCAR SRGNAYDMDYWGQGTSVTVSS (SEQ ID NO: 1975)
VH	EVQLQESGAELVMPGASVKMSCKASGYTFTDYWMHWVKQRPQG LEW IGAIDTSDSYTSYHQNFKGKATLTE DESSSTAYMQLSLTS EDSAIYYCARRDYGGFGYWGQGT TLTVSS (SEQ ID NO: 1976)
VH	EVQLQESGP ELVK PGASVKIPCKASGYTFTDYNMDWVKQSHGKSLEWIGDIDPNNGGTIYNQKFKGKATLTV DKSSRTAYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1977)
VH	EVQLQESGP ELVK PGASVKIPCRASGYIFTDYNMDWVKQSHGKSLEWIGDIDPNNGGTIYNQKFKDKTTLTV DKSSRTAYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1978)
VH	EVQLQESGP ELVK PGASVKIPCKASGYTFTDYNMDWVKQNHGKSLEWIGDIDPNNGGTIYNQKFKGKATLTV DKSSRTAYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1979)
VH	EVQLQESGP ELVK PGASVKIPCKASGYTFTDYNMDWVKQSHGKSLEWIGDIDPNSGGSIYNQKFKGKATLTV DKSSRTVYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1980)
VH	EVQLQESGP ELVK PGASVKITCKASGYTFTDYNMDWVKQSHGKSLEWIGDIDPNNGGTIYNQKFKGKATLTV DKSSNTAYMELRSLAS EDTAVYYCAR WRSSMDYWGQGTSVSVSS (SEQ ID NO: 1981)
VH	EVQLQESGP ELVK PGASVKIPCKASGYTFTDYNMDWVKQSHGKSLEWIGDIDPNNGGTIYNQNFKGKATLTV DKSSSTAYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1982)
VH	EVQLQESGP ELVK PGASVKIPCKASGYTFTDYNMDWVKQSHGKSLEWIGDIDPNNGGTIYNQKFKGAALTV DKSSSTAYMELRSLTS EDTAVYYCAR WRSSMDYWGQGTSVTVSS (SEQ ID NO: 1983)

VH	EVQLQESGPPELVKPGASVKIPCKASGYTFSTDYNMDWVKQSHGKSLEWIGDIDPNNGGIIYNQKFKGKAALTV DKSSSTAYMELRSLTSED TAVYYCTRWRRSSMDYWGQGTSTVTVSS (SEQ ID NO: 1984)
VH	EVQLQESGPPELVKPGASVKIPCKASGYTFSTDYNMDWVKQSHGKSLEWIGDIDPNNGNTIYNQKFKGKATLTV DKSSSTAYMELRSLTSED TAVYYCTKWRSSMDYWGQGTSTVTVSS (SEQ ID NO: 1985)
VH	EVQLQESGPPELVKPGASVKIPCKASGYTFSTDYNMDWVMQSHGKSLEWIGDIDPNNGGTIYNQKFKGKATLTV DKSSSTAYMELRSLTSED TAVYYCTRWRRSSMDYWGQGTSTVTVSS (SEQ ID NO: 1986)
VH	EVQLQESGPPELVKPGASVKIPCKASGYTFSTDYNVDWVKQSHGKSLEWIGDIDPNNGGTIFYNQKFKGKATLTV DKSSSTAHMELRSLTSED TAVYYCVRWRSSMDYWGQGTSTVTVSS (SEQ ID NO: 1987)
VH	EVQLQESGPPELVKPGASVKIPCKASGYTFSTDYNMDWVKQSHGKSLEWIGDIDPNTGTTFYNQDFKGKATLTV DKSSSTAYMELRSLTSED TAVYYCARWRSSMDYWGQGTSLTVSS (SEQ ID NO: 1988)
VH	EVQLQESGAELVRPGASVTLSCASGYTFSTDYEMHWVKQTPVHGLEWIGVIDPGTGGTAYNQKFKVKALLTA DKSSNTAYMELRSLTSEDSAVYYCTSEKFGSNYYFDYWGQGTTLTVSS (SEQ ID NO: 1989)
VH	EVQLQESGAELVRPGASVTLSCASGYTFSTDYEIHWVKQTPVHGLEWIGVIDPETGGTAYNQKFKGKAILTA DKSSSTAYMELRSLTSEDSAVYYCTSEKFGSSYYFDYWGQGTTFVTVSS (SEQ ID NO: 1990)
VH	EVQLQESGPGLVKPSQSLSITCTVTGYSITSDYAWNWIWQFPGNKLEWMGYITYSGSPTYNPSLKSQFSITR DTSKNQFFLQLNSLTEDTATYYCARGLGGHYFDYWGQGTTLTVSS (SEQ ID NO: 1991)
VH	EVQLQESGPGLVKPSQSLSITCTVTGYSITSDYAWNWIWQFPGNKLEWMGYITYSGSPTYNPSLKSQFSITR DTSKNQFFLQLNSVTEDTATYYCARGLGGHYFDYWGQGTTLTVSS (SEQ ID NO: 1992)
VH	EVQLQESGPDLVKPSQSLSITCTVTGYSITSGYSWHWIWQFPGNKLEWMGFIHYSGDTNYPNPSLKSRSFSITR DTSKNQFFLHLNSVTPEDTATYYCASPSRLLFDYWGHTTLTVSS (SEQ ID NO: 1993)
VH	EVQLQESGPGLVAPSQSLSITCTVSGFSLTNYGVDWVRQSPGKGLEWLGVWVGSTNYSNALSRLSISKD NSKSQVFLKMNSLQTD TAMYCCASDGFVYWGQGTTLTVTVSS (SEQ ID NO: 1994)
VH	EVQLQESGPGLVAPSQSLSITCTVSGFSLTNYGVDWVRQSPGKGLEWLGVWVGIGSTNYSNALSRLSISKD NSKSQVFLKMNSLQSD TAMYCCASDGFVYWGQGTTLTVTVSS (SEQ ID NO: 1995)
VH	EVQLQESGPSLVQPSQSLSITCTVSGFSLTNYGVHWVRQSPGKGLEWLGVWVRGGNTDYNAAFMSRLSITKD NSKSQVFFKMNLSLQADDTAIYYCAASMIGGYWGQGTTLTVSS (SEQ ID NO: 1996)
VH	EVQLQESGPSLVQPSQSLSITCTVSGFSLTNYGVHWVRQSPGKGLEWLGVWVRGGNTDYNAAFMSRLSITKD NSKSQVFFKFHSLQTD DTAIYYCAASMIGGYWGQGTTLTVSS (SEQ ID NO: 1997)
VH	QVQLVQSGAEVKKPGASVKVSKASGYTFSTSYWMHWVRQAPGQGLEWMGMIDPSNSETSLNQKFQGRVTMTV DKSTNTVYMELSSSLRSED TAVYYCARSRGNYAYEMDYWGQGTTLTVTVSS (SEQ ID NO: 1998)
VH	EVQLVQSGAEVKKPGASVKVSKASGYTFSTSYWMHWVRQAPGQGLEWMGMIDPSNSETSLNQKFQGRVTNLV DKSTNTAYMELSSSLRSED TAVYYCARSRGNYAYEMDYWGQGTTLTVTVSS (SEQ ID NO: 1999)
VH	EVQLVQSGTEVTKPGASVKVSKASGYTFSTSYWMHWVRQAPGQGLEWLG MIDPSNSETTLNQKFQGRVTMTV DKSTNTVYMELTSLRSED TAVYYCARSRGNYAYEMDYWGQGTTLTVTVSS (SEQ ID NO: 2000)
VH	EVQLVQSGAEVKKPGASVKVSKASGYTFSTDYEMHWVRQAPGQGLEWMGIIDPGTGGTAYNQKFQGRVTMTA DKSTSTVYMELSSSLRSED TAVYYCTSEKFGSNYYFDYWGQGTTLTVTVSS (SEQ ID NO: 2001)

VH	EVQLVQSGAEVKKPGASVKVSCASGYTFTDYEMHWVRQAPGQGLEWMGIIDPGTGGTAYNQKFQGRVTMTA DKSTNTVYMELSSLRSED TAVYYCTSEKFGSNYYFDYWGGTGLVTVSS (SEQ ID NO: 2002)
VH	EVQLVQSGAEVKKPGASVKVSCASGYTFTDYEMHWVRQAPGQRLWGMVIDPGTGGTAYNQKFQGRVTITA DKSASTAYMELSSLRSED TAVYYCTSEKFGSNYYFDYWGGTGLVTVSS (SEQ ID NO: 2003)
VL	DIVLTQTPAIMSASPGEKVTLTCSASSSVSNLYWYQQRPGSSPKLWIYSTSNLASGVPARFSGSGSGTSY SLTISSMEAEDAASYFCHQWSSYPPTFGSGTKLEIK (SEQ ID NO: 2004)
VL	DIVITQTAIMSASPGEKVTLTCSASSSVSNLYWYQQRPGSSPKLWIYSTSNLASGVPARFSGSGSGTSY SLTISSMEAEDAASYFCHQWSSYPPTFGSGTKLEIK (SEQ ID NO: 2005)
VL	DIVMTQTPATMSASPGEKVTLTCSASSSVSNLYWYQQKPGSSPKVWIYSTSNLASGVPARFSGSGSGTSY SLTISSMEAEDAASYFCHQWSSYPPTFGSGTKLEIK (SEQ ID NO: 2006)
VL	DIVMTQTATMSASPGEKVTLTCSASSSVSNLYWYQQKPGSSPKVWIYSTSNLASGVPARFSGSGSGTSY SLTISSMEAEDAASYFCHQWSSYPPTFGSGTKLEIK (SEQ ID NO: 2007)
VL	DIVLTQSTAIMSASPGEKVTLTCSASSGVSNLYWYQQKPGSSPKLWIYSTSNLASGVPARFSGSGSGTSY SLTISSVEAEDAASYFCHQWSSYPPTFGSGTKLEIK (SEQ ID NO: 2008)
VL	DIVLTQTPSSLSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSKLPWTFGGGKLEIK (SEQ ID NO: 2009)
VL	DIVLTQSPSSLSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSKLPWTFGGGKLEIK (SEQ ID NO: 2010)
VL	DIVITQSPSSLSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2011)
VL	DIVMTQSPSSLSASLGDRVTISCSASQGISNYLNWYQQRPDGTGVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSNLPWTFGGGKLEIK (SEQ ID NO: 2012)
VL	DIVMTQSPSSLSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2013)
VL	DIVMTQSTSSLSASLGDRVTISCSASQGISHYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTIRNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2014)
VL	DIVMTQSPSSLSASLGDRVTISCSASQGISHYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTIRNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2015)
VL	DIVMTQSPSSLSASLGDRVTISCSASQDISSYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2016)
VL	DIVMTQTPSSLSASLGDRVTISCSASQGISYLNWYQQKPDGTIKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSELPWTFGGGKLEIK (SEQ ID NO: 2017)
VL	DIVMTQTPSSMSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSYLPWTFGGGKLEIK (SEQ ID NO: 2018)
VL	DIVMTQTPSSLSASLGDRVTISCSASQIGINYLNWYQQKPDGTVKLLIYYTSSLHSGVPSRFSGSGSGTDYS LTISNLEPEDIATYYCQQYSNLPWTFGGGKLEIK (SEQ ID NO: 2019)

VL	DIVMTQSPSSLSASLGDRVTISCSASQGISNYLNWYQQKPDGTVKLLIYYTSNLHSGVPSRFSGSGSGTDYS LTISDLAPEDIATYYCQQYSYLPWTFGGGKLEIK (SEQ ID NO: 2020)
VL	DIVITQSPSLSLPVGLGDQASISCRSSQSLVHSNGNTYLHWYLQKPGQSPKLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDLGVYFCSQSTHVPYTFGGGKLEIK (SEQ ID NO: 2021)
VL	DIVLTQTPLSLPVSLGDQASISCRSSQSIVHSNGNTYLEWYLQKPGQSPKLLIYKVSNRFSGVPDRFSGSGS GTNFTLKISRVEAEDLGVYYCFQGSHVPLTFGAGTKLELK (SEQ ID NO: 2022)
VL	DIVLTQSPSLSLPVSLGDQASISCRSSQSIVHSDGNTYLEWYLQKPGQSPKLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDLGIYYCFQGSHVPLTFGAGTKLELK (SEQ ID NO: 2023)
VL	DIVITQTPLSLPVSLGDQASISCRSSQTIVHSDGNTYLEWYLQKPGQSPKLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDLGVYYCFQGSHVPLTFGAGTKLEIK (SEQ ID NO: 2024)
VL	DIVMTQSTLSLPVSLGDQVSISCRSSQSIVHSDGNTYLEWYLQKPGQSPNLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDLGVYYCFQGSHVPLTFGAGTKLELK (SEQ ID NO: 2025)
VL	DIVLTQDELSNPVTSGESVSISSCRSSKSLLYKDGTKTYLNWFLQRPQGSPQVLIYFMSTRASGVSDRFSGSGS GTDFTLEISRVKAEDVGYYCQQLVDFPFTFGSGTKLELK (SEQ ID NO: 2026)
VL	DIVMTQDELYNPVTSGESVSISSCRSSKSLLYKDGTKTYLNWFLQRPQGSPQVLIYFMSTRASGVSDRFSGSGS GTDFTLEISRVKAEDVGYYCQQLVDFPFTFGSGTKLEIK (SEQ ID NO: 2027)
VL	DIVMTQSPSSLTVTAGEKVTMSCKSSQSLNSGTQKNYLTWYQQKPGQPPKLLIYWASTRESGVPDRFTGSG SGTDFTLTISVQAEDLAVYYCQNDYSYPLTFGAGTKLEIK (SEQ ID NO: 2028)
VL	DIVLTQTATLSVTPGDRVSLSCRASQSISDYLHWYQQKSHESPRLLIKYASQSIGIPSRFSGSGSGSDFT LTVNSVEPEDVGYYCQNGHSYPYTFGGGKLEIK (SEQ ID NO: 2029)
VL	DIVLTQSPDTLSVTPGDRVSLSCRASQSISDYLHWYQQKSHESPRLLIKYASQSIGIPSRFSGSGSGSDFT LSINSVEPEDVGYYCQNGHSYPYTFGGGKLELK (SEQ ID NO: 2030)
VL	EIVLTQSPATLSLSPGERATLSCRASSSVSSNYLYWYQQKPGQAPRLLIYSTSNRATGIPARFSGSGSGTDY TLTISSLEPEDFAVYYCHQWSSYPPTFGQGKLEIK (SEQ ID NO: 2031)
VL	DIVLTQSPATLSLSPGERATLSCRASSSVSSNYLYWYQQKPGQAPRLLIYSTSNLATGIPARFSGSGSGTDY TLTISSLEPEDFAVYFCHQWSSYPPTFGQGKLEIK (SEQ ID NO: 2032)
VL	DIVLTQSPGTLSSLSPGEKVTLSRASSSVSSNYLYWYQQKPGQAPRLVIYSTSNLATGIPDRFSGSGSGTDY TLTISRLEPEDFAVYFCHQWSSYPPTFGQGKVEIK (SEQ ID NO: 2034)
VL	DVMTQSPSLSLPVTLGQPASISCRSSQTIVHSDGNTYLEWYQQRPQGSPRLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDVGYYCFQGSHVPLTFGQGKLEIK (SEQ ID NO: 2035)
VL	DIVMTQSPSLSLPVTLGQPASISCRSSQTIVHSDGNTYLEWYQQRPQGQSPKLLIYKVSNRFSGVPDRFSGSGS GTDFTLKISRVEAEDVGYYCFQGSHVPLTFGQGKLEIK (SEQ ID NO: 2036)
VL	DIVMTQTPLSSPVTLGQPASISCRSSQTIVHSDGNTYLEWYQQRPQGPPRLLIYKVSNRFSGVPDRFSGSGA GTDFTLKISRVEAEDVGYYCFQGSHVPLTFGQGKLEIK (SEQ ID NO: 2037)

Note that (i) the sequences provided for Group B, Group P, Group Q, and Group V and designated as “HC” and “LC” are heavy chain and light chain amino acid sequences; all other

sequences presented in Table 7 are variable heavy chain and variable light chain sequences; (ii) the sequence provided in Group D is a fusion protein comprising a B7-DC polypeptide and an immunoglobulin polypeptide; and (iii) in Group I, the last sequence is a bispecific antibody (BsAb) sequence.

[000572] In some embodiments, the activatable anti-PD-1 antibody includes a CDR sequence shown in Table 8, a combination of VL CDR sequences (VL CDR1, VL CDR2, VL CDR3) selected from the group consisting of those combinations shown in a single row Table 8, a combination of VH CDR sequences (VH CDR1, VH CDR2, VH CDR3) selected from the group consisting of those combinations shown in Table 8, or a combination of VL CDR and VH CDR sequences (VL CDR1, VL CDR2, VL CDR3, VH CDR1, VH CDR2, VH CDR3) selected the group consisting of those combinations shown in Table 8. The CDR regions were defined according to AbM definition as described in the CDR definition table in Andrew C.R. Martin's Bioinformatics Group website at UCL.

Table 8. CDR Sequences for Antibodies and Activatable Antibodies that Bind PD-1

AB Name	VH			VL		
	CDR1 (SEQ ID NO)	CDR2 (SEQ ID NO)	CDR3 (SEQ ID NO)	CDR1 (SEQ ID NO)	CDR2 (SEQ ID NO)	CDR3 (SEQ ID NO)
M13	GFTFSGYAMS (653)	YISNSGGNAH (658)	EDYGTSPFVY (664)	RASESVDNYGIS FMN (669)	AASNQGS (678)	QQSKDVPWT (683)
M19	GYTFTDYMD (654)	YIYPKNGGSS (659)	KVVATDY (665)	KSSQSLLYSSNQ KNYL (670)	WASIRES (679)	QQCDSYPWT (684)
M3	GFTFSNYAMS (655)	YISNGGGDTH (660)	ENYGTSPFVY (666)	RASESVDNYGIS FMN (669)	AASNQGS (678)	QQSKDVPWT (683)
M5	GFSFSSYDMS (656)	TISGGGRYTY (661)	NYYGFDY (667)	KASQDVGTAVA (671)	WASTRHT (680)	QQYSSYPWT (685)
M14	GFTFSSYGMS (657)	TISGGGRDIY (662)	LYLGFDY (668)	LASQTIGTWLA (672)	AATSLAD (681)	QQLYSIPWT (686)
A	GFTFSGYAMS (653)	YISNSGGNAH (658)	EDYGTSPFVY (664)			
Ab	GFTFSGYAMS (653)	YISNSGGNAH (658)	EDYGTSPFVY (664)			
Ae	GFTFSGYAMS (653)	YISNSGGNTH (663)	EDYGTSPFVY (664)			
Af	GFTFSGYAMS (653)	YISNSGGNTH (663)	EDYGTSPFVY (664)			
Ba	GYTFTDYMD (654)	YIYPKNGGSS (659)	KVVATDY (665)			
Bb	GYTFTDYMD (654)	YIYPKNGGSS (659)	KVVATDY (665)			
C	GFTFSNYAMS (655)	YISNGGGDTH (660)	ENYGTSPFVY (666)			
Ca	GFTFSNYAMS (655)	AYISNQGGDT H (2041)	ENYGTSPFVY (666)			
D	GFSFSSYDMS (656)	TISGGGRYTY (661)	NYYGFDY (667)			
1.0				RASESVDNYGIS FMN (673)	AASNQGS (678)	QQSKDVPWT (683)

1.1				RASESVDNYGIS FMN (673)	AASNQGS (678)	QQSKDVPWT (683)
1.2				RASESVDQYGIS FMN (674)	AASNQGS (678)	QQSKDVPWT (683)
1.4				RASESVDSYGIS FMN (675)	AASNQGS (678)	QQSKDVPWT (683)
1.5				RASESVDAYGIS FMN (676)	AASNQGS (678)	QQSKDVPWT (683)
1.6				RASESVDNYGIS FMN (673)	AASDQGS (682)	QQSKDVPWT (683)
c1 1.7				RASESVDAYGIS FMN (676)	AASNQGS (678)	QQSKDVPWT (683)
1.9				RASESVDAYGIS FMN (676)	AASNQGS (678)	QQSKDVPWT (683)
1.10				RASESVDAYGIS FMN (676)	AASNQGS (678)	QQSKDVPWT (683)
2				KSSQSLLYSSNQ KNYLA (677)	WASIRES (679)	QQSDSYPWT (687)
4				KASQDVGTAVA (671)	WASTRHT (680)	QQYSSYPWT (685)

[000573] In some embodiments, the activatable anti-PD-1 antibody comprises or is derived from an antibody that is manufactured, secreted or otherwise produced by a hybridoma, such as, for example, the hybridoma(s) disclosed in US Patent No. 8,927,697 and US Patent Application Publication Nos. US 2011-0171215 and US 2015-0152180, and deposited at the European Collection of Cell Cultures (ECACC) under Accession Number 08090902; 08090903; and 08090901.

[000574] In some embodiments, the activatable anti-PD-1 antibody comprises or is derived from an antibody that is manufactured, secreted or otherwise produced by a hybridoma, such as, for example, the hybridoma(s) disclosed in US Patent Application Publication No. US 2014-0335093, and deposited at the Collection Nationale De Cultures De Microorganismes (CNCM) under deposit number 1-4122.

[000575] In some embodiments, the activatable anti-PD-1 antibody comprises or is derived from an antibody that is manufactured, secreted or otherwise produced by a hybridoma, such as,

for example, the hybridoma(s) disclosed in PCT Publication Nos. WO 2015/058573 or WO 2014/206107, and deposited at the China Culture Collection Committee General Microbiology Center under Accession no. 8351.

[000576] In some embodiments, the activatable anti-PD-1 antibody comprises or is derived from an antibody that is manufactured, secreted or otherwise produced by a hybridoma, such as, for example, the hybridoma(s) disclosed in US Patent Application Publication No. US2011177088, and deposited at the Collection Nationale De Cultures De Microorganismes (CNCM) under deposit number 1-3745.

[000577] In some embodiments, the activatable anti-PD-1 antibody includes a CDR sequence shown in Table 9, a combination of VL CDR sequences selected from the group consisting of those combinations shown in Table 9, and/or a combination of VH CDR sequences selected from the group consisting of those combinations shown in Table 9.

[000578] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group A in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group A in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group A in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group A in Table 9.

[000579] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group B in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group B in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group B in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group B in Table 9.

[000580] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group C in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group C in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group C in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group C in Table 9.

[000581] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group E in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group E in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group E in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group E in Table 9.

[000582] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group F in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group F in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group F in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group F in Table 9.

[000583] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group G in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the

combinations shown in Group G in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group G in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group G in Table 9.

[000584] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group H in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group H in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group H in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group H in Table 9.

[000585] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group L in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group L in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group L in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group L in Table 9.

[000586] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group M in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group M in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group M in Table 9, and a combination of light chain

CDR sequences selected from the group consisting of the combinations shown in Group M in Table 9.

[000587] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group N in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group N in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group N in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group N in Table 9.

[000588] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group O in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group O in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group O in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group O in Table 9.

[000589] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group P in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group P in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group P in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group P in Table 9.

[000590] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in

Group Q in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group Q in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group Q in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group Q in Table 9.

[000591] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group R in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group R in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group R in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group R in Table 9.

[000592] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group T in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group T in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group T in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group T in Table 9.

[000593] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group U in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group U in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group

consisting of the combinations shown in Group U in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group U in Table 9.

[000594] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group V in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group V in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group V in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group V in Table 9.

[000595] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group W in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group W in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group W in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group W in Table 9.

[000596] In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group X in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group X in Table 9. In some embodiments, the activatable anti-PD-1 antibody includes a combination of heavy chain CDR sequences selected from the group consisting of the combinations shown in Group X in Table 9, and a combination of light chain CDR sequences selected from the group consisting of the combinations shown in Group X in Table 9.

Table 9. Additional CDR Sequences for Antibodies and Activatable Antibodies that Bind PD-1

Group A					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
NYGMH (688)	VIWYDGSNKYYADSVKG (694)	NDDY (700)	RASQSVSSYLA (704)	DASNRAT (706)	QQRSNWPLT (710)
NYGFH (689)	VIWYDGSKKYYADSVKG (695)	GDDY (701)	RASQSVSSYLA (704)	DTSNRAT (707)	QQRSNWPLT (710)
NYGMH (688)	LIWYDGSNKYYADSVKG (696)	NVDH (702)	RASQSVSSYLA (704)	DASNRAT (706)	QQSSNWPRP (711)
NSGMH (690)	VIWYDGSKRYYADSVKG (697)	NDDY (700)	RASQSVSSYLA (704)	DASNRAT (706)	QQSSNWPRP (711)
RSSFFWG (691)	SIYYSGSTYYNPSLKS (698)	DYDILTGEDDY (703)	RASQGISSWLA (705)	AASNLR (708)	QQYYSYPRT (712)
SYGFH (692)	VIWYDGSKKYYADSVKG (695)	GDDY (701)	RASQSVSSYLA (704)	DASNRAT (706)	QQRSNWPLT (710)
RSSYFWG (693)	SIFYSGETYFNPSLKS (699)	DYDILTGEDDY (703)	RASQGISSWLA (705)	AASSLQS (709)	QQYYSYPRT (712)
Group B					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
SYLY (713)	GVNPSNGGTNFSEKFK S (715)	RDSNYDGGFDY (717)	RASKSVSTSGFSY LH (719)	LASNLES (721)	QHSWELPLT (723)
NYMY (714)	GINPSNGGTNFNEKFK N (716)	RDYRFDMGFDY (718)	RASKGVSTSGYSY LH (720)	LASYLES (722)	QHSRDLPLT (724)
AASGFTFS SYAMS (627)	TITGGGRNTYYPDSVK G (817)	QGYDGYTFAY (823)	RASESVDNSGISF MN (830)	AASNPGS (836)	QQSKEVPWT (840)
AASGFTFS SFGMS (811)	TISGGGSNTYYPDSVK (818)	IYDVAWFAY (824)	RSSQTIVHSDGNT YLE (831)	AASNQGS (678)	FQGSHPYPT (841)
KSGYSFT DYALH (812)	VISTHYGDTVYNQRFK G (832)	EGYGSIFYFDQ (825)	RASESVDSYGNSF MN (832)	KVSNRFS (837)	QQNNEVPLT (842)
KASGYAFT SYNIY (813)	YIDLINGDTSYNEKF (819)	EGRLSFDY (826)	RSSQSIVQSNGNT YLE (833)	LASNLD (838)	FQGSHPYPT (843)
AASGFTFS NYDMS (814)	YISGGGNTYYPDTL (820)	ISLTGIFYDY (827)	RSSQTIVHGNGNT YLE (834)	SASTLAS (839)	QQGFGTSNVE NP (844)
AASGFTFN SYGMS (815)	TISGGGSYTYYPDSVQ G (821)	GNVYVMDY (828)	QASENIYSSLA (835)		
KSGYIFT DYVMH (816)	VISTYYSNINYNQKFK G (822)	EGFGRPYWYFD V (829)			
Group C					

VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GYSITSDY AWN (845)	YINYSGSTSYNPSLKS (847)	WIGSSAWYFDV (849)	RSGQNIVHSNGNTY LE (851)	KVSNRFF (853)	FQGSHPVFT (855)
GYTFTTY LY (846)	GINPSNGGTNFNEKFK (848)	RDYRYDRGFDY (850)	RASKSVSTSGFNYY H (852)	LASNLES (721)	QHSRELPLT (856)
Group E					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
SYGIS (857)	WISAYNGNTNYAQ KLQG (861)	DADYSSGSGY (865)	SGDALPKQYAY (869)	KDTERPS (872)	QSADNSITYR V (876)
SYIYH (858)	IINPRGATISYAO KFQG (862)	AGIYGDFDY (866)	TGTSNDVGGYNYVS (870)	DVTNRPS (873)	SSYTIVTNFE VL (877)
SGAYYWS (859)	YIYNGNTYNNPS LRS (863)	ASDYVWGGYRYM DAFDI (867)	SGSNSNIGNSNVN (871)	GNNQRPS (874)	AAWDDSLNGP V (878)
SSYWMS (860)	AISGSGGSTYYAD SVKG (864)	ENWGSYFDL (868)	RASQGISSWLA (705)	KASTLES (875)	QQSYSTPWT (879)
Group F					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
SSWIH (880)	YIYPSTGFTEYNQ KFKD (881)	WRDSSGYHAMDY (882)	RASQSVSTSGYSYMH (883)	FGSNLES (884)	QHSWEIPYT (885)
Group L					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GFTFSSYD YYMH (886)	VIWYSGSNTYYND SLKS (887)	AYFGVDV (888)	RASQGIGNTLA (889)	RASQGIG NTLA (889)	QQYDHPVLT (891)
Group G					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GFSLTSYG VH (892)	VIWAGGSTNYNSA LMS (317)	GFSLTSYGVH (896)	KASQSVSNDVA (898)	YAFHRFT (900)	
VIWAGGST NYPNLSKS (893)	ARAYGNYWYIDV (897)	ARAYGNYWYIDV (897)	KSSSVSNDVA (899)		
VIWAGGST NYPNLSKS (894)	HQAYSSPYT (895)				

Group H					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
DYEMH (901)	VIESETGGTAYNQ KFKG (902)	EGITTVATTTYW YFDV (903)	RSSQSIVHSNGNTYLE (904)	KVSNRFS (905)	FQGSHPVPLT (906)
Group M					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GYTFTTYP IE (907)	NFHYPNDDTKYNE KFK (908)		RASSSVISSYLH (909)	STSNLAS (910)	QQYNGYPLT (911)
					QQYNSYPLT (912)
Group N					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
SYLY (713)	GVNPSNGGTNFSE KFK (1596)	RDSNYDGGFDY (717)	RASKSVSTSGFSYLH (719)	LASNLES (721)	QHSWELPLT (723)
Group P					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GYTFTTY (854)	YPGTGG (1597)	WTTGTGAY (1598)	QSLLDSGNQKNF (1599)	WAS (1600)	DYSYPY (1601)
TYWMH (1602)	NIYPGTGGSNFDE KFK (1603)		KSSQSLLDSGNQKNFL (1833)	WASTRES (1604)	QNDYSYPYT (1605)
GYTFTTYW MH (1606)					
Group Q					
VH			VL		
CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)	CDR1 (SEQ ID NO:)	CDR2 (SEQ ID NO:)	CDR3 (SEQ ID NO:)
GFTFSSYT (1607)	ISDTGGNT (1608)	AKDQGGSYPPYF HY (1609)	QSISSY (1610)	AAS (1611)	QQSYSTPPIT (1612)
GFTVSNNY (1613)	IYSGGFT (1614)	ARYYYDTSYWT FFDY (1615)	QSVSSN (1616)	GAS (1617)	QQYNNWPLT (1618)
GFTFSSYG (1619)	IWYDGSNI (1620)	ARPGHWNYFFEY (1621)	QSINNY (1622)	TAS (1623)	QQSYSTPPLT (1624)
GFTFRNYD (1625)	IGSAGDT (1626)	TRDIHCSSTRCY GMDV (1627)	QSISSY (1628)	AAS (1611)	QQSYSTPPIT (1612)
GFKFSNEW (1629)	IKSKTDGGTT (1630)	TTDQDFWSGYT GADYYGMDV (1631)		AAS (1611)	QQSYSTPPIT (1612)
GGSLNGYY (1632)	IDHSGST (1633)	AREGLLPFDY (1634)	QSVYSNY (1635)	AAS (1611)	HQYATSPWT (1636)

DDSISSSTT YY (1637)	MSYNGNN (622)	ARHLGYNGNWYP FDF (1638)	QSVSSSY (1639)	GAS (1640)	QQYGSSPWT (1641)
GFTFDDYA (1642)	INWSGNNI (1643)	TKDISITGTDLDA FDV (1644)	QGISNW (1645)	SAS (1646)	QQAHSFPLT (1647)
GFTFSSYG (1619)	IWSDGDSE (1648)	ARDRDLEDI (1649)	QGIRND (1650)	AAS (1611)	LQHNSYPLT (1651)
GFTFSNFG (1652)	ISGGGRDT (1653)	VKWGNIYFDY (1654)	LSINTF (1655)	AAS (1611)	QQSSNTPFT (1656)
GFTFDDYG (1657)	ISWNDGKT (1658)	ARDWQYLIERYF DY (1659)	QSVSSSY (1639)	GAS (1640)	QQYGSSPWT (1641)
GFTFDDYG (1657)	IGWTGGRS (1660)	ARDRQWLQWYF DY (1661)	QSISSY (1610)	AAS (1611)	QQSYSTPPIT (1612)
GFTFDDYG (1657)	ISWNDGKT (1658)	ARDWQYLIDRYF DF (1662)			
GFTFDDYA (1642)	IGWSSGSI (1663)	AKAYTFMITLYF DY (1664)			
GFTFDDYD (1665)	SGWNRGSL (1666)	AKGFVVVSAAYF DY (1667)			
GVTFRNFA (1668)	IIPFFSAA (1669)	AREGERGHTYGF DY (1670)			
GFTFDDYA (1642)	INWNRGRT (1671)	AKAEQWLDEGYF DY (1672)			
GFTFEDYA (1673)	ISDSGGST (1674)	AKDQGGSYPPYF HY (1609)			
GYTFTSFY (1675)	IGWSNVKI (1676)	VKAYTSMITLYF DY (1677)			
GGSISSYH (1678)	INPSDGST (1679)	ARRVAGDIFDI (1680)			
GFTFEDYG (1681)	IYYIGST (1682)	ARVPVGATGASD V (1683)			
GFTFSAYA (1684)	ISWTGGNT (1685)	TRDRQWLMQWYF DY (1686)			
GFALHDYA (1687)	ISYGGSDK (1688)	AKSAHNWFFFDY (1689)			
GDSISSTA YH (1140)	ISWNSGVI (1717)	AKGSGSYVSWF DP (1718)			
GFTFSTYA (1690)	ITYNGNT (1691)	ARHLGYNSDFFP FDF (1692)			
	IGSGDST (1693)	VKVRNYDGSFDI (1694)			
Group R					
VH			VL		
SYGMS (1751)	TMSGGGRDIYYP SMKG (1752)	QYYDDWFAY (1753)	RASQISNNLH (1754)	YASQIS (1755)	QQNSWPLT (1756)
TYGVH (1757)	VIWGGSTDYNA FIS (1758)	EKSVYGNVYGAM DY (1759)	KASQSVDDVA (1760)	YAFNRYT (1761)	QQDYRSPWT (1762)
SYGMS (1751)	TISGGGRDIYYP SVKG (1763)	QYYDDWFAY (1753)	RASQISNDLH (1764)	YVSQIS (1765)	QQSDSWPLT (1766)
SDYAWN (1767)	YISYSGYTSYNPS LKS (1768)	SLDYDYGTMDY (1769)	RANSSVSSMH (1770)	AINLAF (1771)	QQWSSRPPT (1772)
SYDMS (1773)	TISGGGSYTYQD SVKG (1774)	PYGPYFDY (1775)	HASQINWLS (1776)	ASNLT (1777)	QQGQSYPT (1778)

SDYAWN (1767)	YISYSGYTSYNPS LKS (1768)	SLDYDYGTMDY (1769)	RANSSVSSMH (1770)	AINLAF (1771)	QQWNSRPPT (1779)
YYDMS (1780)	TISGGGRNTYFID SVKG (1781)	PYEGAVDF (1782)	KASQDVNDVA (1783)	WASTRHH (1784)	QQYSTFPYT (1785)
SYGMS (1751)	TISGGGRDYYLD SVKG (1786)	QYYDDWFAY (1753)	RASQSLSNLH (1787)	YASQSIG (1755)	QQSNSWPLT (1756)
NNWIG (1788)	DFYPGGGYTNYNE KFKG (1789)	GYGTNYWYFDV (1790)	KASQSVSNDVA (1791)	YAFTRYI (1792)	QQDYSSPYT (1793)
NFGMN (1794)	WISGYTREPTYAA DFKG (1795)	DVFDY (1796)	RASESVDNYGYSFMN (1797)	RASNLES (1798)	QQSNADPT (1799)
Group T					
VH			VL		
GYTFTSY MY (1800)	GVNPSNGGTNFNE KFKS (1801)	RDYRYDMGFDY (1802)	RASKGVSTSGYSYLH (1803)	LASYLE (1804)	QHSRELPLT (1805)
GYTFTNY MY (1806)	GINPSNGGTNYNE KFKN (890)	RDYRYDMGFDY (1802)			
Group U					
VH			VL		
G (T/R/ I) (F / L) (S/ E/ T/ P/ R) (T/ S/ H/Q/R/ W) (F/Y/ Q) (1881)	W (I/V) S A (Y/H) N G N T (K/N) Y A Q K L Q G (1882)	(Q/-) GY (G/D) (N/V) Y (L/S) (Y/W) (D/A) (Y/V) (1883)	S G D A L (P/T/S) (M/T/E/K) Q Y (G/A) Y (1884)		Q Q (N/S/W) (Y/K/I) (N/E/S) (S/V/D/T) P (Y/W) T (1885)
(T/R/W/Q /H/S) (Y/F/ Q) G (M/I) (1886)	(T/A) I S G (S/G) G (S/G) (S/D/N) T Y Y (A/P/S) D S V (K/Q) G (1887)	D (A/V/S) (D/E) Y (S/G/R) (S/L/ T) (1888), with the proviso that if A at position 2, then not D or S in position 5			
		GYALDY (2038)			
GYIFSSY (1889)	FPGSGS (1890)	GYGNYLYFDV (1891)	KASQSVSDDVA (1760)	YAFKRYI (1892)	QQNYNSPYT (1893)
SYWIG (1894)	KIFPGSGSADYNE NFKG (1895)	DSEYSSSGSY (1896)	SGDALTTQYAY (1897)	KDTERPS (872)	QSADNSITYR V (876)
		QRDSAWFAS (2039)	RASESVDNSGISFMS (1898)	ATSKLAS (1899)	QQWISDPWT (1900)
			SGDALSEQYAY (1901)	TASNQGS (1902)	QQSKEVPWT (840)

			SGDALPKQYAY (869)		
			SGDALPMQYGY (1903)		
			RTSSSVNYMH (1904)		
GYRFTWY (1905)	SAYNGN (1906)	DVDYSSGSGY (1907)			
WYGIS (1908)	WISAYNGNTNYAQ KLQG (861)	DAEYSLGSGY (1909)			
GYRFSTF (1910)	SAHNGN (1911)	DAEYGS GSGY (1912)			
TFGIS (1913)	WVSAHNGNTNYAQ KLQG (1914)	DADYGS GSGY (1915)			
GYRFETY (1916)	SYSGR (1917)	DVDYGTGSGY (1918)			
GYRFRQY (1919)	YISYSGRTSYNPS LTS (1920)	DVDYGS GSGY (1921)			
QYGIS (1922)	SGGSD (1923)	DAEYGS GSGY (1912)			
GYRFTRY (1924)	TISGGGSDTYYPD SVQG (1925)	GYALDY (2038)			
RYGIS (1926)		QGYDVYSWFAY (1927)			
GYRFPHY (1928)					
HYGIS (1929)					
GYRFTRQ (1930)					
RQGIS (1931)					
GHSITS DY (1932)					
SDYAWN (1767)					
GFTFSTF (1933)					
TFGMS (1934)					
Group V					
VH			VL		
GWSLTGPG (1935)	IYGDGST (1936)	AYEYAMDW (1937)	WSVSTSGKSY (1938)	LLS (1939)	YHIRDLT (2040)
Group W					
VH			VL		
GYTFTSY IN (1940)	YPGSSL (1941)	LSTGT Fay (1942)	KSSQSLWDSTNQNFL T (1943)	WTSTRES (1944)	QNDYFYPLT (1945)
GYTFTSY (1946)	NIYPGSSLTNYNE KFKN (1947)	LTTGT Fay (1948)	KSSQSLWDSTNQNFL T (1949)	WTSYRES (1950)	QNDYFY PHT (1951)

SYWIN (1952)	YPGSSI (1953)	LLTGTFAY (1954)	KSSQSLLDSGNQKNFL T (1955)	WTS (T/Y) RES (1956)	QNDYSYPLT (1957)
	NIYPGSSITNYNE KFKN (1958)	L(L/S)TGTFAY (1959)	KSSQSL(W/L)DS(G/ T)NQKNFLT (1960)		QNDY(F/S)Y P(L/H)T (1961)
	WPGSSL (1962)				
	NIWPGSSLTNYNE KFKN (1963)				
	NIYPGSSSTNYNE KFKN (1964)				
	NI(Y/W)PGSS (L/I/S)TNYNEK FKN (1965)				
Group X					
VH			VL		
GLTFSSSG (1966)	IWYDGSKR (1706)	ATNNDY (1967)	RASQSVSSYLA (704)	TASNRAT (1968)	QQYSNWPRT (1969)

[000597] The ABs in the activatable antibodies of the disclosure specifically bind a PD-1 target, such as, for example, mammalian PD-1. In some embodiments, such Abs bind mammalian PD-1. In some embodiments, such Abs bind human PD-1. In some embodiments, such Abs bind non-human primate PD-1. Also included in the disclosure are ABs that bind to the same PD-1 epitope as an antibody of the disclosure and/or an activated activatable antibody described herein. Also included in the disclosure are ABs that compete with an anti-PD-1 antibody and/or an activated anti-PD-1 activatable antibody described herein for binding to a PD-1 target, e.g., human PD-1. Also included in the disclosure are ABs that cross-compete with an anti-PD-1 antibody and/or an activated anti-PD-1 activatable antibody described herein for binding to a PD-1 target, e.g., human PD-1.

[000598] The activatable anti-PD-1 antibodies provided herein include a masking moiety. In some embodiments, the masking moiety is an amino acid sequence that is coupled or otherwise attached to the anti-PD-1 antibody and is positioned within the activatable anti-PD-1 antibody construct such that the masking moiety reduces the ability of the anti-PD-1 antibody to specifically bind PD-1. Suitable masking moieties are identified using any of a variety of known techniques. For example, peptide masking moieties are identified using the methods described in PCT Publication No. WO 2009/025846 by Daugherty et al., the contents of which are hereby incorporated by reference in their entirety.

[000599] The activatable anti-PD-1 antibodies provided herein include a cleavable moiety. In some embodiments, the cleavable moiety includes an amino acid sequence that is a substrate for a protease, usually an extracellular protease. Suitable substrates are identified using any of a variety of known techniques. For example, peptide substrates are identified using the methods described in U.S. Patent No. 7,666,817 by Daugherty et al.; in U.S. Patent No. 8,563,269 by Stagliano et al.; and in PCT Publication No. WO 2014/026136 by La Porte et al., the contents of each of which are hereby incorporated by reference in their entirety. (*See also* Boulware et al. “Evolutionary optimization of peptide substrates for proteases that exhibit rapid hydrolysis kinetics.” *Biotechnol Bioeng.* 106.3 (2010): 339-46).

[000600] Exemplary substrates include but are not limited to substrates cleavable by one or more of the following enzymes or proteases listed in Table 3.

Table 3: Exemplary Proteases and/or Enzymes

ADAMS, ADAMTS, <i>e.g.</i> ADAM8 ADAM9 ADAM10 ADAM12 ADAM15 ADAM17/TACE ADAMDEC1 ADAMTS1 ADAMTS4 ADAMTS5	Cysteine proteinases, <i>e.g.</i> , Cruzipain Legumain Otubain-2	Serine proteases, <i>e.g.</i> , activated protein C Cathepsin A Cathepsin G Chymase coagulation factor proteases (<i>e.g.</i> , FVIIa, FIXa, FXa, FXIa, FXIIa)
		Elastase
		Granzyme B
		Guanidinobenzoatase
		HtrA1
Aspartate proteases, <i>e.g.</i> , BACE Renin	KLKs, <i>e.g.</i> , KLK4 KLK5 KLK6 KLK7 KLK8 KLK10 KLK11 KLK13 KLK14	Human Neutrophil Elastase Lactoferrin
		Marapsin NS3/4A PACE4 Plasmin PSA tPA Thrombin Trypsin uPA
Aspartic cathepsins, <i>e.g.</i> , Cathepsin D Cathepsin E	Metallo proteinases, <i>e.g.</i> , Meprin Neprilysin PSMA BMP-1	Type II Transmembrane Serine Proteases (TTSPs), <i>e.g.</i> , DESC1
Caspases, <i>e.g.</i> , Caspase 1 Caspase 2 Caspase 3 Caspase 4 Caspase 5 Caspase 6	MMPs, <i>e.g.</i> , MMP1 MMP2 MMP3 MMP7	

Caspase 7	MMP8	DPP-4
Caspase 8	MMP9	FAP
Caspase 9	MMP10	Hepsin
Caspase 10	MMP11	Matriptase-2
Caspase 14	MMP12	MT-SP1/Matriptase
	MMP13	TMPRSS2
Cysteine cathepsins, <i>e.g.</i> ,	MMP14	TMPRSS3
Cathepsin B	MMP15	TMPRSS4
Cathepsin C	MMP16	
Cathepsin K	MMP17	
Cathepsin L	MMP19	
Cathepsin S	MMP20	
Cathepsin V/L2	MMP23	
Cathepsin X/Z/P	MMP24	
	MMP26	
	MMP27	

[000601] The activatable anti-PD-1 antibodies described herein overcome a limitation of antibody therapeutics, particularly antibody therapeutics that are known to be toxic to at least some degree *in vivo*. Target-mediated toxicity constitutes a major limitation for the development of therapeutic antibodies. The activatable anti-PD-1 antibodies provided herein are designed to address the toxicity associated with the inhibition of the target in normal tissues by traditional therapeutic antibodies. These activatable anti-PD-1 antibodies remain masked until proteolytically activated at the site of disease. Starting with an anti-PD-1 antibody as a parental therapeutic antibody, the activatable anti-PD-1 antibodies of the invention were engineered by coupling the antibody to an inhibitory mask through a linker that incorporates a protease substrate.

[000602] When the AB is modified with a MM and is in the presence of the target, specific binding of the AB to its target is reduced or inhibited, as compared to the specific binding of the AB not modified with an MM or the specific binding of the parental AB to the target.

[000603] The K_d of the AB modified with a MM towards the target is at least 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 50,000, 100,000, 500,000, 1,000,000, 5,000,000, 10,000,000, 50,000,000 or greater, or between 5-10, 10-100, 10-1,000, 10-10,000, 10-100,000, 10-1,000,000, 10-10,000,000, 100-1,000, 100-10,000, 100-100,000, 100-1,000,000, 100-10,000,000, 1,000-10,000, 1,000-100,000, 1,000-1,000,000, 1000-10,000,000, 10,000-100,000, 10,000-1,000,000, 10,000-10,000,000, 100,000-1,000,000, or 100,000-10,000,000 times greater

than the K_d of the AB not modified with an MM or of the parental AB towards the target.

Conversely, the binding affinity of the AB modified with a MM towards the target is at least 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 50,000, 100,000, 500,000, 1,000,000, 5,000,000, 10,000,000, 50,000,000 or greater, or between 5-10, 10-100, 10-1,000, 10-10,000, 10-100,000, 10-1,000,000, 10-10,000,000, 100-1,000, 100-10,000, 100-100,000, 100-1,000,000, 100-10,000,000, 1,000-10,000, 1,000-100,000, 1,000-1,000,000, 1000-10,000,000, 10,000-100,000, 10,000-1,000,000, 10,000-10,000,000, 100,000-1,000,000, or 100,000-10,000,000 times lower than the binding affinity of the AB not modified with an MM or of the parental AB towards the target.

[000604] In some embodiments, the dissociation constant (K_d) of the MM towards the AB is generally greater than the K_d of the AB towards the target. The K_d of the MM towards the AB can be at least 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 100,000, 1,000,000 or even 10,000,000 times greater than the K_d of the AB towards the target. Conversely, the binding affinity of the MM towards the AB is generally lower than the binding affinity of the AB towards the target. The binding affinity of MM towards the AB can be at least 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 100,000, 1,000,000 or even 10,000,000 times lower than the binding affinity of the AB towards the target.

[000605] In some embodiments, the dissociation constant (K_d) of the MM towards the AB is approximately equal to the K_d of the AB towards the target. In some embodiments, the dissociation constant (K_d) of the MM towards the AB is no more than the dissociation constant of the AB towards the target.

[000606] In some embodiments, the dissociation constant (K_d) of the MM towards the AB is less than the dissociation constant of the AB towards the target.

[000607] In some embodiments, the dissociation constant (K_d) of the MM towards the AB is greater than the dissociation constant of the AB towards the target.

[000608] In some embodiments, the MM has a K_d for binding to the AB that is no more than the K_d for binding of the AB to the target.

[000609] In some embodiments, the MM has a K_d for binding to the AB that is no less than the K_d for binding of the AB to the target.

[000610] In some embodiments, the MM has a K_d for binding to the AB that is approximately equal to the K_d for binding of the AB to the target.

[000611] In some embodiments, the MM has a K_d for binding to the AB that is less than the K_d for binding of the AB to the target.

[000612] In some embodiments, the MM has a K_d for binding to the AB that is greater than the K_d for binding of the AB to the target.

[000613] In some embodiments, the MM has a K_d for binding to the AB that is no more than 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, or 1,000 fold greater than the K_d for binding of the AB to the target. In some embodiments, the MM has a K_d for binding to the AB that is between 1-5, 2-5, 2-10, 5-10, 5-20, 5-50, 5-100, 10-100, 10-1,000, 20-100, 20-1000, or 100-1,000 fold greater than the K_d for binding of the AB to the target.

[000614] In some embodiments, the MM has an affinity for binding to the AB that is less than the affinity of binding of the AB to the target.

[000615] In some embodiments, the MM has an affinity for binding to the AB that is no more than the affinity of binding of the AB to the target.

[000616] In some embodiments, the MM has an affinity for binding to the AB that is approximately equal of the affinity of binding of the AB to the target.

[000617] In some embodiments, the MM has an affinity for binding to the AB that is no less than the affinity of binding of the AB to the target.

[000618] In some embodiments, the MM has an affinity for binding to the AB that is greater than the affinity of binding of the AB to the target.

[000619] In some embodiments, the MM has an affinity for binding to the AB that is 2, 3, 4, 5, 10, 25, 50, 100, 250, 500, or 1,000 less than the affinity of binding of the AB to the target. In some embodiments, the MM has an affinity for binding to the AB that is between 1-5, 2-5, 2-10, 5-10, 5-20, 5-50, 5-100, 10-100, 10-1,000, 20-100, 20-1000, or 100-1,000 fold less than the affinity of binding of the AB to the target. In some embodiments, the MM has an affinity for binding to the AB that is 2 to 20 fold less than the affinity of binding of the AB to the target. In some embodiments, a MM not covalently linked to the AB and at equimolar concentration to the AB does not inhibit the binding of the AB to the target.

[000620] When the AB is modified with a MM and is in the presence of the target specific binding of the AB to its target is reduced or inhibited, as compared to the specific binding of the AB not modified with an MM or the specific binding of the parental AB to the target. When compared to the binding of the AB not modified with an MM or the binding of the parental AB

to the target the AB's ability to bind the target when modified with an MM can be reduced by at least 50%, 60%, 70%, 80%, 90%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% and even 100% for at least 2, 4, 6, 8, 12, 28, 24, 30, 36, 48, 60, 72, 84, or 96 hours, or 5, 10, 15, 30, 45, 60, 90, 120, 150, or 180 days, or 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 months or more when measured *in vivo* or in an *in vitro* assay.

[000621] The MM inhibits the binding of the AB to the target. The MM binds the antigen binding domain of the AB and inhibits binding of the AB to the target. The MM can sterically inhibit the binding of the AB to the target. The MM can allosterically inhibit the binding of the AB to its target. In these embodiments when the AB is modified or coupled to a MM and in the presence of target there is no binding or substantially no binding of the AB to the target, or no more than 0.001%, 0.01%, 0.1%, 1%, 2%, 3%, 4%, 5%, 6%, 7%, 8%, 9%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, or 50% binding of the AB to the target, as compared to the binding of the AB not modified with an MM, the parental AB, or the AB not coupled to an MM to the target, for at least 2, 4, 6, 8, 12, 28, 24, 30, 36, 48, 60, 72, 84, or 96 hours, or 5, 10, 15, 30, 45, 60, 90, 120, 150, or 180 days, or 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 months or longer when measured *in vivo* or in an *in vitro* assay.

[000622] When an AB is coupled to or modified by a MM, the MM 'masks' or reduces or otherwise inhibits the specific binding of the AB to the target. When an AB is coupled to or modified by a MM, such coupling or modification can effect a structural change that reduces or inhibits the ability of the AB to specifically bind its target.

[000623] An AB coupled to or modified with an MM can be represented by the following formulae (in order from an amino (N) terminal region to carboxyl (C) terminal region:

(MM)-(AB)

(AB)-(MM)

(MM)-L-(AB)

(AB)-L-(MM)

where MM is a masking moiety, the AB is an antibody or antibody fragment thereof, and the L is a linker. In many embodiments, it can be desirable to insert one or more linkers, *e.g.*, flexible linkers, into the composition so as to provide for flexibility.

[000624] In certain embodiments, the MM is not a natural binding partner of the AB. In some embodiments, the MM contains no or substantially no homology to any natural binding

partner of the AB. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% similar to any natural binding partner of the AB. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 25% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 50% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 20% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 10% identical to any natural binding partner of the AB.

[000625] In some embodiments, the activatable antibodies include an AB that is modified by an MM and also includes one or more cleavable moieties (CM). Such activatable antibodies exhibit activatable/switchable binding, to the AB's target. Activatable antibodies generally include an antibody or antibody fragment (AB), modified by or coupled to a masking moiety (MM) and a modifiable or cleavable moiety (CM). In some embodiments, the CM contains an amino acid sequence that serves as a substrate for at least one protease.

[000626] The elements of the activatable antibodies are arranged so that the MM and CM are positioned such that in a cleaved (or relatively active) state and in the presence of a target, the AB binds a target while the activatable antibody is in an uncleaved (or relatively inactive) state in the presence of the target, specific binding of the AB to its target is reduced or inhibited. The specific binding of the AB to its target can be reduced due to the inhibition or masking of the AB's ability to specifically bind its target by the MM.

[000627] The K_d of the AB modified with a MM and a CM towards the target is at least 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 50,000, 100,000, 500,000, 1,000,000, 5,000,000, 10,000,000, 50,000,000 or greater, or between 5-10, 10-100, 10-1,000, 10-10,000, 10-100,000, 10-1,000,000, 10-10,000,000, 100-1,000, 100-10,000, 100-100,000, 100-1,000,000, 100-10,000,000, 1,000-10,000, 1,000-100,000, 1,000-1,000,000, 1,000-10,000,000, 10,000-100,000, 10,000-1,000,000, 10,000-10,000,000, 100,000-1,000,000, or 100,000-10,000,000 times greater than the K_d of the AB not modified with an MM and a CM or of the parental AB towards the target. Conversely, the binding affinity of the AB modified with a MM and a CM towards the target is at least 5, 10, 25, 50, 100, 250, 500, 1,000, 2,500, 5,000, 10,000, 50,000, 100,000, 500,000, 1,000,000, 5,000,000, 10,000,000, 50,000,000 or greater, or between 5-10, 10-

100, 10-1,000, 10-10,000, 10-100,000, 10-1,000,000, 10-10,000,000, 100-1,000, 100-10,000, 100-100,000, 100-1,000,000, 100-10,000,000, 1,000-10,000, 1,000-100,000, 1,000-1,000,000, 1000-10,000,000, 10,000-100,000, 10,000-1,000,000, 10,000-10,000,000, 100,000-1,000,000, or 100,000-10,000,000 times lower than the binding affinity of the AB not modified with an MM and a CM or of the parental AB towards the target.

[000628] When the AB is modified with a MM and a CM and is in the presence of the target but not in the presence of a modifying agent (for example at least one protease), specific binding of the AB to its target is reduced or inhibited, as compared to the specific binding of the AB not modified with an MM and a CM or of the parental AB to the target. When compared to the binding of the parental AB or the binding of an AB not modified with an MM and a CM to its target, the AB's ability to bind the target when modified with an MM and a CM can be reduced by at least 50%, 60%, 70%, 80%, 90%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% and even 100% for at least 2, 4, 6, 8, 12, 28, 24, 30, 36, 48, 60, 72, 84, or 96 hours or 5, 10, 15, 30, 45, 60, 90, 120, 150, or 180 days, or 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12 months or longer when measured *in vivo* or in an *in vitro* assay.

[000629] As used herein, the term cleaved state refers to the condition of the activatable antibodies following modification of the CM by at least one protease. The term uncleaved state, as used herein, refers to the condition of the activatable antibodies in the absence of cleavage of the CM by a protease. As discussed above, the term "activatable antibodies" is used herein to refer to an activatable antibody in both its uncleaved (native) state, as well as in its cleaved state. It will be apparent to the ordinarily skilled artisan that in some embodiments a cleaved activatable antibody can lack an MM due to cleavage of the CM by protease, resulting in release of at least the MM (*e.g.*, where the MM is not joined to the activatable antibodies by a covalent bond (*e.g.*, a disulfide bond between cysteine residues).

[000630] By activatable or switchable is meant that the activatable antibody exhibits a first level of binding to a target when the activatable antibody is in a inhibited, masked or uncleaved state (*i.e.*, a first conformation), and a second level of binding to the target in the uninhibited, unmasked and/or cleaved state (*i.e.*, a second conformation), where the second level of target binding is greater than the first level of binding. In general, the access of target to the AB of the activatable antibody is greater in the presence of a cleaving agent capable of cleaving the CM, *i.e.*, a protease, than in the absence of such a cleaving agent. Thus, when the activatable antibody

is in the uncleaved state, the AB is inhibited from target binding and can be masked from target binding (*i.e.*, the first conformation is such the AB cannot bind the target), and in the cleaved state the AB is not inhibited or is unmasked to target binding.

[000631] The CM and AB of the activatable antibodies are selected so that the AB represents a binding moiety for a given target, and the CM represents a substrate for a protease. In some embodiments, the protease is co-localized with the target at a treatment site or diagnostic site in a subject. As used herein, co-localized refers to being at the same site or relatively close nearby. In some embodiments, a protease cleaves a CM yielding an activated antibody that binds to a target located nearby the cleavage site. The activatable antibodies disclosed herein find particular use where, for example, a protease capable of cleaving a site in the CM, *i.e.*, a protease, is present at relatively higher levels in or in close proximity to target-containing tissue of a treatment site or diagnostic site than in tissue of non-treatment sites (for example in healthy tissue). In some embodiments, a CM of the disclosure is also cleaved by one or more other proteases. In some embodiments, it is the one or more other proteases that is co-localized with the target and that is responsible for cleavage of the CM *in vivo*.

[000632] In some embodiments activatable antibodies provide for reduced toxicity and/or adverse side effects that could otherwise result from binding of the AB at non-treatment sites if the AB were not masked or otherwise inhibited from binding to the target.

[000633] In general, an activatable antibody can be designed by selecting an AB of interest and constructing the remainder of the activatable antibody so that, when conformationally constrained, the MM provides for masking of the AB or reduction of binding of the AB to its target. Structural design criteria can be taken into account to provide for this functional feature.

[000634] Activatable antibodies exhibiting a switchable phenotype of a desired dynamic range for target binding in an inhibited versus an uninhibited conformation are provided. Dynamic range generally refers to a ratio of (a) a maximum detected level of a parameter under a first set of conditions to (b) a minimum detected value of that parameter under a second set of conditions. For example, in the context of an activatable antibody, the dynamic range refers to the ratio of (a) a maximum detected level of target protein binding to an activatable antibody in the presence of at least one protease capable of cleaving the CM of the activatable antibodies to (b) a minimum detected level of target protein binding to an activatable antibody in the absence

of the protease. The dynamic range of an activatable antibody can be calculated as the ratio of the dissociation constant of an activatable antibody cleaving agent (*e.g.*, enzyme) treatment to the dissociation constant of the activatable antibodies cleaving agent treatment. The greater the dynamic range of an activatable antibody, the better the switchable phenotype of the activatable antibody. Activatable antibodies having relatively higher dynamic range values (*e.g.*, greater than 1) exhibit more desirable switching phenotypes such that target protein binding by the activatable antibodies occurs to a greater extent (*e.g.*, predominantly occurs) in the presence of a cleaving agent (*e.g.*, enzyme) capable of cleaving the CM of the activatable antibodies than in the absence of a cleaving agent.

[000635] Activatable antibodies can be provided in a variety of structural configurations. Exemplary formulae for activatable antibodies are provided below. It is specifically contemplated that the N- to C-terminal order of the AB, MM and CM can be reversed within an activatable antibody. It is also specifically contemplated that the CM and MM can overlap in amino acid sequence, *e.g.*, such that the CM is contained within the MM.

[000636] For example, activatable antibodies can be represented by the following formula (in order from an amino (N) terminal region to carboxyl (C) terminal region:

(MM)-(CM)-(AB)

(AB)-(CM)-(MM)

where MM is a masking moiety, CM is a cleavable moiety, and AB is an antibody or fragment thereof. It should be noted that although MM and CM are indicated as distinct components in the formulae above, in all exemplary embodiments (including formulae) disclosed herein it is contemplated that the amino acid sequences of the MM and the CM could overlap, *e.g.*, such that the CM is completely or partially contained within the MM. In addition, the formulae above provide for additional amino acid sequences that can be positioned N-terminal or C-terminal to the activatable antibodies elements.

[000637] In certain embodiments, the MM is not a natural binding partner of the AB. In some embodiments, the MM contains no or substantially no homology to any natural binding partner of the AB. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% similar to any natural binding partner of the AB. In some embodiments, the MM is no more than 5%, 10%, 15%, 20%, 25%, 30%, 35%, 40%, 45%, 50%, 55%, 60%, 65%, 70%, 75%, or 80% identical to any natural binding

partner of the AB. In some embodiments, the MM is no more than 50% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 25% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 20% identical to any natural binding partner of the AB. In some embodiments, the MM is no more than 10% identical to any natural binding partner of the AB.

[000638] In many embodiments it can be desirable to insert one or more linkers, *e.g.*, flexible linkers, into the activatable antibody construct so as to provide for flexibility at one or more of the MM-CM junction, the CM-AB junction, or both. For example, the AB, MM, and/or CM may not contain a sufficient number of residues (*e.g.*, Gly, Ser, Asp, Asn, especially Gly and Ser, particularly Gly) to provide the desired flexibility. As such, the switchable phenotype of such activatable antibody constructs can benefit from introduction of one or more amino acids to provide for a flexible linker. In addition, as described below, where the activatable antibody is provided as a conformationally constrained construct, a flexible linker can be operably inserted to facilitate formation and maintenance of a cyclic structure in the uncleaved activatable antibody.

[000639] For example, in certain embodiments an activatable antibody comprises one of the following formulae (where the formula below represent an amino acid sequence in either N- to C-terminal direction or C- to N-terminal direction):

$$(MM)-L1-(CM)-(AB)$$

$$(MM)-(CM)-L2-(AB)$$

$$(MM)-L1-(CM)-L2-(AB)$$

wherein MM, CM, and AB are as defined above; wherein L1 and L2 are each independently and optionally present or absent, are the same or different flexible linkers that include at least 1 flexible amino acid (*e.g.*, Gly). In addition, the formulae above provide for additional amino acid sequences that can be positioned N-terminal or C-terminal to the activatable antibodies elements. Examples include, but are not limited to, targeting moieties (*e.g.*, a ligand for a receptor of a cell present in a target tissue) and serum half-life extending moieties (*e.g.*, polypeptides that bind serum proteins, such as immunoglobulin (*e.g.*, IgG) or serum albumin (*e.g.*, human serum albumin (HAS))).

[000640] The CM is specifically cleaved by at least one protease at a rate of about 0.001-1500 x 10⁴ M⁻¹S⁻¹ or at least 0.001, 0.005, 0.01, 0.05, 0.1, 0.5, 1, 2.5, 5, 7.5, 10, 15, 20, 25, 50,

75, 100, 125, 150, 200, 250, 500, 750, 1000, 1250, or $1500 \times 10^4 \text{ M}^{-1}\text{S}^{-1}$. In some embodiments, the CM is specifically cleaved at a rate of about $100,000 \text{ M}^{-1}\text{S}^{-1}$. In some embodiments, the CM is specifically cleaved at a rate from about 1×10^2 to about $1 \times 10^6 \text{ M}^{-1}\text{S}^{-1}$ (i.e., from about 1×10^2 to about $1 \times 10^6 \text{ M}^{-1}\text{S}^{-1}$).

[000641] For specific cleavage by an enzyme, contact between the enzyme and CM is made. When the activatable antibody comprising an AB coupled to a MM and a CM is in the presence of target and sufficient enzyme activity, the CM can be cleaved. Sufficient enzyme activity can refer to the ability of the enzyme to make contact with the CM and effect cleavage. It can readily be envisioned that an enzyme can be in the vicinity of the CM but unable to cleave because of other cellular factors or protein modification of the enzyme.

[000642] Linkers suitable for use in compositions described herein are generally ones that provide flexibility of the modified AB or the activatable antibodies to facilitate the inhibition of the binding of the AB to the target. Such linkers are generally referred to as flexible linkers. Suitable linkers can be readily selected and can be of any of a suitable of different lengths, such as from 1 amino acid (*e.g.*, Gly) to 20 amino acids, from 2 amino acids to 15 amino acids, from 3 amino acids to 12 amino acids, including 4 amino acids to 10 amino acids, 5 amino acids to 9 amino acids, 6 amino acids to 8 amino acids, or 7 amino acids to 8 amino acids, and can be 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, or 20 amino acids in length.

[000643] Exemplary flexible linkers include glycine polymers (G)_n, glycine-serine polymers (including, for example, (GS)_n, (GSGGS)_n (SEQ ID NO: 363) and (GGGS)_n (SEQ ID NO: 364), where n is an integer of at least one), glycine-alanine polymers, alanine-serine polymers, and other flexible linkers known in the art. Glycine and glycine-serine polymers are relatively unstructured, and therefore can be able to serve as a neutral tether between components. Glycine accesses significantly more phi-psi space than even alanine, and is much less restricted than residues with longer side chains (see Scheraga, *Rev. Computational Chem.* 11173-142 (1992)). Exemplary flexible linkers include, but are not limited to GGSG (SEQ ID NO: 365), GGSGG (SEQ ID NO: 366), GSGSG (SEQ ID NO: 367), GSGGG (SEQ ID NO: 368), GGGSG (SEQ ID NO: 369), GSSSG (SEQ ID NO: 370), and the like. The ordinarily skilled artisan will recognize that design of an activatable antibodies can include linkers that are all or partially flexible, such that the linker can include a flexible linker as well as one or more

portions that confer less flexible structure to provide for a desired activatable antibodies structure.

[000644] The disclosure also provides compositions and methods that include an activatable anti-PD-1 antibody that includes an antibody or antibody fragment (AB) that specifically binds PD-1, where the AB is coupled to a masking moiety (MM) that decreases the ability of the AB to bind its target. In some embodiments, the activatable anti-PD-1 antibody further includes a cleavable moiety (CM) that is a substrate for a protease. The compositions and methods provided herein enable the attachment of one or more agents to one or more cysteine residues in the AB without compromising the activity (e.g., the masking, activating or binding activity) of the activatable anti-PD-1 antibody. In some embodiments, the compositions and methods provided herein enable the attachment of one or more agents to one or more cysteine residues in the AB without reducing or otherwise disturbing one or more disulfide bonds within the MM. The compositions and methods provided herein produce an activatable anti-PD-1 antibody that is conjugated to one or more agents, *e.g.*, any of a variety of therapeutic, diagnostic and/or prophylactic agents, for example, in some embodiments, without any of the agent(s) being conjugated to the MM of the activatable anti-PD-1 antibody. The compositions and methods provided herein produce conjugated activatable anti-PD-1 antibodies in which the MM retains the ability to effectively and efficiently mask the AB of the activatable antibody in an uncleaved state. The compositions and methods provided herein produce conjugated activatable anti-PD-1 antibodies in which the activatable antibody is still activated, *i.e.*, cleaved, in the presence of a protease that can cleave the CM.

[000645] The activatable anti-PD-1 antibodies have at least one point of conjugation for an agent, but in the methods and compositions provided herein less than all possible points of conjugation are available for conjugation to an agent. In some embodiments, the one or more points of conjugation are sulfur atoms involved in disulfide bonds. In some embodiments, the one or more points of conjugation are sulfur atoms involved in interchain disulfide bonds. In some embodiments, the one or more points of conjugation are sulfur atoms involved in interchain sulfide bonds, but not sulfur atoms involved in intrachain disulfide bonds. In some embodiments, the one or more points of conjugation are sulfur atoms of cysteine or other amino acid residues containing a sulfur atom. Such residues can occur naturally in the antibody

structure or can be incorporated into the antibody by site-directed mutagenesis, chemical conversion, or mis-incorporation of non-natural amino acids.

[000646] Also provided are methods of preparing a conjugate of an activatable anti-PD-1 antibody having one or more interchain disulfide bonds in the AB and one or more intrachain disulfide bonds in the MM, and a drug reactive with free thiols is provided. The method generally includes partially reducing interchain disulfide bonds in the activatable antibody with a reducing agent, such as, for example, TCEP; and conjugating the drug reactive with free thiols to the partially reduced activatable antibody. As used herein, the term partial reduction refers to situations where an activatable anti-PD-1 antibody is contacted with a reducing agent and less than all disulfide bonds, *e.g.*, less than all possible sites of conjugation are reduced. In some embodiments, less than 99%, 98%, 97%, 96%, 95%, 90%, 85%, 80%, 75%, 70%, 65%, 60%, 55%, 50%, 45%, 40%, 35%, 30%, 25%, 20%, 15%, 10% or less than 5% of all possible sites of conjugation are reduced.

[000647] In yet other embodiments, a method of reducing and conjugating an agent, *e.g.*, a drug, to an activatable anti-PD-1 antibody resulting in selectivity in the placement of the agent is provided. The method generally includes partially reducing the activatable anti-PD-1 antibody with a reducing agent such that any conjugation sites in the masking moiety or other non-AB portion of the activatable antibody are not reduced, and conjugating the agent to interchain thiols in the AB. The conjugation site(s) are selected so as to allow desired placement of an agent to allow conjugation to occur at a desired site. The reducing agent is, for example, TCEP. The reduction reaction conditions such as, for example, the ratio of reducing agent to activatable antibody, the length of incubation, the temperature during the incubation, the pH of the reducing reaction solution, etc., are determined by identifying the conditions that produce a conjugated activatable antibody in which the MM retains the ability to effectively and efficiently mask the AB of the activatable antibody in an uncleaved state. The ratio of reduction agent to activatable anti-PD-1 antibody will vary depending on the activatable antibody. In some embodiments, the ratio of reducing agent to activatable anti-PD-1 antibody will be in a range from about 20:1 to 1:1, from about 10:1 to 1:1, from about 9:1 to 1:1, from about 8:1 to 1:1, from about 7:1 to 1:1, from about 6:1 to 1:1, from about 5:1 to 1:1, from about 4:1 to 1:1, from about 3:1 to 1:1, from about 2:1 to 1:1, from about 20:1 to 1:1.5, from about 10:1 to 1:1.5, from about 9:1 to 1:1.5, from about 8:1 to 1:1.5, from about 7:1 to 1:1.5, from about 6:1 to 1:1.5, from about 5:1 to 1:1.5, from

about 4:1 to 1:1.5, from about 3:1 to 1:1.5, from about 2:1 to 1:1.5, from about 1.5:1 to 1:1.5, or from about 1:1 to 1:1.5. In some embodiments, the ratio is in a range of from about 5:1 to 1:1. In some embodiments, the ratio is in a range of from about 5:1 to 1.5:1. In some embodiments, the ratio is in a range of from about 4:1 to 1:1. In some embodiments, the ratio is in a range from about 4:1 to 1.5:1. In some embodiments, the ratio is in a range from about 8:1 to about 1:1. In some embodiments, the ratio is in a range of from about 2.5:1 to 1:1.

[000648] In some embodiments, a method of reducing interchain disulfide bonds in the AB of an activatable anti-PD-1 antibody and conjugating an agent, e.g., a thiol-containing agent such as a drug, to the resulting interchain thiols to selectively locate agent(s) on the AB is provided. The method generally includes partially reducing the AB with a reducing agent to form at least two interchain thiols without forming all possible interchain thiols in the activatable antibody; and conjugating the agent to the interchain thiols of the partially reduced AB. For example, the AB of the activatable antibody is partially reduced for about 1 hour at about 37°C at a desired ratio of reducing agent:activatable antibody. In some embodiments, the ratio of reducing agent to activatable antibody will be in a range from about 20:1 to 1:1, from about 10:1 to 1:1, from about 9:1 to 1:1, from about 8:1 to 1:1, from about 7:1 to 1:1, from about 6:1 to 1:1, from about 5:1 to 1:1, from about 4:1 to 1:1, from about 3:1 to 1:1, from about 2:1 to 1:1, from about 20:1 to 1:1.5, from about 10:1 to 1:1.5, from about 9:1 to 1:1.5, from about 8:1 to 1:1.5, from about 7:1 to 1:1.5, from about 6:1 to 1:1.5, from about 5:1 to 1:1.5, from about 4:1 to 1:1.5, from about 3:1 to 1:1.5, from about 2:1 to 1:1.5, from about 1.5:1 to 1:1.5, or from about 1:1 to 1:1.5. In some embodiments, the ratio is in a range of from about 5:1 to 1:1. In some embodiments, the ratio is in a range of from about 5:1 to 1.5:1. In some embodiments, the ratio is in a range of from about 4:1 to 1:1. In some embodiments, the ratio is in a range from about 4:1 to 1.5:1. In some embodiments, the ratio is in a range from about 8:1 to about 1:1. In some embodiments, the ratio is in a range of from about 2.5:1 to 1:1.

[000649] The thiol-containing reagent can be, for example, cysteine or N-acetyl cysteine. The reducing agent can be, for example, TCEP. In some embodiments, the reduced activatable antibody can be purified prior to conjugation, using for example, column chromatography, dialysis, or diafiltration. Alternatively, the reduced antibody is not purified after partial reduction and prior to conjugation.

[000650] The invention also provides partially reduced activatable anti-PD-1 antibodies in which at least one interchain disulfide bond in the activatable antibody has been reduced with a reducing agent without disturbing any intrachain disulfide bonds in the activatable antibody, wherein the activatable antibody includes an antibody or an antigen binding fragment thereof (AB) that specifically binds to PD-1, a masking moiety (MM) that inhibits the binding of the AB of the activatable antibody in an uncleaved state to the PD-1 target, and a cleavable moiety (CM) coupled to the AB, wherein the CM is a polypeptide that functions as a substrate for a protease. In some embodiments the MM is coupled to the AB via the CM. In some embodiments, one or more intrachain disulfide bond(s) of the activatable antibody is not disturbed by the reducing agent. In some embodiments, one or more intrachain disulfide bond(s) of the MM within the activatable antibody is not disturbed by the reducing agent. In some embodiments, the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM. In some embodiments, reducing agent is TCEP.

[000651] The disclosure also provides partially reduced activatable antibodies in which at least one interchain disulfide bond in the activatable antibody has been reduced with a reducing agent without disturbing any intrachain disulfide bonds in the activatable antibody, wherein the activatable antibody includes an antibody or an antigen binding fragment thereof (AB) that specifically binds to the target, e.g., PD-1, a masking moiety (MM) that inhibits the binding of the AB of the activatable antibody in an uncleaved state to the target, and a cleavable moiety (CM) coupled to the AB, wherein the CM is a polypeptide that functions as a substrate for at least one protease. In some embodiments, the MM is coupled to the AB via the CM. In some embodiments, one or more intrachain disulfide bond(s) of the activatable antibody is not disturbed by the reducing agent. In some embodiments, one or more intrachain disulfide bond(s) of the MM within the activatable antibody is not disturbed by the reducing agent. In some embodiments, the activatable antibody in the uncleaved state has the structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM. In some embodiments, reducing agent is TCEP.

[000652] In some embodiments, the activatable antibodies described herein also include an agent conjugated to the activatable antibody. In some embodiments, the conjugated agent is a therapeutic agent, such as an anti-inflammatory. In such embodiments, the agent is conjugated to

a carbohydrate moiety of the activatable antibody, for example, in some embodiments, where the carbohydrate moiety is located outside the antigen-binding region of the antibody or antigen-binding fragment in the activatable antibody. In some embodiments, the agent is conjugated to a sulfhydryl group of the antibody or antigen-binding fragment in the activatable antibody.

[000653] In some embodiments, the agent is a detectable moiety such as, for example, a label or other marker. For example, the agent is or includes a radiolabeled amino acid, one or more biotinyl moieties that can be detected by marked avidin (*e.g.*, streptavidin containing a fluorescent marker or enzymatic activity that can be detected by optical or calorimetric methods), one or more radioisotopes or radionuclides, one or more fluorescent labels, one or more enzymatic labels, and/or one or more chemiluminescent agents. In some embodiments, detectable moieties are attached by spacer molecules.

[000654] Those of ordinary skill in the art will recognize that a large variety of possible moieties can be coupled to the resultant antibodies of the disclosure. (*See, for example*, "Conjugate Vaccines", Contributions to Microbiology and Immunology, J. M. Cruse and R. E. Lewis, Jr (eds), Carger Press, New York, (1989), the entire contents of which are incorporated herein by reference).

[000655] Coupling can be accomplished by any chemical reaction that will bind the two molecules so long as the antibody and the other moiety retain their respective activities. This linkage can include many chemical mechanisms, for instance covalent binding, affinity binding, intercalation, coordinate binding and complexation. In some embodiments, the binding is, however, covalent binding. Covalent binding can be achieved either by direct condensation of existing side chains or by the incorporation of external bridging molecules. Many bivalent or polyvalent linking agents are useful in coupling protein molecules, such as the antibodies of the present disclosure, to other molecules. For example, representative coupling agents can include organic compounds such as thioesters, carbodiimides, succinimide esters, diisocyanates, glutaraldehyde, diazobenzenes and hexamethylene diamines. This listing is not intended to be exhaustive of the various classes of coupling agents known in the art but, rather, is exemplary of the more common coupling agents. (*See* Killen and Lindstrom, Jour. Immun. 133:1335-2549 (1984); Jansen et al., Immunological Reviews 62:185-216 (1982); and Vitetta et al., Science 238:1098 (1987).

[000656] In some embodiments, in addition to the compositions and methods provided herein, the conjugated activatable antibody can also be modified for site-specific conjugation through modified amino acid sequences inserted or otherwise included in the activatable antibody sequence. These modified amino acid sequences are designed to allow for controlled placement and/or dosage of the conjugated agent within a conjugated activatable antibody. For example, the activatable antibody can be engineered to include cysteine substitutions at positions on light and heavy chains that provide reactive thiol groups and do not negatively impact protein folding and assembly, nor alter antigen binding. In some embodiments, the activatable antibody can be engineered to include or otherwise introduce one or more non-natural amino acid residues within the activatable antibody to provide suitable sites for conjugation. In some embodiments, the activatable antibody can be engineered to include or otherwise introduce enzymatically activatable peptide sequences within the activatable antibody sequence.

[000657] Suitable linkers are described in the literature. (*See, for example*, Ramakrishnan, S. et al., Cancer Res. 44:201-208 (1984) describing use of MBS (M-maleimidobenzoyl-N-hydroxysuccinimide ester). *See also*, U.S. Patent No. 5,030,719, describing use of halogenated acetyl hydrazide derivative coupled to an antibody by way of an oligopeptide linker. In some embodiments, suitable linkers include: (i) EDC (1-ethyl-3-(3-dimethylamino-propyl) carbodiimide hydrochloride; (ii) SMPT (4-succinimidylloxycarbonyl-alpha-methyl-alpha-(2-pyridyl-dithio)-toluene (Pierce Chem. Co., Cat. (21558G); (iii) SPDP (succinimidyl-6 [3-(2-pyridyl-dithio) propionamido]hexanoate (Pierce Chem. Co., Cat #21651G); (iv) Sulfo-LC-SPDP (sulfosuccinimidyl 6 [3-(2-pyridyl-dithio)-propionamide] hexanoate (Pierce Chem. Co. Cat. #2165-G); and (v) sulfo-NHS (N-hydroxysulfo-succinimide: Pierce Chem. Co., Cat. #24510) conjugated to EDC. Additional linkers include, but are not limited to, SMCC, sulfo-SMCC, SPDB, or sulfo-SPDB.

[000658] The linkers described above contain components that have different attributes, thus leading to conjugates with differing physio-chemical properties. For example, sulfo-NHS esters of alkyl carboxylates are more stable than sulfo-NHS esters of aromatic carboxylates. NHS-ester containing linkers are less soluble than sulfo-NHS esters. Further, the linker SMPT contains a sterically hindered disulfide bond, and can form conjugates with increased stability. Disulfide linkages, are in general, less stable than other linkages because the disulfide linkage is cleaved *in vitro*, resulting in less conjugate available. Sulfo-NHS, in particular, can enhance the

stability of carbodimide couplings. Carbodimide couplings (such as EDC) when used in conjunction with sulfo-NHS, forms esters that are more resistant to hydrolysis than the carbodimide coupling reaction alone.

[000659] In some embodiments, the linkers are cleavable. In some embodiments, the linkers are non-cleavable. In some embodiments, two or more linkers are present. The two or more linkers are all the same, *i.e.*, cleavable or non-cleavable, or the two or more linkers are different, *i.e.*, at least one cleavable and at least one non-cleavable.

[000660] The present disclosure utilizes several methods for attaching agents to ABs: (a) attachment to the carbohydrate moieties of the AB, or (b) attachment to sulfhydryl groups of the AB, or (c) attachment to amino groups of the AB, or (d) attachment to carboxylate groups of the AB. According to the disclosure, ABs can be covalently attached to an agent through an intermediate linker having at least two reactive groups, one to react with AB and one to react with the agent. The linker, which can include any compatible organic compound, can be chosen such that the reaction with AB (or agent) does not adversely affect AB reactivity and selectivity. Furthermore, the attachment of linker to agent might not destroy the activity of the agent. Suitable linkers for reaction with oxidized antibodies or oxidized antibody fragments include those containing an amine selected from the group consisting of primary amine, secondary amine, hydrazine, hydrazide, hydroxylamine, phenylhydrazine, semicarbazide and thiosemicarbazide groups. Such reactive functional groups can exist as part of the structure of the linker, or can be introduced by suitable chemical modification of linkers not containing such groups.

[000661] According to the present disclosure, suitable linkers for attachment to reduced ABs include those having certain reactive groups capable of reaction with a sulfhydryl group of a reduced antibody or fragment. Such reactive groups include, but are not limited to: reactive haloalkyl groups (including, for example, haloacetyl groups), p-mercuribenzoate groups and groups capable of Michael-type addition reactions (including, for example, maleimides and groups of the type described by Mitra and Lawton, 1979, J. Amer. Chem. Soc. 101: 3097-3110).

[000662] According to the present disclosure, suitable linkers for attachment to neither oxidized nor reduced Abs include those having certain functional groups capable of reaction with the primary amino groups present in unmodified lysine residues in the Ab. Such reactive groups include, but are not limited to, NHS carboxylic or carbonic esters, sulfo-NHS carboxylic or

carbonic esters, 4-nitrophenyl carboxylic or carbonic esters, pentafluorophenyl carboxylic or carbonic esters, acyl imidazoles, isocyanates, and isothiocyanates.

[000663] According to the present disclosure, suitable linkers for attachment to neither oxidized nor reduced Abs include those having certain functional groups capable of reaction with the carboxylic acid groups present in aspartate or glutamate residues in the Ab, which have been activated with suitable reagents. Suitable activating reagents include EDC, with or without added NHS or sulfo-NHS, and other dehydrating agents utilized for carboxamide formation. In these instances, the functional groups present in the suitable linkers would include primary and secondary amines, hydrazines, hydroxylamines, and hydrazides.

[000664] The agent can be attached to the linker before or after the linker is attached to the AB. In certain applications it can be desirable to first produce an AB-linker intermediate in which the linker is free of an associated agent. Depending upon the particular application, a specific agent can then be covalently attached to the linker. In some embodiments, the AB is first attached to the MM, CM and associated linkers and then attached to the linker for conjugation purposes.

[000665] *Branched Linkers:* In specific embodiments, branched linkers that have multiple sites for attachment of agents are utilized. For multiple site linkers, a single covalent attachment to an AB would result in an AB-linker intermediate capable of binding an agent at a number of sites. The sites can be aldehyde or sulfhydryl groups or any chemical site to which agents can be attached.

[000666] In some embodiments, higher specific activity (or higher ratio of agents to AB) can be achieved by attachment of a single site linker at a plurality of sites on the AB. This plurality of sites can be introduced into the AB by either of two methods. First, one can generate multiple aldehyde groups and/or sulfhydryl groups in the same AB. Second, one can attach to an aldehyde or sulfhydryl of the AB a "branched linker" having multiple functional sites for subsequent attachment to linkers. The functional sites of the branched linker or multiple site linker can be aldehyde or sulfhydryl groups, or can be any chemical site to which linkers can be attached. Still higher specific activities can be obtained by combining these two approaches, that is, attaching multiple site linkers at several sites on the AB.

[000667] *Cleavable Linkers:* Peptide linkers that are susceptible to cleavage by enzymes of the complement system, such as but not limited to u-plasminogen activator, tissue plasminogen

activator, trypsin, plasmin, or another enzyme having proteolytic activity can be used in one embodiment of the present disclosure. According to one method of the present disclosure, an agent is attached via a linker susceptible to cleavage by complement. The antibody is selected from a class that can activate complement. The antibody-agent conjugate, thus, activates the complement cascade and releases the agent at the target site. According to another method of the present disclosure, an agent is attached via a linker susceptible to cleavage by enzymes having a proteolytic activity such as a u-plasminogen activator, a tissue plasminogen activator, plasmin, or trypsin.

[000668] Non-limiting examples of cleavable linker sequences are provided in Table 5.

Table 5: Exemplary Linker Sequences for Conjugation

Types of Cleavable Sequences	Amino Acid Sequence
<u>Plasmin cleavable sequences</u>	
Pro-urokinase	PRFKIIGG (SEQ ID NO: 587) PRFRIIGG (SEQ ID NO: 588)
TGF β	SSRHRRALD (SEQ ID NO: 589)
Plasminogen	RKSSIIIRMRDVVL (SEQ ID NO: 590)
Staphylokinase	SSSFDKGKYKKGDDA (SEQ ID NO: 591) SSSFDKGKYKRGDDA (SEQ ID NO: 592)
<u>Factor Xa cleavable sequences</u>	
	IEGR (SEQ ID NO: 593) IDGR (SEQ ID NO: 594) GGSIDGR (SEQ ID NO: 595)
<u>MMP cleavable sequences</u>	
Gelatinase A	PLGLWA (SEQ ID NO: 596)
<u>Collagenase cleavable sequences</u>	
Calf skin collagen (α 1(I) chain)	GPQGIAGQ (SEQ ID NO: 597)
Calf skin collagen (α 2(I) chain)	GPQGLLGA (SEQ ID NO: 598)
Bovine cartilage collagen (α 1(II) chain)	GIAGQ (SEQ ID NO: 599)
Human liver collagen (α 1(III) chain)	GPLGIAGI (SEQ ID NO: 600)
Human α ₂ M	GPEGLRVG (SEQ ID NO: 601)
Human PZP	YGAGLGVV (SEQ ID NO: 602) AGLGVVER (SEQ ID NO: 603) AGLGISST (SEQ ID NO: 604)
Rat α ₁ M	EPQALAMS (SEQ ID NO: 605)
	QALAMSAI (SEQ ID NO: 606)
Rat α ₂ M	AAYHLVSQ (SEQ ID NO: 607)
	MDAFLESS (SEQ ID NO: 608)
Rat α ₁ I ₃ (2J)	ESLPVVAV (SEQ ID NO: 609)

Rat $\alpha_1\text{I}_3$ (27J)
Human fibroblast collagenase
(autolytic cleavages)

SAPAVESE (SEQ ID NO: 610)
DVAQFVLT (SEQ ID NO: 611)
VAQFVLTE (SEQ ID NO: 612)
AQFVLTEG (SEQ ID NO: 613)
PVQPIGPQ (SEQ ID NO: 614)

[000669] In addition, agents can be attached via disulfide bonds (for example, the disulfide bonds on a cysteine molecule) to the AB. Since many tumors naturally release high levels of glutathione (a reducing agent) this can reduce the disulfide bonds with subsequent release of the agent at the site of delivery. In some embodiments, the reducing agent that would modify a CM would also modify the linker of the conjugated activatable antibody.

[000670] *Spacers and Cleavable Elements:* In some embodiments, it can be necessary to construct the linker in such a way as to optimize the spacing between the agent and the AB of the activatable antibody. This can be accomplished by use of a linker of the general structure:



wherein

W is either --NH--CH₂-- or --CH₂--;

Q is an amino acid, peptide; and

n is an integer from 0 to 20.

[000671] In some embodiments, the linker can comprise a spacer element and a cleavable element. The spacer element serves to position the cleavable element away from the core of the AB such that the cleavable element is more accessible to the enzyme responsible for cleavage. Certain of the branched linkers described above can serve as spacer elements.

[000672] Throughout this discussion, it should be understood that the attachment of linker to agent (or of spacer element to cleavable element, or cleavable element to agent) need not be particular mode of attachment or reaction. Any reaction providing a product of suitable stability and biological compatibility is acceptable.

[000673] *Serum Complement and Selection of Linkers:* According to one method of the present disclosure, when release of an agent is desired, an AB that is an antibody of a class that can activate complement is used. The resulting conjugate retains both the ability to bind antigen and activate the complement cascade. Thus, according to this embodiment of the present disclosure, an agent is joined to one end of the cleavable linker or cleavable element and the

other end of the linker group is attached to a specific site on the AB. For example, if the agent has an hydroxy group or an amino group, it can be attached to the carboxy terminus of a peptide, amino acid or other suitably chosen linker via an ester or amide bond, respectively. For example, such agents can be attached to the linker peptide via a carbodimide reaction. If the agent contains functional groups that would interfere with attachment to the linker, these interfering functional groups can be blocked before attachment and deblocked once the product conjugate or intermediate is made. The opposite or amino terminus of the linker is then used either directly or after further modification for binding to an AB that is capable of activating complement.

[000674] Linkers (or spacer elements of linkers) can be of any desired length, one end of which can be covalently attached to specific sites on the AB of the activatable antibody. The other end of the linker or spacer element can be attached to an amino acid or peptide linker.

[000675] Thus when these conjugates bind to antigen in the presence of complement the amide or ester bond that attaches the agent to the linker will be cleaved, resulting in release of the agent in its active form. These conjugates, when administered to a subject, will accomplish delivery and release of the agent at the target site, and are particularly effective for the in vivo delivery of pharmaceutical agents, antibiotics, antimetabolites, antiproliferative agents and the like as presented in but not limited to those in Table 5.

[000676] *Linkers for Release without Complement Activation:* In yet another application of targeted delivery, release of the agent without complement activation is desired since activation of the complement cascade will ultimately lyse the target cell. Hence, this approach is useful when delivery and release of the agent should be accomplished without killing the target cell. Such is the goal when delivery of cell mediators such as hormones, enzymes, corticosteroids, neurotransmitters, genes or enzymes to target cells is desired. These conjugates can be prepared by attaching the agent to an AB that is not capable of activating complement via a linker that is mildly susceptible to cleavage by serum proteases. When this conjugate is administered to an individual, antigen-antibody complexes will form quickly whereas cleavage of the agent will occur slowly, thus resulting in release of the compound at the target site.

[000677] *Biochemical Cross Linkers:* In some embodiments, the activatable antibody can be conjugated to one or more therapeutic agents using certain biochemical cross-linkers. Cross-linking reagents form molecular bridges that tie together functional groups of two different

molecules. To link two different proteins in a step-wise manner, hetero-bifunctional cross-linkers can be used that eliminate unwanted homopolymer formation.

[000678] Peptidyl linkers cleavable by lysosomal proteases are also useful, for example, Val-Cit, Val-Ala or other dipeptides. In addition, acid-labile linkers cleavable in the low-pH environment of the lysosome can be used, for example: bis-sialyl ether. Other suitable linkers include cathepsin-labile substrates, particularly those that show optimal function at an acidic pH.

[000679] Exemplary hetero-bifunctional cross-linkers are referenced in Table 6.

Table 6: Exemplary Hetero-Bifunctional Cross Linkers

<u>HETERO-BIFUNCTIONAL CROSS-LINKERS</u>			
Linker	Reactive Toward	Advantages and Applications	Spacer Arm Length after cross-linking (Angstroms)
SMPT	Primary amines	Greater stability	11.2 Å
	Sulfhydryls		
SPDP	Primary amines	Thiolation	6.8 Å
	Sulfhydryls	Cleavable cross-linking	
LC-SPDP	Primary amines	Extended spacer arm	15.6 Å
	Sulfhydryls		
Sulfo-LC-SPDP	Primary amines	Extender spacer arm	15.6 Å
	Sulfhydryls	Water-soluble	
SMCC	Primary amines	Stable maleimide reactive group	11.6 Å
	Sulfhydryls	Enzyme-antibody conjugation	
		Hapten-carrier protein conjugation	
Sulfo-SMCC	Primary amines	Stable maleimide reactive group	11.6 Å
	Sulfhydryls	Water-soluble	
		Enzyme-antibody conjugation	
MBS	Primary amines	Enzyme-antibody conjugation	9.9 Å
	Sulfhydryls	Hapten-carrier protein conjugation	
Sulfo-MBS	Primary amines	Water-soluble	9.9 Å
	Sulfhydryls		
SIAB	Primary amines	Enzyme-antibody conjugation	10.6 Å
	Sulfhydryls		
Sulfo-SIAB	Primary amines	Water-soluble	10.6 Å
	Sulfhydryls		
SMPB	Primary amines	Extended spacer arm	14.5 Å
	Sulfhydryls	Enzyme-antibody conjugation	
Sulfo-SMPB	Primary amines	Extended spacer arm	14.5 Å

EDE/Sulfo-NHS	Sulfhydryls Primary amines Carboxyl groups	Water-soluble Hapten-Carrier conjugation	0
ABH	Carbohydrates Nonselective	Reacts with sugar groups	11.9 Å

[000680] *Non-Cleavable Linkers or Direct Attachment:* In some embodiments of the disclosure, the conjugate can be designed so that the agent is delivered to the target but not released. This can be accomplished by attaching an agent to an AB either directly or via a non-cleavable linker.

[000681] These non-cleavable linkers can include amino acids, peptides, D-amino acids or other organic compounds that can be modified to include functional groups that can subsequently be utilized in attachment to ABs by the methods described herein. A general formula for such an organic linker could be



wherein

W is either --NH--CH₂-- or --CH₂--;

Q is an amino acid, peptide; and

n is an integer from 0 to 20.

[000682] *Non-Cleavable Conjugates:* In some embodiments, a compound can be attached to ABs that do not activate complement. When using ABs that are incapable of complement activation, this attachment can be accomplished using linkers that are susceptible to cleavage by activated complement or using linkers that are not susceptible to cleavage by activated complement.

[000683] The antibodies disclosed herein can also be formulated as immunoliposomes. Liposomes containing the antibody are prepared by methods known in the art, such as described in Epstein et al., Proc. Natl. Acad. Sci. USA, 82: 3688 (1985); Hwang et al., Proc. Natl. Acad. Sci. USA, 77: 4030 (1980); and U.S. Pat. Nos. 4,485,045 and 4,544,545. Liposomes with enhanced circulation time are disclosed in U.S. Patent No. 5,013,556.

[000684] Particularly useful liposomes can be generated by the reverse-phase evaporation method with a lipid composition comprising phosphatidylcholine, cholesterol, and PEG-derivatized phosphatidylethanolamine (PEG-PE). Liposomes are extruded through filters of defined pore size to yield liposomes with the desired diameter. Fab' fragments of the antibody of

the present disclosure can be conjugated to the liposomes as described in Martin et al., J. Biol. Chem., 257: 286-288 (1982) via a disulfide-interchange reaction.

Definitions:

[000685] Unless otherwise defined, scientific and technical terms used in connection with the present disclosure shall have the meanings that are commonly understood by those of ordinary skill in the art. The term “a” entity or “an” entity refers to one or more of that entity. For example, a compound refers to one or more compounds. As such, the terms “a”, “an”, “one or more” and “at least one” can be used interchangeably. Further, unless otherwise required by context, singular terms shall include pluralities and plural terms shall include the singular. Generally, nomenclatures utilized in connection with, and techniques of, cell and tissue culture, molecular biology, and protein and oligo- or polynucleotide chemistry and hybridization described herein are those well-known and commonly used in the art. Standard techniques are used for recombinant DNA, oligonucleotide synthesis, and tissue culture and transformation (*e.g.*, electroporation, lipofection). Enzymatic reactions and purification techniques are performed according to manufacturer’s specifications or as commonly accomplished in the art or as described herein. The foregoing techniques and procedures are generally performed according to conventional methods well known in the art and as described in various general and more specific references that are cited and discussed throughout the present specification. *See e.g.*, Sambrook *et al.* Molecular Cloning: A Laboratory Manual (2d ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N.Y. (1989)). The nomenclatures utilized in connection with, and the laboratory procedures and techniques of, analytical chemistry, synthetic organic chemistry, and medicinal and pharmaceutical chemistry described herein are those well-known and commonly used in the art. Standard techniques are used for chemical syntheses, chemical analyses, pharmaceutical preparation, formulation, and delivery, and treatment of patients.

[000686] As utilized in accordance with the present disclosure, the following terms, unless otherwise indicated, shall be understood to have the following meanings:

[000687] As used herein, the term “antibody” refers to immunoglobulin molecules and antigen-binding portions of immunoglobulin (Ig) molecules, *i.e.*, molecules that contain an antigen binding site that specifically binds (immunoreacts with) an antigen. By “specifically bind” or “immunoreacts with” or “immunospecifically bind” is meant that the antibody reacts with one or more antigenic determinants of the desired antigen and does not react with other

polypeptides or binds at much lower affinity ($K_d > 10^{-6}$). Antibodies include, but are not limited to, polyclonal, monoclonal, chimeric, domain antibody, single chain, Fab, and $F(ab')_2$ fragments, scFvs, and an Fab expression library.

[000688] The basic antibody structural unit is known to comprise a tetramer. Each tetramer is composed of two identical pairs of polypeptide chains, each pair having one “light” (about 25 kDa) and one “heavy” chain (about 50-70 kDa). The amino-terminal portion of each chain includes a variable region of about 100 to 110 or more amino acids primarily responsible for antigen recognition. The carboxy-terminal portion of each chain defines a constant region primarily responsible for effector function. In general, antibody molecules obtained from humans relate to any of the classes IgG, IgM, IgA, IgE and IgD, which differ from one another by the nature of the heavy chain present in the molecule. Certain classes have subclasses as well, such as IgG₁, IgG₂, and others. Furthermore, in humans, the light chain can be a kappa chain or a lambda chain.

[000689] The term “monoclonal antibody” (mAb) or “monoclonal antibody composition”, as used herein, refers to a population of antibody molecules that contain only one molecular species of antibody molecule consisting of a unique light chain gene product and a unique heavy chain gene product. In particular, the complementarity determining regions (CDRs) of the monoclonal antibody are identical in all the molecules of the population. MAbs contain an antigen binding site capable of immunoreacting with a particular epitope of the antigen characterized by a unique binding affinity for it.

[000690] The term “antigen-binding site” or “binding portion” refers to the part of the immunoglobulin molecule that participates in antigen binding. The antigen binding site is formed by amino acid residues of the N-terminal variable (“V”) regions of the heavy (“H”) and light (“L”) chains. Three highly divergent stretches within the V regions of the heavy and light chains, referred to as “hypervariable regions,” are interposed between more conserved flanking stretches known as “framework regions,” or “FRs”. Thus, the term “FR” refers to amino acid sequences that are naturally found between, and adjacent to, hypervariable regions in immunoglobulins. In an antibody molecule, the three hypervariable regions of a light chain and the three hypervariable regions of a heavy chain are disposed relative to each other in three dimensional space to form an antigen-binding surface. The antigen-binding surface is complementary to the three-dimensional surface of a bound antigen, and the three hypervariable regions of each of the heavy and light

chains are referred to as “complementarity-determining regions,” or “CDRs.” The assignment of amino acids to each domain is in accordance with the definitions of Kabat Sequences of Proteins of Immunological Interest (National Institutes of Health, Bethesda, Md. (1987 and 1991)), or Chothia & Lesk J. Mol. Biol. 196:901-917 (1987), Chothia *et al.* Nature 342:878-883 (1989).

[000691] As used herein, the term “epitope” includes any protein determinant capable of specific binding to an immunoglobulin, an scFv, or a T-cell receptor. The term “epitope” includes any protein determinant capable of specific binding to an immunoglobulin or T-cell receptor. Epitopic determinants usually consist of chemically active surface groupings of molecules such as amino acids or sugar side chains and usually have specific three dimensional structural characteristics, as well as specific charge characteristics. For example, antibodies can be raised against N-terminal or C-terminal peptides of a polypeptide. An antibody is said to specifically bind an antigen when the dissociation constant is $\leq 1 \mu\text{M}$; in some embodiments, $\leq 100 \text{ nM}$ and in some embodiments, $\leq 10 \text{ nM}$.

[000692] As used herein, the terms “specific binding,” “immunological binding,” and “immunological binding properties” refer to the non-covalent interactions of the type which occur between an immunoglobulin molecule and an antigen for which the immunoglobulin is specific. The strength, or affinity of immunological binding interactions can be expressed in terms of the dissociation constant (K_d) of the interaction, wherein a smaller K_d represents a greater affinity. Immunological binding properties of selected polypeptides can be quantified using methods well known in the art. One such method entails measuring the rates of antigen-binding site/antigen complex formation and dissociation, wherein those rates depend on the concentrations of the complex partners, the affinity of the interaction, and geometric parameters that equally influence the rate in both directions. Thus, both the “on rate constant” (K_{on}) and the “off rate constant” (K_{off}) can be determined by calculation of the concentrations and the actual rates of association and dissociation. (*See* Nature 361:186-87 (1993)). The ratio of K_{off}/K_{on} is equal to the dissociation constant K_d . (*See, generally*, Davies et al. (1990) Annual Rev Biochem 59:439-473). An antibody of the present disclosure is said to specifically bind to the target, when the binding constant (K_d) is $\leq 1 \mu\text{M}$, in some embodiments $\leq 100 \text{ nM}$, in some embodiments $\leq 10 \text{ nM}$, and in some embodiments $\leq 100 \text{ pM}$ to about 1 pM , as measured by assays such as radioligand binding assays or similar assays known to those skilled in the art.

[000693] The term “isolated polynucleotide” as used herein shall mean a polynucleotide of genomic, cDNA, or synthetic origin or some combination thereof, which by virtue of its origin the “isolated polynucleotide” (1) is not associated with all or a portion of a polynucleotide in which the “isolated polynucleotide” is found in nature, (2) is operably linked to a polynucleotide which it is not linked to in nature, or (3) does not occur in nature as part of a larger sequence. Polynucleotides in accordance with the disclosure include the nucleic acid molecules encoding the heavy chain immunoglobulin molecules shown herein, and nucleic acid molecules encoding the light chain immunoglobulin molecules shown herein.

[000694] The term “isolated protein” referred to herein means a protein of cDNA, recombinant RNA, or synthetic origin or some combination thereof, which by virtue of its origin, or source of derivation, the “isolated protein” (1) is not associated with proteins found in nature, (2) is free of other proteins from the same source, *e.g.*, free of murine proteins, (3) is expressed by a cell from a different species, or (4) does not occur in nature.

[000695] The term “polypeptide” is used herein as a generic term to refer to native protein, fragments, or analogs of a polypeptide sequence. Hence, native protein fragments, and analogs are species of the polypeptide genus. Polypeptides in accordance with the disclosure comprise the heavy chain immunoglobulin molecules shown herein, and the light chain immunoglobulin molecules shown herein, as well as antibody molecules formed by combinations comprising the heavy chain immunoglobulin molecules with light chain immunoglobulin molecules, such as kappa light chain immunoglobulin molecules, and vice versa, as well as fragments and analogs thereof.

[000696] The term “naturally-occurring” as used herein as applied to an object refers to the fact that an object can be found in nature. For example, a polypeptide or polynucleotide sequence that is present in an organism (including viruses) that can be isolated from a source in nature and that has not been intentionally modified by man in the laboratory or otherwise is naturally-occurring.

[000697] The term “operably linked” as used herein refers to positions of components so described are in a relationship permitting them to function in their intended manner. A control sequence “operably linked” to a coding sequence is ligated in such a way that expression of the coding sequence is achieved under conditions compatible with the control sequences.

[000698] The term “control sequence” as used herein refers to polynucleotide sequences that are necessary to effect the expression and processing of coding sequences to which they are ligated. The nature of such control sequences differs depending upon the host organism in prokaryotes, such control sequences generally include promoter, ribosomal binding site, and transcription termination sequence in eukaryotes, generally, such control sequences include promoters and transcription termination sequence. The term “control sequences” is intended to include, at a minimum, all components whose presence is essential for expression and processing, and can also include additional components whose presence is advantageous, for example, leader sequences and fusion partner sequences. The term “polynucleotide” as referred to herein means nucleotides of at least 10 bases in length, either ribonucleotides or deoxynucleotides or a modified form of either type of nucleotide. The term includes single and double stranded forms of DNA.

[000699] The term oligonucleotide referred to herein includes naturally occurring, and modified nucleotides linked together by naturally occurring, and non-naturally occurring oligonucleotide linkages. Oligonucleotides are a polynucleotide subset generally comprising a length of 200 bases or fewer. In some embodiments, oligonucleotides are 10 to 60 bases in length and in some embodiments, 12, 13, 14, 15, 16, 17, 18, 19, or 20 to 40 bases in length. Oligonucleotides are usually single stranded, *e.g.*, for probes, although oligonucleotides can be double stranded, *e.g.*, for use in the construction of a gene mutant. Oligonucleotides of the disclosure are either sense or antisense oligonucleotides.

[000700] The term “naturally occurring nucleotides” referred to herein includes deoxyribonucleotides and ribonucleotides. The term “modified nucleotides” referred to herein includes nucleotides with modified or substituted sugar groups and the like. The term “oligonucleotide linkages” referred to herein includes oligonucleotide linkages such as phosphorothioate, phosphorodithioate, phosphoroselenoate, phosphorodiselenoate, phosphoroanilothioate, phosphoraniladate, phosphoronmidate, and the like. *See e.g.*, LaPlanche *et al.* Nucl. Acids Res. 14:9081 (1986); Stec *et al.* J. Am. Chem. Soc. 106:6077 (1984), Stein *et al.* Nucl. Acids Res. 16:3209 (1988), Zon *et al.* Anti Cancer Drug Design 6:539 (1991); Zon *et al.* Oligonucleotides and Analogues: A Practical Approach, pp. 87-108 (F. Eckstein, Ed., Oxford University Press, Oxford England (1991)); Stec *et al.* U.S. Patent No. 5,151,510; Uhlmann and

Peyman Chemical Reviews 90:543 (1990). An oligonucleotide can include a label for detection, if desired.

[000701] As used herein, the twenty conventional amino acids and their abbreviations follow conventional usage. *See Immunology - A Synthesis* (2nd Edition, E.S. Golub and D.R. Green, Eds., Sinauer Associates, Sunderland, Mass. (1991)). Stereoisomers (*e.g.*, D- amino acids) of the twenty conventional amino acids, unnatural amino acids such as α -, α -disubstituted amino acids, N-alkyl amino acids, lactic acid, and other unconventional amino acids can also be suitable components for polypeptides of the present disclosure. Examples of unconventional amino acids include: 4 hydroxyproline, γ -carboxyglutamate, ϵ -N,N,N-trimethyllysine, ϵ -N-acetyllysine, O-phosphoserine, N- acetylserine, N-formylmethionine, 3-methylhistidine, 5-hydroxylysine, σ -N-methylarginine, and other similar amino acids and imino acids (*e.g.*, 4-hydroxyproline). In the polypeptide notation used herein, the left-hand direction is the amino terminal direction and the right-hand direction is the carboxy-terminal direction, in accordance with standard usage and convention.

[000702] Similarly, unless specified otherwise, the left-hand end of single- stranded polynucleotide sequences 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' addition of nascent RNA transcripts is referred to as the transcription direction sequence regions on the DNA strand having the same sequence as the RNA and 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 and that are 3' to the 3' end of the RNA transcript are referred to as "downstream sequences".

[000703] As applied to polypeptides, the term "substantial identity" means that two peptide sequences, when optimally aligned, such as by the programs GAP or BESTFIT using default gap weights, share at least 80 percent sequence identity, in some embodiments, at least 90 percent sequence identity, in some embodiments, at least 95 percent sequence identity, and in some embodiments, at least 99 percent sequence identity.

[000704] In some embodiments, residue positions that are not identical differ by conservative amino acid substitutions.

[000705] As discussed herein, minor variations in the amino acid sequences of antibodies or immunoglobulin molecules are contemplated as being encompassed by the present disclosure,

providing that the variations in the amino acid sequence maintain at least 75%, in some embodiments, at least 80%, 90%, 95%, and in some embodiments, 99%. In particular, conservative amino acid replacements are contemplated. Conservative replacements are those that take place within a family of amino acids that are related in their side chains. Genetically encoded amino acids are generally divided into families: (1) acidic amino acids are aspartate, glutamate; (2) basic amino acids are lysine, arginine, histidine; (3) non-polar amino acids are alanine, valine, leucine, isoleucine, proline, phenylalanine, methionine, tryptophan, and (4) uncharged polar amino acids are glycine, asparagine, glutamine, cysteine, serine, threonine, tyrosine. The hydrophilic amino acids include arginine, asparagine, aspartate, glutamine, glutamate, histidine, lysine, serine, and threonine. The hydrophobic amino acids include alanine, cysteine, isoleucine, leucine, methionine, phenylalanine, proline, tryptophan, tyrosine and valine. Other families of amino acids include (i) serine and threonine, which are the aliphatic-hydroxy family; (ii) asparagine and glutamine, which are the amide containing family; (iii) alanine, valine, leucine and isoleucine, which are the aliphatic family; and (iv) phenylalanine, tryptophan, and tyrosine, which are the aromatic family. For example, it is reasonable to expect that an isolated replacement of a leucine with an isoleucine or valine, an aspartate with a glutamate, a threonine with a serine, or a similar replacement of an amino acid with a structurally related amino acid will not have a major effect on the binding or properties of the resulting molecule, especially if the replacement does not involve an amino acid within a framework site. Whether an amino acid change results in a functional peptide can readily be determined by assaying the specific activity of the polypeptide derivative. Assays are described in detail herein. Fragments or analogs of antibodies or immunoglobulin molecules can be readily prepared by those of ordinary skill in the art. Suitable amino- and carboxy-termini of fragments or analogs occur near boundaries of functional domains. Structural and functional domains can be identified by comparison of the nucleotide and/or amino acid sequence data to public or proprietary sequence databases. In some embodiments, computerized comparison methods are used to identify sequence motifs or predicted protein conformation domains that occur in other proteins of known structure and/or function. Methods to identify protein sequences that fold into a known three-dimensional structure are known. Bowie *et al.* Science 253:164 (1991). Thus, the foregoing examples demonstrate that those of skill in the art can recognize sequence motifs and structural

conformations that can be used to define structural and functional domains in accordance with the disclosure.

[000706] Suitable amino acid substitutions are those that: (1) reduce susceptibility to proteolysis, (2) reduce susceptibility to oxidation, (3) alter binding affinity for forming protein complexes, (4) alter binding affinities, and (5) confer or modify other physicochemical or functional properties of such analogs. Analogs can include various muteins of a sequence other than the naturally-occurring peptide sequence. For example, single or multiple amino acid substitutions (for example, conservative amino acid substitutions) can be made in the naturally-occurring sequence (for example, in the portion of the polypeptide outside the domain(s) forming intermolecular contacts. A conservative amino acid substitution should not substantially change the structural characteristics of the parent sequence (*e.g.*, a replacement amino acid should not tend to break a helix that occurs in the parent sequence, or disrupt other types of secondary structure that characterizes the parent sequence). Examples of art-recognized polypeptide secondary and tertiary structures are described in *Proteins, Structures and Molecular Principles* (Creighton, Ed., W. H. Freeman and Company, New York (1984)); *Introduction to Protein Structure* (C. Branden and J. Tooze, eds., Garland Publishing, New York, N.Y. (1991)); and Thornton et al. *Nature* 354:105 (1991).

[000707] The term “polypeptide fragment” as used herein refers to a polypeptide that has an amino terminal and/or carboxy-terminal deletion and/or one or more internal deletion(s), but where the remaining amino acid sequence is identical to the corresponding positions in the naturally-occurring sequence deduced, for example, from a full length cDNA sequence. Fragments typically are at least 5, 6, 8 or 10 amino acids long, in some embodiments, at least 14 amino acids long, in some embodiments, at least 20 amino acids long, usually at least 50 amino acids long, and in some embodiments, at least 70 amino acids long. The term “analog” as used herein refers to polypeptides that are comprised of a segment of at least 25 amino acids that has substantial identity to a portion of a deduced amino acid sequence and that has specific binding to the target, under suitable binding conditions. Typically, polypeptide analogs comprise a conservative amino acid substitution (or addition or deletion) with respect to the naturally-occurring sequence. Analogs typically are at least 20 amino acids long, in some embodiments, at least 50 amino acids long or longer, and can often be as long as a full-length naturally-occurring polypeptide.

[000708] The term “agent” is used herein to denote a chemical compound, a mixture of chemical compounds, a biological macromolecule, or an extract made from biological materials.

[000709] As used herein, the terms “label” or “labeled” refers to incorporation of a detectable marker, *e.g.*, by incorporation of a radiolabeled amino acid or attachment to a polypeptide of biotinyl moieties that can be detected by marked avidin (*e.g.*, streptavidin containing a fluorescent marker or enzymatic activity that can be detected by optical or calorimetric methods). In certain situations, the label or marker can also be therapeutic. Various methods of labeling polypeptides and glycoproteins are known in the art and can be used. Examples of labels for polypeptides include, but are not limited to, the following: radioisotopes or radionuclides (*e.g.*, ^3H , ^{14}C , ^{15}N , ^{35}S , ^{90}Y , ^{99}Tc , ^{111}In , ^{125}I , ^{131}I), fluorescent labels (*e.g.*, FITC, rhodamine, lanthanide phosphors), enzymatic labels (*e.g.*, horseradish peroxidase, p-galactosidase, luciferase, alkaline phosphatase), chemiluminescent, biotinyl groups, predetermined polypeptide epitopes recognized by a secondary reporter (*e.g.*, leucine zipper pair sequences, binding sites for secondary antibodies, metal binding domains, epitope tags). In some embodiments, labels are attached by spacer arms of various lengths to reduce potential steric hindrance. The term “pharmaceutical agent or drug” as used herein refers to a chemical compound or composition capable of inducing a desired therapeutic effect when properly administered to a patient.

[000710] Other chemistry terms herein are used according to conventional usage in the art, as exemplified by The McGraw-Hill Dictionary of Chemical Terms (Parker, S., Ed., McGraw-Hill, San Francisco (1985)).

[000711] As used herein, “substantially pure” means an object species is the predominant species present (*i.e.*, on a molar basis it is more abundant than any other individual species in the composition), and in some embodiments, a substantially purified fraction is a composition wherein the object species comprises at least about 50 percent (on a molar basis) of all macromolecular species present.

[000712] Generally, a substantially pure composition will comprise more than about 80 percent of all macromolecular species present in the composition, in some embodiments, more than about 85%, 90%, 95%, and 99%. In some embodiments, the object species is purified to essential homogeneity (contaminant species cannot be detected in the composition by

conventional detection methods) wherein the composition consists essentially of a single macromolecular species.

[000713] The term patient includes human and veterinary subjects.

[000714] Antibodies and/or activatable antibodies of the disclosure specifically bind a given target, e.g., a human target protein such as human PD-1. Also included in the disclosure are antibodies and/or activatable antibodies that bind to the same epitope as the antibodies and/or activatable antibodies described herein. Also included in the disclosure are antibodies and/or antibodies activatable antibodies that compete with an anti-PD-1 antibody and/or an anti-PD-1 activatable antibody described herein for binding to PD-1, e.g., human PD-1. Also included in the disclosure are antibodies and/or antibodies activatable antibodies that cross-compete with an anti-PD-1 antibody and/or an anti-PD-1 activatable antibody described herein for binding to PD-1, e.g., human PD-1.

[000715] Those skilled in the art will recognize that it is possible to determine, without undue experimentation, if a monoclonal antibody (*e.g.*, a murine monoclonal or humanized antibody) has the same specificity as a monoclonal antibody used in the methods described herein by ascertaining whether the former prevents the latter from binding to the target. If the monoclonal antibody being tested competes with the monoclonal antibody of the disclosure, as shown by a decrease in binding by the monoclonal antibody of the disclosure, then the two monoclonal antibodies bind to the same, or a closely related, epitope. An alternative method for determining whether a monoclonal antibody has the specificity of a monoclonal antibody of the disclosure is to pre-incubate the monoclonal antibody of the disclosure with the target and then add the monoclonal antibody being tested to determine if the monoclonal antibody being tested is inhibited in its ability to bind the target. If the monoclonal antibody being tested is inhibited then, in all likelihood, it has the same, or functionally equivalent, epitopic specificity as the monoclonal antibody of the disclosure.

Multispecific Activatable Antibodies

[000716] The disclosure also provides multispecific anti-PD-1 activatable antibodies. The multispecific activatable antibodies provided herein are multispecific antibodies that recognize PD-1 and at least one or more different antigens or epitopes and that include at least one masking moiety (MM) linked to at least one antigen- or epitope-binding domain of the multispecific antibody such that coupling of the MM reduces the ability of the antigen- or epitope-binding

domain to bind its target. In some embodiments, the MM is coupled to the antigen- or epitope-binding domain of the multispecific antibody via a cleavable moiety (CM) that functions as a substrate for at least one protease. The activatable multispecific antibodies provided herein are stable in circulation, activated at intended sites of therapy and/or diagnosis but not in normal, *i.e.*, healthy tissue, and, when activated, exhibit binding to a target that is at least comparable to the corresponding, unmodified multispecific antibody.

[000717] In some embodiments, the multispecific activatable antibodies are designed to engage immune effector cells, also referred to herein as immune-effector cell engaging multispecific activatable antibodies. In some embodiments, the multispecific activatable antibodies are designed to engage leukocytes, also referred to herein as leukocyte engaging multispecific activatable antibodies. In some embodiments, the multispecific activatable antibodies are designed to engage T cells, also referred to herein as T-cell engaging multispecific activatable antibodies. In some embodiments, the multispecific activatable antibodies engage a surface antigen on a leukocyte, such as on a T cell, on a natural killer (NK) cell, on a myeloid mononuclear cell, on a macrophage, and/or on another immune effector cell. In some embodiments, the immune effector cell is a leukocyte. In some embodiments, the immune effector cell is a T cell. In some embodiments, the immune effector cell is a NK cell. In some embodiments, the immune effector cell is a mononuclear cell, such as a myeloid mononuclear cell. In some embodiments, the multispecific activatable antibodies are designed to bind or otherwise interact with more than one target and/or more than one epitope, also referred to herein as multi-antigen targeting activatable antibodies. As used herein, the terms “target” and “antigen” are used interchangeably.

[000718] In some embodiments, immune effector cell engaging multispecific activatable antibodies of the disclosure include a targeting antibody or antigen-binding fragment thereof that binds PD-1 and an immune effector cell engaging antibody or antigen-binding portion thereof, where at least one of the targeting antibody or antigen-binding fragment thereof and/or the immune effector cell engaging antibody or antigen-binding portion thereof is masked. In some embodiments, the immune effector cell engaging antibody or antigen binding fragment thereof includes a first antibody or antigen-binding fragment thereof (AB1) that binds a first, immune effector cell engaging target, where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind the first target. In some

embodiments, the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1. In some embodiments, the immune effector cell engaging antibody or antigen binding fragment thereof includes a first antibody or antigen-binding fragment thereof (AB1) that binds a first, immune effector cell engaging target, where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind the first target, and the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1. In some embodiments, the non-immune effector cell engaging antibody is a cancer targeting antibody. In some embodiments the non-immune cell effector antibody is an IgG. In some embodiments the immune effector cell engaging antibody is a scFv. In some embodiments the PD-1-targeting antibody (e.g., non-immune cell effector antibody) is an IgG and the immune effector cell engaging antibody is a scFv. In some embodiments, the immune effector cell is a leukocyte. In some embodiments, the immune effector cell is a T cell. In some embodiments, the immune effector cell is a NK cell. In some embodiments, the immune effector cell is a myeloid mononuclear cell.

[000719] In some embodiments, T-cell engaging multispecific activatable antibodies of the disclosure include a PD-1-targeting antibody or antigen-binding fragment thereof and a T-cell engaging antibody or antigen-binding portion thereof, where at least one of the PD-1-targeting antibody or antigen-binding fragment thereof and/or the T-cell engaging antibody or antigen-binding portion thereof is masked. In some embodiments, the T-cell engaging antibody or antigen binding fragment thereof includes a first antibody or antigen-binding fragment thereof (AB1) that binds a first, T-cell engaging target, where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind the first target. In some embodiments, the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1. In some embodiments, the T-

cell engaging antibody or antigen binding fragment thereof includes a first antibody or antigen-binding fragment thereof (AB1) that binds a first, T-cell engaging target, where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind the first target, and the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1.

[000720] In some embodiments of an immune effector cell engaging multispecific activatable antibody, one antigen is PD-1, and another antigen is typically a stimulatory or inhibitory receptor present on the surface of a T-cell, natural killer (NK) cell, myeloid mononuclear cell, macrophage, and/or other immune effector cell, such as, but not limited to, B7-H4, BTLA, CD3, CD4, CD8, CD16a, CD25, CD27, CD28, CD32, CD56, CD137, CTLA-4, GITR, HVEM, ICOS, LAG3, NKG2D, OX40, TIGIT, TIM3, or VISTA. In some embodiments, the antigen is a stimulatory receptor present on the surface of a T cell or NK cell; examples of such stimulatory receptors include, but are not limited to, CD3, CD27, CD28, CD137 (also referred to as 4-1BB), GITR, HVEM, ICOS, NKG2D, and OX40. In some embodiments, the antigen is an inhibitory receptor present on the surface of a T-cell; examples of such inhibitory receptors include, but are not limited to, BTLA, CTLA-4, LAG3, TIGIT, TIM3, and NK-expressed KIRs. The antibody domain conferring specificity to the T-cell surface antigen can also be substituted by a ligand or ligand domain that binds to a T-cell receptor, a NK-cell receptor, a macrophage receptor, and/or other immune effector cell receptor, such as, but not limited to, B7-1, B7-2, B7H3, PDL1, PDL2, or TNFSF9.

[000721] In some embodiments, the T-cell engaging multispecific activatable antibody includes an anti-CD3 epsilon (CD3 ϵ , also referred to herein as CD3e and CD3) scFv and a targeting antibody or antigen-binding fragment thereof, where at least one of the anti-CD3 ϵ scFv and/or the targeting antibody or antigen-binding portion thereof is masked. In some embodiments, the CD3 ϵ scFv includes a first antibody or antigen-binding fragment thereof (AB1) that binds CD3 ϵ , where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind CD3 ϵ . In some embodiments, the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds

PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1. In some embodiments, the CD3 ϵ scFv includes a first antibody or antigen-binding fragment thereof (AB1) that binds CD3 ϵ , where the AB1 is attached to a masking moiety (MM1) such that coupling of the MM1 reduces the ability of the AB1 to bind CD3 ϵ , and the targeting antibody or antigen-binding fragment thereof includes a second antibody or fragment thereof that includes a second antibody or antigen-binding fragment thereof (AB2) that binds PD-1, where the AB2 is attached to a masking moiety (MM2) such that coupling of the MM2 reduces the ability of the AB2 to bind PD-1.

[000722] In some embodiments, the multi-antigen targeting antibodies and/or multi-antigen targeting activatable antibodies include at least a first antibody or antigen-binding fragment thereof that binds a first target and/or first epitope and a second antibody or antigen-binding fragment thereof that binds a second target and/or a second epitope. In some embodiments, the multi-antigen targeting antibodies and/or multi-antigen targeting activatable antibodies bind two or more different targets. In some embodiments, the multi-antigen targeting antibodies and/or multi-antigen targeting activatable antibodies bind two or more different epitopes on the same target. In some embodiments, the multi-antigen targeting antibodies and/or multi-antigen targeting activatable antibodies bind a combination of two or more different targets and two or more different epitopes on the same target.

[000723] In some embodiments, a multispecific activatable antibody comprising an IgG has the IgG variable domains masked. In some embodiments, a multispecific activatable antibody comprising a scFv has the scFv domains masked. In some embodiments, a multispecific activatable antibody has both IgG variable domains and scFv domains, where at least one of the IgG variable domains is coupled to a masking moiety. In some embodiments, a multispecific activatable antibody has both IgG variable domains and scFv domains, where at least one of the scFv domains is coupled to a masking moiety. In some embodiments, a multispecific activatable antibody has both IgG variable domains and scFv domains, where at least one of the IgG variable domains is coupled to a masking moiety and at least one of the scFv domains is coupled to a masking moiety. In some embodiments, a multispecific activatable antibody has both IgG variable domains and scFv domains, where each of the IgG variable domains and the scFv domains is coupled to its own masking moiety. In some embodiments, one antibody domain of a multispecific activatable antibody has specificity for a target antigen and another antibody

domain has specificity for a T-cell surface antigen. In some embodiments, one antibody domain of a multispecific activatable antibody has specificity for a target antigen and another antibody domain has specificity for another target antigen. In some embodiments, one antibody domain of a multispecific activatable antibody has specificity for an epitope of a target antigen and another antibody domain has specificity for another epitope of the target antigen.

[000724] In a multispecific activatable antibody, a scFv can be fused to the carboxyl terminus of the heavy chain of an IgG activatable antibody, to the carboxyl terminus of the light chain of an IgG activatable antibody, or to the carboxyl termini of both the heavy and light chains of an IgG activatable antibody. In a multispecific activatable antibody, a scFv can be fused to the amino terminus of the heavy chain of an IgG activatable antibody, to the amino terminus of the light chain of an IgG activatable antibody, or to the amino termini of both the heavy and light chains of an IgG activatable antibody. In a multispecific activatable antibody, a scFv can be fused to any combination of one or more carboxyl termini and one or more amino termini of an IgG activatable antibody. In some embodiments, a masking moiety (MM) linked to a cleavable moiety (CM) is attached to and masks an antigen binding domain of the IgG. In some embodiments, a masking moiety (MM) linked to a cleavable moiety (CM) is attached to and masks an antigen binding domain of at least one scFv. In some embodiments, a masking moiety (MM) linked to a cleavable moiety (CM) is attached to and masks an antigen binding domain of an IgG and a masking moiety (MM) linked to a cleavable moiety (CM) is attached to and masks an antigen binding domain of at least one scFv.

[000725] The disclosure provides examples of multispecific activatable antibody structures which include, but are not limited to, the following: (VL-CL)₂:(VH-CH1-CH2-CH3-L4-VH*-L3-VL*-L2-CM-L1-MM)₂; (VL-CL)₂:(VH-CH1-CH2-CH3-L4-VL*-L3-VH*-L2-CM-L1-MM)₂; (MM-L1-CM-L2-VL-CL)₂:(VH-CH1-CH2-CH3-L4-VH*-L3-VL*)₂; (MM-L1-CM-L2-VL-CL)₂:(VH-CH1-CH2-CH3-L4-VL*-L3-VH*)₂; (VL-CL)₂:(MM-L1-CM-L2-VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL)₂:(MM-L1-CM-L2-VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (MM-L1-CM-L2-VL-CL)₂:(VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (MM-L1-CM-L2-VL-CL)₂:(VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*-L2-CM-L1-MM)₂:(VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*-L2-CM-L1-MM)₂:(VH-CH1-CH2-CH3)₂; (MM-L1-CM-L2-VL*-L3-VH*-L4-VL-CL)₂:(VH-CH1-CH2-CH3)₂; (MM-L1-CM-L2-VH*-L3-VL*-L4-VL-CL)₂:(VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*-L2-CM-L1-

MM)₂: (MM-L1-CM-L2-VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*-L2-CM-L1-MM)₂: (MM-L1-CM-L2-VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*-L2-CM-L1-MM)₂: (MM-L1-CM-L2-VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*-L2-CM-L1-MM)₂: (MM-L1-CM-L2-VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*)₂: (MM-L1-CM-L2-VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*)₂: (MM-L1-CM-L2-VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*)₂: (MM-L1-CM-L2-VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*)₂: (MM-L1-CM-L2-VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*-L2-CM-L1-MM)₂: (VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VH*-L3-VL*-L2-CM-L1-MM)₂: (VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂; (VL-CL-L4-VL*-L3-VH*-L2-CM-L1-MM)₂: (VL*-L3-VH*-L4-VH-CH1-CH2-CH3)₂; or (VL-CL-L4-VL*-L3-VH*-L2-CM-L1-MM)₂: (VH*-L3-VL*-L4-VH-CH1-CH2-CH3)₂, wherein: VL and VH represent the light and heavy variable domains of the first specificity, contained in the IgG; VL* and VH* represent the variable domains of the second specificity, contained in the scFv; L1 is a linker peptide connecting the masking moiety (MM) and the cleavable moiety (CM); L2 is a linker peptide connecting the cleavable moiety (CM), and the antibody; L3 is a linker peptide connecting the variable domains of the scFv; L4 is a linker peptide connecting the antibody of the first specificity to the antibody of the second specificity; CL is the light-chain constant domain; and CH1, CH2, CH3 are the heavy chain constant domains. The first and second specificities can be toward any antigen or epitope.

[000726] In some embodiments of a T-cell engaging multispecific activatable antibody, one antigen is PD-1, and another antigen is typically a stimulatory (also referred to herein as activating) or inhibitory receptor present on the surface of a T-cell, natural killer (NK) cell, myeloid mononuclear cell, macrophage, and/or other immune effector cell, such as, but not limited to, B7-H4, BTLA, CD3, CD4, CD8, CD16a, CD25, CD27, CD28, CD32, CD56, CD137 (also referred to as TNFRSF9), CTLA-4, GITR, HVEM, ICOS, LAG3, NKG2D, OX40, PD-1, TIGIT, TIM3, or VISTA. The antibody domain conferring specificity to the T-cell surface antigen can also be substituted by a ligand or ligand domain that binds to a T-cell receptor, a NK-cell receptor, a macrophage receptor, and/or other immune effector cell receptor.

[000727] In some embodiments, the targeting antibody is an anti-PD-1 antibody disclosed herein. In some embodiments, the targeting antibody can be in the form an activatable antibody.

In some embodiments, the scFv(s) can be in the form of a Pro-scFv (see, e.g., WO 2009/025846, WO 2010/081173).

[000728] In some embodiments, the scFv is specific for binding CD3 ϵ , and comprises or is derived from an antibody or fragment thereof that binds CD3 ϵ , e.g., CH2527, FN18, H2C, OKT3, 2C11, UCHT1, or V9. In some embodiments, the scFv is specific for binding CTLA-4 (also referred to herein as CTLA and CTLA4).

[000729] In some embodiments, the anti-CTLA-4 scFv includes the amino acid sequence:

GGGSGGGGSGSGGGSGGGGSGGGGEIVLTQSPGTLSSLSPGERATLSCRASQSVSSSYLAWYQQKP
GQAPRLLIYGASSRATGIPDRFSGSGSGTDFTLTISRLEPEDFAVYYCQQYGSSPLTFGGGTKV
EIKRSGGSTITSYNVYYTKLSSSGTQVQLVQVTGGGVVQPGRSLRLSCAASGSTFSSYAMSWVRQ
APGKGLEWVSAISGSGGSTYYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCATNSLYW
YFDLWGRGTLTVSSAS (SEQ ID NO: 585)

[000730] In some embodiments, the anti-CTLA-4 scFv includes the amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the amino acid sequence of SEQ ID NO: 585.

[000731] In some embodiments, the anti-CD3 ϵ scFv includes the amino acid sequence:

GGGSGGGGSGSGGGSGGGGSGGGGQVQLQQSGAELARPGASVKMSCKASGYTFTRYTMHWVKQRP
GQGLEWIGYINPSRGYTNYNQKFKDKATLTDDKSSSTAYMQLSSLTSEDSAVYYCARYYDDHYC
LDYWGQGTTLTVSSGGGGSGGGGSGGGGSQIVLTQSPAISASPGKEKVTMTCSASSSVSYMNWY
QQKSGTSPKRWIYDTSKLASGVPAHFRGSGSGTSYSLTISGMEAEDAATYYCQQWSSNPFTFGS
GTKLEINR (SEQ ID NO: 586)

[000732] In some embodiments, the anti-CD3 ϵ scFv includes the amino acid sequence that is at least 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99% or more identical to the amino acid sequence of SEQ ID NO: 586.

[000733] In some embodiments, the scFv is specific for binding one or more T-cells, one or more NK-cells and/or one or more macrophages. In some embodiments, the scFv is specific for binding a target selected from the group consisting of B7-H4, BTLA, CD3, CD4, CD8, CD16a, CD25, CD27, CD28, CD32, CD56, CD137, CTLA-4, GITR, HVEM, ICOS, LAG3, NKG2D, OX40, TIGIT, TIM3, or VISTA.

[000734] In some embodiments, the multispecific activatable antibody also includes an agent conjugated to the AB. In some embodiments, the agent is a therapeutic agent. In some embodiments, the agent is conjugated to the multispecific activatable antibody via a linker. In some embodiments, the agent is conjugated to the AB via a cleavable linker. In some embodiments, the linker is a non-cleavable linker.

[000735] In some embodiments, the multispecific activatable antibody also includes a detectable moiety. In some embodiments, the detectable moiety is a diagnostic agent.

[000736] In some embodiments, the multispecific activatable antibody naturally contains one or more disulfide bonds. In some embodiments, the multispecific activatable antibody can be engineered to include one or more disulfide bonds.

[000737] The disclosure also provides an isolated nucleic acid molecule encoding a multispecific activatable antibody described herein, as well as vectors that include these isolated nucleic acid sequences. The disclosure provides methods of producing a multispecific activatable antibody by culturing a cell under conditions that lead to expression of the activatable antibody, wherein the cell comprises such a nucleic acid molecule. In some embodiments, the cell comprises such a vector.

[000738] The disclosure also provides a method of manufacturing multispecific activatable antibodies of the disclosure by (a) culturing a cell comprising a nucleic acid construct that encodes the multispecific activatable antibody under conditions that lead to expression of the multispecific activatable, and (b) recovering the multispecific activatable antibody. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

[000739] The disclosure also provides multispecific activatable antibodies and/or multispecific activatable antibody compositions that include at least a first antibody or antigen-binding fragment thereof (AB1) that specifically binds a first target or first epitope and a second antibody or antigen-binding fragment thereof (AB2) that binds a second target or a second epitope, where at least AB1 is coupled or otherwise attached to a masking moiety (MM1), such that coupling of the MM1 reduces the ability of AB1 to bind its target. In some embodiments, the MM1 is coupled to AB1 via a first cleavable moiety (CM1) sequence that includes a substrate for a protease, for example, a protease that is co-localized with the target of AB1 at a treatment site or a diagnostic site in a subject. The multispecific activatable antibodies provided herein are stable in circulation, activated at intended sites of therapy and/or diagnosis but not in normal, *i.e.*,

healthy tissue, and, when activated, exhibit binding to the target of AB1 that is at least comparable to the corresponding, unmodified multispecific antibody. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

[000740] The disclosure also provides compositions and methods that include a multispecific activatable antibody that includes at least a first antibody or antibody fragment (AB1) that specifically binds a target and a second antibody or antibody fragment (AB2), where at least the first AB in the multispecific activatable antibody is coupled to a masking moiety (MM1) that decreases the ability of AB1 to bind its target. In some embodiments, each AB is coupled to a MM that decreases the ability of its corresponding AB to each target. For example, in bispecific activatable antibody embodiments, AB1 is coupled to a first masking moiety (MM1) that decreases the ability of AB1 to bind its target, and AB2 is coupled to a second masking moiety (MM2) that decreases the ability of AB2 to bind its target. In some embodiments, the multispecific activatable antibody comprises more than two AB regions; in such embodiments, AB1 is coupled to a first masking moiety (MM1) that decreases the ability of AB1 to bind its target, AB2 is coupled to a second masking moiety (MM2) that decreases the ability of AB2 to bind its target, AB3 is coupled to a third masking moiety (MM3) that decreases the ability of AB3 to bind its target, and so on for each AB in the multispecific activatable antibody. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

[000741] In some embodiments, the multispecific activatable antibody further includes at least one cleavable moiety (CM) that is a substrate for a protease, where the CM links a MM to an AB. For example, in some embodiments, the multispecific activatable antibody includes at least a first antibody or antibody fragment (AB1) that specifically binds a target and a second antibody or antibody fragment (AB2), where at least the first AB in the multispecific activatable antibody is coupled via a first cleavable moiety (CM1) to a masking moiety (MM1) that decreases the ability of AB1 to bind its target. In some bispecific activatable antibody embodiments, AB1 is coupled via CM1 to MM1, and AB2 is coupled via a second cleavable moiety (CM2) to a second masking moiety (MM2) that decreases the ability of AB2 to bind its target. In some embodiments, the multispecific activatable antibody comprises more than two AB regions; in some of these embodiments, AB1 is coupled via CM1 to MM1, AB2 is coupled via CM2 to MM2, and AB3 is coupled via a third cleavable moiety (CM3) to a third masking

moiety (MM3) that decreases the ability of AB3 to bind its target, and so on for each AB in the multispecific activatable antibody. Suitable AB, MM, and/or CM include any of the AB, MM, and/or CM disclosed herein.

Activatable antibodies Having Non-Binding Steric Moieties or Binding Partners for Non-Binding Steric Moieties

[000742] The disclosure also provides activatable antibodies that include non-binding steric moieties (NB) or binding partners (BP) for non-binding steric moieties, where the BP recruits or otherwise attracts the NB to the activatable antibody. The activatable antibodies provided herein include, for example, an activatable antibody that includes a non-binding steric moiety (NB), a cleavable linker (CL) and antibody or antibody fragment (AB) that binds a target; an activatable antibody that includes a binding partner for a non-binding steric moiety (BP), a CL and an AB; and an activatable antibody that includes a BP to which an NB has been recruited, a CL and an AB that binds the target. Activatable antibodies in which the NB is covalently linked to the CL and AB of the activatable antibody or is associated by interaction with a BP that is covalently linked to the CL and AB of the activatable antibody are referred to herein as “NB-containing activatable antibodies.” By activatable or switchable is meant that the activatable antibody exhibits a first level of binding to a target when the activatable antibody is in an inhibited, masked or uncleaved state (*i.e.*, a first conformation), and a second level of binding to the target when the activatable antibody is in an uninhibited, unmasked and/or cleaved state (*i.e.*, a second conformation, *i.e.*, activated antibody), where the second level of target binding is greater than the first level of target binding. The activatable antibody compositions can exhibit increased bioavailability and more favorable biodistribution compared to conventional antibody therapeutics.

[000743] In some embodiments, activatable antibodies provide for reduced toxicity and/or adverse side effects that could otherwise result from binding of the at non-treatment sites and/or non-diagnostic sites if the AB were not masked or otherwise inhibited from binding to such a site.

[000744] Anti-PD-1 activatable antibodies that include a non-binding steric moiety (NB) can be made using the methods set forth in PCT Publication No. WO 2013/192546, the contents of which are hereby incorporated by reference in their entirety.

Use Of Antibodies and Activatable Antibodies

[000745] It will be appreciated that administration of therapeutic entities in accordance with the disclosure will be administered with suitable carriers, excipients, and other agents that are incorporated into formulations to provide improved transfer, delivery, tolerance, and the like. A multitude of appropriate formulations can be found in the formulary known to all pharmaceutical chemists: Remington's Pharmaceutical Sciences (15th ed, Mack Publishing Company, Easton, PA (1975)), particularly Chapter 87 by Blaug, Seymour, therein. These formulations include, for example, powders, pastes, ointments, jellies, waxes, oils, lipids, lipid (cationic or anionic) containing vesicles (such as Lipofectin™), DNA conjugates, anhydrous absorption pastes, oil-in-water and water-in-oil emulsions, emulsions carbowax (polyethylene glycols of various molecular weights), semi-solid gels, and semi-solid mixtures containing carbowax. Any of the foregoing mixtures can be appropriate in treatments and therapies in accordance with the present disclosure, provided that the active ingredient in the formulation is not inactivated by the formulation and the formulation is physiologically compatible and tolerable with the route of administration. *See also* Baldrick P. "Pharmaceutical excipient development: the need for preclinical guidance." Regul. Toxicol Pharmacol. 32(2):210-8 (2000), Wang W. "Lyophilization and development of solid protein pharmaceuticals." Int. J. Pharm. 203(1-2):1-60 (2000), Charman WN "Lipids, lipophilic drugs, and oral drug delivery-some emerging concepts." J Pharm Sci. 89(8):967-78 (2000), Powell *et al.* "Compendium of excipients for parenteral formulations" PDA J Pharm Sci Technol. 52:238-311 (1998) and the citations therein for additional information related to formulations, excipients and carriers well known to pharmaceutical chemists.

[000746] Therapeutic formulations of the disclosure, which include an anti-PD-1 antibody and/or activatable anti-PD-1 antibody, such as by way of non-limiting example, an antibody and/or an activatable antibody, are used to prevent, treat or otherwise ameliorate a disease or disorder associated with aberrant target expression and/or activity. For example, therapeutic formulations of the disclosure, which include an antibody and/or an activatable antibody, are used to treat or otherwise ameliorate a cancer or other neoplastic condition, inflammation, an inflammatory disorder, and/or an autoimmune disease. In some embodiments, the cancer is a solid tumor or a hematologic malignancy where the target is expressed. In some embodiments, the cancer is a solid tumor where the target is expressed. In some embodiments, the cancer is a

hematologic malignancy where the target is expressed. In some embodiments, the target is expressed on parenchyma (e.g., in cancer, the portion of an organ or tissue that often carries out function(s) of the organ or tissue). In some embodiments, the target is expressed on a cell, tissue, or organ. In some embodiments, the target is expressed on stroma (i.e., the connective supportive framework of a cell, tissue, or organ). In some embodiments, the target is expressed on an osteoblast. In some embodiments, the target is expressed on the endothelium (vasculature). In some embodiments, the target is expressed on a cancer stem cell.

[000747] Efficaciousness of prevention, amelioration or treatment is determined in association with any known method for diagnosing or treating the disease or disorder associated with target expression and/or activity, such as, for example, aberrant target expression and/or activity. Prolonging the survival of a subject or otherwise delaying the progression of the disease or disorder associated with target expression and/or activity, e.g., aberrant target expression and/or activity, in a subject indicates that the antibody and/or activatable antibody confers a clinical benefit.

[000748] An antibody and/or an activatable antibody can be administered in the form of pharmaceutical compositions. Principles and considerations involved in preparing such compositions, as well as guidance in the choice of components are provided, for example, in Remington : The Science And Practice Of Pharmacy 19th ed. (Alfonso R. Gennaro, et al., editors) Mack Pub. Co., Easton, Pa.: 1995; Drug Absorption Enhancement: Concepts, Possibilities, Limitations, And Trends, Harwood Academic Publishers, Langhorne, Pa., 1994; and Peptide And Protein Drug Delivery (Advances In Parenteral Sciences, Vol. 4), 1991, M. Dekker, New York.

[000749] In some embodiments where antibody fragments are used, the smallest fragment that specifically binds to the binding domain of the target protein is selected. For example, based upon the variable-region sequences of an antibody, peptide molecules can be designed that retain the ability to bind the target protein sequence. Such peptides can be synthesized chemically and/or produced by recombinant DNA technology. (See, e.g., Marasco et al., Proc. Natl. Acad. Sci. USA, 90: 7889-7893 (1993)). The formulation can also contain more than one active compounds as necessary for the particular indication being treated, for example, in some embodiments, those with complementary activities that do not adversely affect each other. In some embodiments, or in addition, the composition can comprise an agent that enhances its

function, such as, for example, a cytotoxic agent, cytokine, chemotherapeutic agent, or growth-inhibitory agent. Such molecules are suitably present in combination in amounts that are effective for the purpose intended.

[000750] The active ingredients can also be entrapped in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization, for example, hydroxymethylcellulose or gelatin-microcapsules and poly-(methacrylate) microcapsules, respectively, in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles, and nanocapsules) or in macroemulsions.

[000751] The formulations to be used for *in vivo* administration must be sterile. This is readily accomplished by filtration through sterile filtration membranes.

[000752] Sustained-release preparations can be prepared. Suitable examples of sustained-release preparations include semipermeable matrices of solid hydrophobic polymers containing the antibody, which matrices are in the form of shaped articles, *e.g.*, films, or microcapsules. Examples of sustained-release matrices include polyesters, hydrogels (for example, poly(2-hydroxyethyl-methacrylate), or poly(vinylalcohol)), polylactides (U.S. Pat. No. 3,773,919), copolymers of L-glutamic acid and γ ethyl-L-glutamate, non-degradable ethylene-vinyl acetate, degradable lactic acid-glycolic acid copolymers such as the LUPRON DEPOTTM (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and poly-D-(-)-3-hydroxybutyric acid. While polymers such as ethylene-vinyl acetate and lactic acid-glycolic acid enable release of molecules for over 100 days, certain hydrogels release proteins for shorter time periods.

[000753] In some embodiments, the antibody and/or activatable antibody contains a detectable label. An intact antibody, or a fragment thereof (*e.g.*, Fab, scFv, or F(ab)₂) is used. The term “labeled”, with regard to the probe or antibody, is intended to encompass direct labeling of the probe or antibody by coupling (*i.e.*, physically linking) a detectable substance to the probe or antibody, as well as indirect labeling of the probe or antibody by reactivity with another reagent that is directly labeled. Examples of indirect labeling include detection of a primary antibody using a fluorescently-labeled secondary antibody and end-labeling of a DNA probe with biotin such that it can be detected with fluorescently-labeled streptavidin. The term “biological sample” is intended to include tissues, cells and biological fluids isolated from a subject, as well as tissues, cells and fluids present within a subject. Included within the usage of

the term “biological sample”, therefore, is blood and a fraction or component of blood including blood serum, blood plasma, or lymph. That is, the detection method of the disclosure can be used to detect an analyte mRNA, protein, or genomic DNA in a biological sample *in vitro* as well as *in vivo*. For example, *in vitro* techniques for detection of an analyte mRNA include Northern hybridizations and *in situ* hybridizations. *In vitro* techniques for detection of an analyte protein include enzyme linked immunosorbent assays (ELISAs), Western blots, immunoprecipitations, immunochemical staining, and immunofluorescence. *In vitro* techniques for detection of an analyte genomic DNA include Southern hybridizations. Procedures for conducting immunoassays are described, for example in “ELISA: Theory and Practice: Methods in Molecular Biology”, Vol. 42, J. R. Crowther (Ed.) Human Press, Totowa, NJ, 1995; “Immunoassay”, E. Diamandis and T. Christopoulos, Academic Press, Inc., San Diego, CA, 1996; and “Practice and Theory of Enzyme Immunoassays”, P. Tijssen, Elsevier Science Publishers, Amsterdam, 1985. Furthermore, *in vivo* techniques for detection of an analyte protein include introducing into a subject a labeled anti-analyte protein antibody. For example, the antibody can be labeled with a radioactive marker whose presence and location in a subject can be detected by standard imaging techniques.

[000754] The antibodies and/or activatable antibodies of the disclosure are also useful in a variety of diagnostic and prophylactic formulations. In one embodiment, an antibody and/or an activatable antibody is administered to patients that are at risk of developing one or more of the aforementioned disorders. A patient’s or organ’s predisposition to one or more of the aforementioned disorders can be determined using genotypic, serological or biochemical markers.

[000755] In some embodiments of the disclosure, an antibody and/or an activatable antibody is administered to human individuals diagnosed with a clinical indication associated with one or more of the aforementioned disorders. Upon diagnosis, an antibody and/or an activatable antibody is administered to mitigate or reverse the effects of the clinical indication.

[000756] An antibody and/or an activatable antibody of the disclosure is also useful in the detection of a target in patient samples and accordingly are useful as diagnostics. For example, the antibodies and/or activatable antibodies, and conjugated versions thereof, of the disclosure are used in *in vitro* assays, *e.g.*, ELISA, to detect target levels in a patient sample.

[000757] In some embodiments, anti-PDL1 antibodies are used as a diagnostic for patients who are more likely to respond favorably to treatment with an anti-PD-1 antibody and/or activatable antibody of the disclosure. In these embodiments, PD-L1 expression in tumor biopsies on both tumor cells and infiltrating immune cells and PD-L1 expression on immune cells in blood are used to indicate the presence of active immunity and, thus, potential for suppression of T cell activity through the engagement of PD-1 by PD-L1.

[000758] In one embodiment, an antibody, a conjugated antibody, an activatable antibody and/or a conjugated activatable antibody of the disclosure is immobilized on a solid support (*e.g.*, the well(s) of a microtiter plate). The immobilized antibody, conjugated antibody, activatable antibody and/or conjugated activatable antibody serves as a capture antibody for any target that can be present in a test sample. Prior to contacting the immobilized antibody and/or activatable antibody, and/or conjugated versions thereof, with a patient sample, the solid support is rinsed and treated with a blocking agent such as milk protein or albumin to prevent nonspecific adsorption of the analyte.

[000759] Subsequently the wells are treated with a test sample suspected of containing the antigen, or with a solution containing a standard amount of the antigen. Such a sample is, *e.g.*, a serum sample from a subject suspected of having levels of circulating antigen considered to be diagnostic of a pathology. After rinsing away the test sample or standard, the solid support is treated with a second antibody that is detectably labeled. The labeled second antibody serves as a detecting antibody. The level of detectable label is measured, and the concentration of target antigen in the test sample is determined by comparison with a standard curve developed from the standard samples.

[000760] It will be appreciated that based on the results obtained using the antibodies and activatable antibodies of the disclosure, and conjugated versions thereof, in an *in vitro* diagnostic assay, it is possible to stage a disease in a subject based on expression levels of the target antigen. For a given disease, samples of blood are taken from subjects diagnosed as being at various stages in the progression of the disease, and/or at various points in the therapeutic treatment of the disease. Using a population of samples that provides statistically significant results for each stage of progression or therapy, a range of concentrations of the antigen that can be considered characteristic of each stage is designated.

[000761] An antibody, a conjugated antibody, an activatable antibody and/or a conjugated activatable antibody can also be used in diagnostic and/or imaging methods. In some embodiments, such methods are *in vitro* methods. In some embodiments, such methods are *in vivo* methods. In some embodiments, such methods are *in situ* methods. In some embodiments, such methods are *ex vivo* methods. For example, activatable antibodies having an enzymatically cleavable CM can be used to detect the presence or absence of an enzyme that is capable of cleaving the CM. Such activatable antibodies can be used in diagnostics, which can include *in vivo* detection (*e.g.*, qualitative or quantitative) of enzyme activity (or, in some embodiments, an environment of increased reduction potential such as that which can provide for reduction of a disulfide bond) through measured accumulation of activated antibodies (*i.e.*, antibodies resulting from cleavage of an activatable antibody) in a given cell or tissue of a given host organism. Such accumulation of activated antibodies indicates not only that the tissue expresses enzymatic activity (or an increased reduction potential depending on the nature of the CM) but also that the tissue expresses target to which the activated antibody binds.

[000762] For example, the CM can be selected to be substrate for at least one protease found at the site of a tumor, at the site of a viral or bacterial infection at a biologically confined site (*e.g.*, such as in an abscess, in an organ, and the like), and the like. The AB can be one that binds a target antigen. Using methods as disclosed herein, or when appropriate, methods familiar to one skilled in the art, a detectable label (*e.g.*, a fluorescent label or radioactive label or radiotracer) can be conjugated to an AB or other region of an antibody and/or activatable antibody. Suitable detectable labels are discussed in the context of the above screening methods and additional specific examples are provided below. Using an AB specific to a protein or peptide of the disease state, along with at least one protease whose activity is elevated in the disease tissue of interest, activatable antibodies will exhibit an increased rate of binding to disease tissue relative to tissues where the CM specific enzyme is not present at a detectable level or is present at a lower level than in disease tissue or is inactive (*e.g.*, in zymogen form or in complex with an inhibitor). Since small proteins and peptides are rapidly cleared from the blood by the renal filtration system, and because the enzyme specific for the CM is not present at a detectable level (or is present at lower levels in non-disease tissues or is present in inactive conformation), accumulation of activated antibodies in the disease tissue is enhanced relative to non-disease tissues.

[000763] In another example, activatable antibodies can be used to detect the presence or absence of a cleaving agent in a sample. For example, where the activatable antibodies contain a CM susceptible to cleavage by an enzyme, the activatable antibodies can be used to detect (either qualitatively or quantitatively) the presence of an enzyme in the sample. In another example, where the activatable antibodies contain a CM susceptible to cleavage by reducing agent, the activatable antibodies can be used to detect (either qualitatively or quantitatively) the presence of reducing conditions in a sample. To facilitate analysis in these methods, the activatable antibodies can be detectably labeled, and can be bound to a support (*e.g.*, a solid support, such as a slide or bead). The detectable label can be positioned on a portion of the activatable antibody that is not released following cleavage, for example, the detectable label can be a quenched fluorescent label or other label that is not detectable until cleavage has occurred. The assay can be conducted by, for example, contacting the immobilized, detectably labeled activatable antibodies with a sample suspected of containing an enzyme and/or reducing agent for a time sufficient for cleavage to occur, then washing to remove excess sample and contaminants. The presence or absence of the cleaving agent (*e.g.*, enzyme or reducing agent) in the sample is then assessed by a change in detectable signal of the activatable antibodies prior to contacting with the sample *e.g.*, the presence of and/or an increase in detectable signal due to cleavage of the activatable antibody by the cleaving agent in the sample.

[000764] Such detection methods can be adapted to also provide for detection of the presence or absence of a target that is capable of binding the AB of the activatable antibodies when cleaved. Thus, the assays can be adapted to assess the presence or absence of a cleaving agent and the presence or absence of a target of interest. The presence or absence of the cleaving agent can be detected by the presence of and/or an increase in detectable label of the activatable antibodies as described above, and the presence or absence of the target can be detected by detection of a target-AB complex *e.g.*, by use of a detectably labeled anti-target antibody.

[000765] Activatable antibodies are also useful in *in situ* imaging for the validation of activatable antibody activation, *e.g.*, by protease cleavage, and binding to a particular target. *In situ* imaging is a technique that enables localization of proteolytic activity and target in biological samples such as cell cultures or tissue sections. Using this technique, it is possible to confirm both binding to a given target and proteolytic activity based on the presence of a detectable label (*e.g.*, a fluorescent label).

[000766] These techniques are useful with any frozen cells or tissue derived from a disease site (*e.g.* tumor tissue) or healthy tissues. These techniques are also useful with fresh cell or tissue samples.

[000767] In these techniques, an activatable antibody is labeled with a detectable label. The detectable label can be a fluorescent dye, (*e.g.* a fluorophore, Fluorescein Isothiocyanate (FITC), Rhodamine Isothiocyanate (TRITC), an Alexa Fluor® label), a near infrared (NIR) dye (*e.g.*, Qdot® nanocrystals), a colloidal metal, a hapten, a radioactive marker, biotin and an amplification reagent such as streptavidin, or an enzyme (*e.g.* horseradish peroxidase or alkaline phosphatase).

[000768] Detection of the label in a sample that has been incubated with the labeled, activatable antibody indicates that the sample contains the target and contains a protease that is specific for the CM of the activatable antibody. In some embodiments, the presence of the protease can be confirmed using broad spectrum protease inhibitors such as those described herein, and/or by using an agent that is specific for the protease, for example, an antibody such as A11, which is specific for the protease matriptase and inhibits the proteolytic activity of matriptase; see *e.g.*, International Publication Number WO 2010/129609, published 11 November 2010. The same approach of using broad spectrum protease inhibitors such as those described herein, and/or by using a more selective inhibitory agent can be used to identify a protease that is specific for the CM of the activatable antibody. In some embodiments, the presence of the target can be confirmed using an agent that is specific for the target, *e.g.*, another antibody, or the detectable label can be competed with unlabeled target. In some embodiments, unlabeled activatable antibody could be used, with detection by a labeled secondary antibody or more complex detection system.

[000769] Similar techniques are also useful for *in vivo* imaging where detection of the fluorescent signal in a subject, *e.g.*, a mammal, including a human, indicates that the disease site contains the target and contains a protease that is specific for the CM of the activatable antibody.

[000770] These techniques are also useful in kits and/or as reagents for the detection, identification or characterization of protease activity in a variety of cells, tissues, and organisms based on the protease-specific CM in the activatable antibody.

[000771] The disclosure provides methods of using the antibodies and/or activatable antibodies in a variety of diagnostic and/or prophylactic indications. For example, the disclosure

provides methods of detecting presence or absence of a cleaving agent and a target of interest in a subject or a sample by (i) contacting a subject or sample with an activatable antibody, wherein the activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, e.g., a protease, and an antigen binding domain or fragment thereof (AB) that specifically binds the target of interest, wherein the activatable antibody in an uncleaved, non-activated state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB and is not a modified form of a natural binding partner of the AB; and (b) wherein, in an uncleaved, non-activated state, the MM interferes with specific binding of the AB to the target, and in a cleaved, activated state the MM does not interfere or compete with specific binding of the AB to the target; and (ii) measuring a level of activated activatable antibody in the subject or sample, wherein a detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent and the target are present in the subject or sample and wherein no detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent, the target or both the cleaving agent and the target are absent and/or not sufficiently present in the subject or sample. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000772] The disclosure also provides methods of detecting presence or absence of a cleaving agent in a subject or a sample by (i) contacting a subject or sample with an activatable antibody in the presence of a target of interest, e.g., the target, wherein the activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, e.g., a protease, and an antigen binding domain or fragment thereof (AB) that specifically binds the target of interest, wherein the activatable antibody in an uncleaved, non-activated state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or

AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB and is not a modified form of a natural binding partner of the AB; and (b) wherein, in an uncleaved, non-activated state, the MM interferes with specific binding of the AB to the target, and in a cleaved, activated state the MM does not interfere or compete with specific binding of the AB to the target; and (ii) measuring a level of activated activatable antibody in the subject or sample, wherein a detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent is present in the subject or sample and wherein no detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or sample. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000773] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent and the target in a subject or a sample, where the kits include at least an activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, e.g., a protease, and an antigen binding domain or fragment thereof (AB) that specifically binds the target of interest, wherein the activatable antibody in an uncleaved, non-activated state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB and is not a modified form of a natural binding partner of the AB; and (b) wherein, in an uncleaved, non-activated state, the MM interferes with specific binding of the AB to the target, and in a cleaved, activated state the MM does not interfere or compete with specific binding of the AB to the target; and (ii) measuring a level of activated activatable antibody in the subject or sample, wherein a detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent is present in the

subject or sample and wherein no detectable level of activated activatable antibody in the subject or sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or sample. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000774] The disclosure also provides methods of detecting presence or absence of a cleaving agent in a subject or a sample by (i) contacting a subject or sample with an activatable antibody, wherein the activatable antibody comprises a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, e.g., a protease, an antigen binding domain (AB) that specifically binds the target, and a detectable label, wherein the activatable antibody in an uncleaved, non-activated state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB and is not a modified form of a natural binding partner of the AB; wherein, in an uncleaved, non-activated state, the MM interferes with specific binding of the AB to the target, and in a cleaved, activated state the MM does not interfere or compete with specific binding of the AB to the target; and wherein the detectable label is positioned on a portion of the activatable antibody that is released following cleavage of the CM; and (ii) measuring a level of detectable label in the subject or sample, wherein a detectable level of the detectable label in the subject or sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or sample and wherein no detectable level of the detectable label in the subject or sample indicates that the cleaving agent is present in the subject or sample. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent

that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000775] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent and the target in a subject or a sample, where the kits include at least an activatable antibody and/or conjugated activatable antibody (e.g., an activatable antibody to which a therapeutic agent is conjugated) described herein for use in contacting a subject or biological sample and means for detecting the level of activated activatable antibody and/or conjugated activatable antibody in the subject or biological sample, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent and the target are present in the subject or biological sample and wherein no detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent, the target or both the cleaving agent and the target are absent and/or not sufficiently present in the subject or biological sample, such that the target binding and/or protease cleavage of the activatable antibody cannot be detected in the subject or biological sample.

[000776] The disclosure also provides methods of detecting presence or absence of a cleaving agent in a subject or a sample by (i) contacting a subject or biological sample with an activatable antibody in the presence of the target, and (ii) measuring a level of activated activatable antibody in the subject or biological sample, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is present in the subject or biological sample and wherein no detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or biological sample at a detectable level, such that protease cleavage of the activatable antibody cannot be detected in the subject or biological sample. Such an activatable antibody includes a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, e.g., a protease, and an antigen binding domain or fragment thereof (AB) that specifically binds the target, wherein the activatable antibody in an uncleaved (*i.e.*, non-activated) state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB; and (b) wherein the MM of the activatable antibody in an

uncleaved state interferes with specific binding of the AB to the target, and wherein the MM of an activatable antibody in a cleaved (*i.e.*, activated) state does not interfere or compete with specific binding of the AB to the target. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the detectable label is attached to the masking moiety. In some embodiments, the detectable label is attached to the cleavable moiety N-terminal to the protease cleavage site. In some embodiments, a single antigen binding site of the AB is masked. In some embodiments wherein an antibody of the disclosure has at least two antigen binding sites, at least one antigen binding site is masked and at least one antigen binding site is not masked. In some embodiments all antigen binding sites are masked. In some embodiments, the measuring step includes use of a secondary reagent comprising a detectable label.

[000777] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent and the target in a subject or a sample, where the kits include at least an activatable antibody and/or conjugated activatable antibody described herein for use in contacting a subject or biological sample with an activatable antibody in the presence of the target, and measuring a level of activated activatable antibody in the subject or biological sample, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is present in the subject or biological sample and wherein no detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or biological sample at a detectable level, such that protease cleavage of the activatable antibody cannot be detected in the subject or biological sample. Such an activatable antibody includes a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, *e.g.*, a protease, and an antigen binding domain or fragment thereof (AB) that specifically binds the target, wherein the activatable antibody in an uncleaved (*i.e.*, non-activated) state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB; and (b) wherein the MM of the activatable antibody in an uncleaved state interferes with specific binding of the AB to the target, and wherein the MM of an activatable antibody in a

cleaved (*i.e.*, activated) state does not interfere or compete with specific binding of the AB to the target. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the detectable label is attached to the masking moiety. In some embodiments, the detectable label is attached to the cleavable moiety N-terminal to the protease cleavage site. In some embodiments, a single antigen binding site of the AB is masked. In some embodiments wherein an antibody of the disclosure has at least two antigen binding sites, at least one antigen binding site is masked and at least one antigen binding site is not masked. In some embodiments all antigen binding sites are masked. In some embodiments, the measuring step includes use of a secondary reagent comprising a detectable label.

[000778] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent in a subject or a sample, where the kits include at least an activatable antibody and/or conjugated activatable antibody described herein for use in contacting a subject or biological sample and means for detecting the level of activated activatable antibody and/or conjugated activatable antibody in the subject or biological sample, wherein the activatable antibody includes a detectable label that is positioned on a portion of the activatable antibody that is released following cleavage of the CM, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or biological sample such that the target binding and/or protease cleavage of the activatable antibody cannot be detected in the subject or biological sample, and wherein no detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent is present in the subject or biological sample at a detectable level.

[000779] The disclosure provides methods of detecting presence or absence of a cleaving agent and the target in a subject or a sample by (i) contacting a subject or biological sample with an activatable antibody, wherein the activatable antibody includes a detectable label that is positioned on a portion of the activatable antibody that is released following cleavage of the CM and (ii) measuring a level of activated activatable antibody in the subject or biological sample, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent, the target or both the cleaving agent and the target are absent and/or not sufficiently present in the subject or biological sample, such that the target binding

and/or protease cleavage of the activatable antibody cannot be detected in the subject or biological sample, and wherein a reduced detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent and the target are present in the subject or biological sample. A reduced level of detectable label is, for example, a reduction of about 5%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95% and/or about 100%. Such an activatable antibody includes a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, and an antigen binding domain or fragment thereof (AB) that specifically binds the target, wherein the activatable antibody in an uncleaved (*i.e.*, non-activated) state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB; and (b) wherein the MM of the activatable antibody in an uncleaved state interferes with specific binding of the AB to the target, and wherein the MM of an activatable antibody in a cleaved (*i.e.*, activated) state does not interfere or compete with specific binding of the AB to the target. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000780] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent and the target in a subject or a sample, where the kits include at least an activatable antibody and/or conjugated activatable antibody described herein for use in contacting a subject or biological sample and means for detecting the level of activated activatable antibody and/or conjugated activatable antibody in the subject or biological sample, wherein a detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent, the target or both the cleaving agent and the target are absent and/or not sufficiently present in the subject or biological sample, such that the target binding

and/or protease cleavage of the activatable antibody cannot be detected in the subject or biological sample, and wherein a reduced detectable level of activated activatable antibody in the subject or biological sample indicates that the cleaving agent and the target are present in the subject or biological sample. A reduced level of detectable label is, for example, a reduction of about 5%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95% and/or about 100%.

[000781] The disclosure also provides methods of detecting presence or absence of a cleaving agent in a subject or a sample by (i) contacting a subject or biological sample with an activatable antibody, wherein the activatable antibody includes a detectable label that is positioned on a portion of the activatable antibody that is released following cleavage of the CM; and (ii) measuring a level of detectable label in the subject or biological sample, wherein a detectable level of the detectable label in the subject or biological sample indicates that the cleaving agent is absent and/or not sufficiently present in the subject or biological sample at a detectable level, such that protease cleavage of the activatable antibody cannot be detected in the subject or biological sample, and wherein a reduced detectable level of the detectable label in the subject or biological sample indicates that the cleaving agent is present in the subject or biological sample. A reduced level of detectable label is, for example, a reduction of about 5%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95% and/or about 100%. Such an activatable antibody includes a masking moiety (MM), a cleavable moiety (CM) that is cleaved by the cleaving agent, and an antigen binding domain or fragment thereof (AB) that specifically binds the target, wherein the activatable antibody in an uncleaved (*i.e.*, non-activated) state comprises a structural arrangement from N-terminus to C-terminus as follows: MM-CM-AB or AB-CM-MM; (a) wherein the MM is a peptide that inhibits binding of the AB to the target, and wherein the MM does not have an amino acid sequence of a naturally occurring binding partner of the AB; and (b) wherein the MM of the activatable antibody in an uncleaved state interferes with specific binding of the AB to the target, and wherein the MM of an activatable antibody in a cleaved (*i.e.*, activated) state does not interfere or compete with specific binding of the AB to the target. In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic

agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000782] The disclosure also provides kits for use in methods of detecting presence or absence of a cleaving agent of interest in a subject or a sample, where the kits include at least an activatable antibody and/or conjugated activatable antibody described herein for use in contacting a subject or biological sample and means for detecting the level of activated activatable antibody and/or conjugated activatable antibody in the subject or biological sample, wherein the activatable antibody includes a detectable label that is positioned on a portion of the activatable antibody that is released following cleavage of the CM, wherein a detectable level of the detectable label in the subject or biological sample indicates that the cleaving agent, the target, or both the cleaving agent and the target are absent and/or not sufficiently present in the subject or biological sample, such that the target binding and/or protease cleavage of the activatable antibody cannot be detected in the subject or biological sample, and wherein a reduced detectable level of the detectable label in the subject or biological sample indicates that the cleaving agent and the target are present in the subject or biological sample. A reduced level of detectable label is, for example, a reduction of about 5%, about 10%, about 15%, about 20%, about 25%, about 30%, about 35%, about 40%, about 45%, about 50%, about 55%, about 60%, about 65%, about 70%, about 75%, about 80%, about 85%, about 90%, about 95% and/or about 100%.

[000783] In some embodiments of these methods and kits, the activatable antibody includes a detectable label. In some embodiments of these methods and kits, the detectable label includes an imaging agent, a contrasting agent, an enzyme, a fluorescent label, a chromophore, a dye, one or more metal ions, or a ligand-based label. In some embodiments of these methods and kits, the imaging agent comprises a radioisotope. In some embodiments of these methods and kits, the radioisotope is indium or technetium. In some embodiments of these methods and kits, the contrasting agent comprises iodine, gadolinium or iron oxide. In some embodiments of these methods and kits, the enzyme comprises horseradish peroxidase, alkaline phosphatase, or β -

galactosidase. In some embodiments of these methods and kits, the fluorescent label comprises yellow fluorescent protein (YFP), cyan fluorescent protein (CFP), green fluorescent protein (GFP), modified red fluorescent protein (mRFP), red fluorescent protein tdimer2 (RFP tdimer2), HCRED, or a europium derivative. In some embodiments of these methods and kits, the luminescent label comprises an N- methylacrydium derivative. In some embodiments of these methods, the label comprises an Alexa Fluor[®] label, such as Alex Fluor[®] 680 or Alexa Fluor[®] 750. In some embodiments of these methods and kits, the ligand-based label comprises biotin, avidin, streptavidin or one or more haptens.

[000784] In some embodiments of these methods and kits, the subject is a mammal. In some embodiments of these methods and kits, the subject is a human. In some embodiments, the subject is a non-human mammal, such as a non-human primate, companion animal (e.g., cat, dog, horse), farm animal, work animal, or zoo animal. In some embodiments, the subject is a rodent.

[000785] In some embodiments of these methods, the method is an *in vivo* method. In some embodiments of these methods, the method is an *in situ* method. In some embodiments of these methods, the method is an *ex vivo* method. In some embodiments of these methods, the method is an *in vitro* method.

[000786] In some embodiments, *in situ* imaging and/or *in vivo* imaging are useful in methods to identify which patients to treat. For example, in *in situ* imaging, the activatable antibodies are used to screen patient samples to identify those patients having the appropriate protease(s) and target(s) at the appropriate location, e.g., at a tumor site.

[000787] In some embodiments *in situ* imaging is used to identify or otherwise refine a patient population suitable for treatment with an activatable antibody of the disclosure. For example, patients that test positive for both the target (e.g., the target) and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody being tested (e.g., accumulate activated antibodies at the disease site) are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. Likewise, patients that test negative for either or both of the target (e.g., the target) and the protease that cleaves the substrate in the CM in the activatable antibody being tested using these methods might be identified as suitable candidates for another form of therapy. In some embodiments, such patients that test negative with respect to a first activatable antibody can be tested with other activatable

antibodies comprising different CMs until a suitable activatable antibody for treatment is identified (e.g., an activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, the patient is then administered a therapeutically effective amount of the activatable antibody for which the patient tested positive.

[000788] In some embodiments *in vivo* imaging is used to identify or otherwise refine a patient population suitable for treatment with an activatable antibody of the disclosure. For example, patients that test positive for both the target (e.g., the target) and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody being tested (e.g., accumulate activated antibodies at the disease site) are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. Likewise, patients that test negative might be identified as suitable candidates for another form of therapy. In some embodiments, such patients that test negative with respect to a first activatable antibody can be tested with other activatable antibodies comprising different CMs until a suitable activatable antibody for treatment is identified (e.g., an activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, the patient is then administered a therapeutically effective amount of the activatable antibody for which the patient tested positive.

[000789] In some embodiments of the methods and kits, the method or kit is used to identify or otherwise refine a patient population suitable for treatment with an activatable antibody of the disclosure. For example, patients that test positive for both the target (e.g., the target) and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody being tested in these methods are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. Likewise, patients that test negative for both of the targets (e.g., the target) and the protease that cleaves the substrate in the CM in the activatable antibody being tested using these methods might be identified as suitable candidates for another form of therapy. In some embodiments, such patients can be tested with other activatable antibodies until a suitable activatable antibody for treatment is identified (e.g., an activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, patients that test negative for either of the target (e.g., the target) are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. In some embodiments, patients that test negative for either of the target (e.g., the target) are identified as not being suitable candidates for treatment with such an activatable antibody

comprising such a CM. In some embodiments, such patients can be tested with other activatable antibodies until a suitable activatable antibody for treatment is identified (e.g., an activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, the activatable antibody is an activatable antibody to which a therapeutic agent is conjugated. In some embodiments, the activatable antibody is not conjugated to an agent. In some embodiments, the activatable antibody comprises a detectable label. In some embodiments, the detectable label is positioned on the AB. In some embodiments, measuring the level of activatable antibody in the subject or sample is accomplished using a secondary reagent that specifically binds to the activated antibody, wherein the reagent comprises a detectable label. In some embodiments, the secondary reagent is an antibody comprising a detectable label.

[000790] In some embodiments, a method or kit is used to identify or otherwise refine a patient population suitable for treatment with an anti-the target activatable antibody and/or conjugated activatable antibody (e.g., activatable antibody to which a therapeutic agent is conjugated) of the disclosure, followed by treatment by administering that activatable antibody and/or conjugated activatable antibody to a subject in need thereof. For example, patients that test positive for both the targets (e.g., the target) and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody and/or conjugated activatable antibody being tested in these methods are identified as suitable candidates for treatment with such antibody and/or such a conjugated activatable antibody comprising such a CM, and the patient is then administered a therapeutically effective amount of the activatable antibody and/or conjugated activatable antibody that was tested. Likewise, patients that test negative for either or both of the target (e.g., the target) and the protease that cleaves the substrate in the CM in the activatable antibody being tested using these methods might be identified as suitable candidates for another form of therapy. In some embodiments, such patients can be tested with other antibody and/or conjugated activatable antibody until a suitable antibody and/or conjugated activatable antibody for treatment is identified (e.g., an activatable antibody and/or conjugated activatable antibody comprising a CM that is cleaved by the patient at the site of disease). In some embodiments, the patient is then administered a therapeutically effective amount of the activatable antibody and/or conjugated activatable antibody for which the patient tested positive.

[000791] In some embodiments of these methods and kits, the MM is a peptide having a length from about 4 to 40 amino acids. In some embodiments of these methods and kits, the

activatable antibody comprises a linker peptide, wherein the linker peptide is positioned between the MM and the CM. In some embodiments of these methods and kits, the activatable antibody comprises a linker peptide, where the linker peptide is positioned between the AB and the CM. In some embodiments of these methods and kits, the activatable antibody comprises a first linker peptide (L1) and a second linker peptide (L2), wherein the first linker peptide is positioned between the MM and the CM and the second linker peptide is positioned between the AB and the CM. In some embodiments of these methods and kits, each of L1 and L2 is a peptide of about 1 to 20 amino acids in length, and wherein each of L1 and L2 need not be the same linker. In some embodiments of these methods and kits, one or both of L1 and L2 comprises a glycine-serine polymer. In some embodiments of these methods and kits, at least one of L1 and L2 comprises an amino acid sequence selected from the group consisting of (GS)_n, (GSGGS)_n (SEQ ID NO: 363) and (GGGS)_n (SEQ ID NO: 364), where n is an integer of at least one. In some embodiments of these methods and kits, at least one of L1 and L2 comprises an amino acid sequence having the formula (GGS)_n, where n is an integer of at least one. In some embodiments of these methods and kits, at least one of L1 and L2 comprises an amino acid sequence selected from the group consisting of GGSG (SEQ ID NO: 365), GGS GG (SEQ ID NO: 366), GSGSG (SEQ ID NO: 367), GSGGG (SEQ ID NO: 368), GGGSG (SEQ ID NO: 369), and GSSSG (SEQ ID NO: 370).

[000792] In some embodiments of these methods and kits, the AB comprises an antibody or antibody fragment sequence selected from the cross-reactive antibody sequences presented herein. In some embodiments of these methods and kits, the AB comprises a Fab fragment, a scFv or a single chain antibody (scAb).

[000793] In some embodiments of these methods and kits, the cleaving agent is a protease that is co-localized in the subject or sample with the target and the CM is a polypeptide that functions as a substrate for the protease, wherein the protease cleaves the CM in the activatable antibody when the activatable antibody is exposed to the protease. In some embodiments of these methods and kits, the CM is a polypeptide of up to 15 amino acids in length. In some embodiments of these methods and kits, the CM is coupled to the N-terminus of the AB. In some embodiments of these methods and kits, the CM is coupled to the C-terminus of the AB. In some embodiments of these methods and kits, the CM is coupled to the N-terminus of a VL chain of the AB.

[000794] The antibodies and/or activatable antibodies of the disclosure are used in diagnostic and prophylactic formulations. In one embodiment, an activatable antibody is administered to patients that are at risk of developing one or more of the aforementioned inflammation, inflammatory disorders, cancer or other disorders.

[000795] A patient's or organ's predisposition to one or more of the aforementioned disorders can be determined using genotypic, serological or biochemical markers.

[000796] In some embodiments of the disclosure, an antibody and/or an activatable antibody is administered to human individuals diagnosed with a clinical indication associated with one or more of the aforementioned disorders. Upon diagnosis, an antibody and/or an activatable antibody is administered to mitigate or reverse the effects of the clinical indication.

[000797] Antibodies and/or activatable antibodies of the disclosure are also useful in the detection of the target in patient samples and accordingly are useful as diagnostics. For example, the antibodies and/or activatable antibodies of the disclosure are used in *in vitro* assays, *e.g.*, ELISA, to detect target levels in a patient sample.

[000798] In some embodiments, anti-PDL1 antibodies are used as a diagnostic for patients who are more likely to respond favorably to treatment with an anti-PD-1 antibody and/or activatable antibody of the disclosure. In these embodiments, PD-L1 expression in tumor biopsies on both tumor cells and infiltrating immune cells and PD-L1 expression on immune cells in blood are used to indicate the presence of active immunity and, thus, potential for suppression of T cell activity through the engagement of PD-1 by PD-L1.

[000799] In one embodiment, an antibody and/or activatable antibody of the disclosure is immobilized on a solid support (*e.g.*, the well(s) of a microtiter plate). The immobilized antibody and/or activatable antibody serves as a capture antibody for any target that can be present in a test sample. Prior to contacting the immobilized antibody and/or activatable antibody with a patient sample, the solid support is rinsed and treated with a blocking agent such as milk protein or albumin to prevent nonspecific adsorption of the analyte.

[000800] Subsequently the wells are treated with a test sample suspected of containing the antigen, or with a solution containing a standard amount of the antigen. Such a sample is, *e.g.*, a serum sample from a subject suspected of having levels of circulating antigen considered to be diagnostic of a pathology. After rinsing away the test sample or standard, the solid support is treated with a second antibody that is detectably labeled. The labeled second antibody serves as a

detecting antibody. The level of detectable label is measured, and the concentration of target antigen in the test sample is determined by comparison with a standard curve developed from the standard samples.

[000801] It will be appreciated that based on the results obtained using the antibodies and/or activatable antibodies of the disclosure in an *in vitro* diagnostic assay, it is possible to stage a disease in a subject based on expression levels of the target antigen. For a given disease, samples of blood are taken from subjects diagnosed as being at various stages in the progression of the disease, and/or at various points in the therapeutic treatment of the disease. Using a population of samples that provides statistically significant results for each stage of progression or therapy, a range of concentrations of the antigen that can be considered characteristic of each stage is designated.

[000802] Antibodies, conjugated antibodies, activatable antibodies and/or conjugated activatable antibodies can also be used in diagnostic and/or imaging methods. In some embodiments, such methods are *in vitro* methods. In some embodiments, such methods are *in vivo* methods. In some embodiments, such methods are *in situ* methods. In some embodiments, such methods are *ex vivo* methods. For example, activatable antibodies having an enzymatically cleavable CM can be used to detect the presence or absence of an enzyme that is capable of cleaving the CM. Such activatable antibodies can be used in diagnostics, which can include *in vivo* detection (*e.g.*, qualitative or quantitative) of enzyme activity (or, in some embodiments, an environment of increased reduction potential such as that which can provide for reduction of a disulfide bond) through measured accumulation of activated antibodies (*i.e.*, antibodies resulting from cleavage of an activatable antibody) in a given cell or tissue of a given host organism. Such accumulation of activated antibodies indicates not only that the tissue expresses enzymatic activity (or an increased reduction potential depending on the nature of the CM) but also that the tissue expresses target to which the activated antibody binds.

[000803] For example, the CM can be selected to be a protease substrate for a protease found at the site of a tumor, at the site of a viral or bacterial infection at a biologically confined site (*e.g.*, such as in an abscess, in an organ, and the like), and the like. The AB can be one that binds a target antigen. Using methods familiar to one skilled in the art, a detectable label (*e.g.*, a fluorescent label or radioactive label or radiotracer) can be conjugated to an AB or other region of an activatable antibody. Suitable detectable labels are discussed in the context of the above

screening methods and additional specific examples are provided below. Using an AB specific to a protein or peptide of the disease state, along with a protease whose activity is elevated in the disease tissue of interest, activatable antibodies will exhibit an increased rate of binding to disease tissue relative to tissues where the CM specific enzyme is not present at a detectable level or is present at a lower level than in disease tissue or is inactive (e.g., in zymogen form or in complex with an inhibitor). Since small proteins and peptides are rapidly cleared from the blood by the renal filtration system, and because the enzyme specific for the CM is not present at a detectable level (or is present at lower levels in non-disease tissues or is present in inactive conformation), accumulation of activated antibodies in the disease tissue is enhanced relative to non-disease tissues.

[000804] In another example, activatable antibodies can be used to detect the presence or absence of a cleaving agent in a sample. For example, where the activatable antibodies contain a CM susceptible to cleavage by an enzyme, the activatable antibodies can be used to detect (either qualitatively or quantitatively) the presence of an enzyme in the sample. In another example, where the activatable antibodies contain a CM susceptible to cleavage by reducing agent, the activatable antibodies can be used to detect (either qualitatively or quantitatively) the presence of reducing conditions in a sample. To facilitate analysis in these methods, the activatable antibodies can be detectably labeled, and can be bound to a support (e.g., a solid support, such as a slide or bead). The detectable label can be positioned on a portion of the activatable antibody that is not released following cleavage, for example, the detectable label can be a quenched fluorescent label or other label that is not detectable until cleavage has occurred. The assay can be conducted by, for example, contacting the immobilized, detectably labeled activatable antibodies with a sample suspected of containing an enzyme and/or reducing agent for a time sufficient for cleavage to occur, then washing to remove excess sample and contaminants. The presence or absence of the cleaving agent (e.g., enzyme or reducing agent) in the sample is then assessed by a change in detectable signal of the activatable antibodies prior to contacting with the sample e.g., the presence of and/or an increase in detectable signal due to cleavage of the activatable antibody by the cleaving agent in the sample.

[000805] Such detection methods can be adapted to also provide for detection of the presence or absence of a target that is capable of binding the AB of the activatable antibodies when cleaved. Thus, the assays can be adapted to assess the presence or absence of a cleaving

agent and the presence or absence of a target of interest. The presence or absence of the cleaving agent can be detected by the presence of and/or an increase in detectable label of the activatable antibodies as described above, and the presence or absence of the target can be detected by detection of a target-AB complex *e.g.*, by use of a detectably labeled anti-target antibody.

[000806] Activatable antibodies are also useful in *in situ* imaging for the validation of activatable antibody activation, *e.g.*, by protease cleavage, and binding to a particular target. *In situ* imaging is a technique that enables localization of proteolytic activity and target in biological samples such as cell cultures or tissue sections. Using this technique, it is possible to confirm both binding to a given target and proteolytic activity based on the presence of a detectable label (*e.g.*, a fluorescent label).

[000807] These techniques are useful with any frozen cells or tissue derived from a disease site (*e.g.* tumor tissue) or healthy tissues. These techniques are also useful with fresh cell or tissue samples.

[000808] In these techniques, an activatable antibody is labeled with a detectable label. The detectable label can be a fluorescent dye, (*e.g.* Fluorescein Isothiocyanate (FITC), Rhodamine Isothiocyanate (TRITC), a near infrared (NIR) dye (*e.g.*, Qdot® nanocrystals), a colloidal metal, a hapten, a radioactive marker, biotin and an amplification reagent such as streptavidin, or an enzyme (*e.g.* horseradish peroxidase or alkaline phosphatase).

[000809] Detection of the label in a sample that has been incubated with the labeled, activatable antibody indicates that the sample contains the target and contains a protease that is specific for the CM of the activatable antibody. In some embodiments, the presence of the protease can be confirmed using broad spectrum protease inhibitors such as those described herein, and/or by using an agent that is specific for the protease, for example, an antibody such as A11, which is specific for the protease matriptase and inhibits the proteolytic activity of matriptase; see *e.g.*, International Publication Number WO 2010/129609, published 11 November 2010. The same approach of using broad spectrum protease inhibitors such as those described herein, and/or by using a more selective inhibitory agent can be used to identify a protease or class of proteases specific for the CM of the activatable antibody. In some embodiments, the presence of the target can be confirmed using an agent that is specific for the target, *e.g.*, another antibody, or the detectable label can be competed with unlabeled target. In

some embodiments, unlabeled activatable antibody could be used, with detection by a labeled secondary antibody or more complex detection system.

[000810] Similar techniques are also useful for *in vivo* imaging where detection of the fluorescent signal in a subject, *e.g.*, a mammal, including a human, indicates that the disease site contains the target and contains a protease that is specific for the CM of the activatable antibody.

[000811] These techniques are also useful in kits and/or as reagents for the detection, identification or characterization of protease activity in a variety of cells, tissues, and organisms based on the protease-specific CM in the activatable antibody.

[000812] In some embodiments, *in situ* imaging and/or *in vivo* imaging are useful in methods to identify which patients to treat. For example, in *in situ* imaging, the activatable antibodies are used to screen patient samples to identify those patients having the appropriate protease(s) and target(s) at the appropriate location, *e.g.*, at a tumor site.

[000813] In some embodiments *in situ* imaging is used to identify or otherwise refine a patient population suitable for treatment with an activatable antibody of the disclosure. For example, patients that test positive for both the target and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody being tested (*e.g.*, accumulate activated antibodies at the disease site) are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. Likewise, patients that test negative for either or both of the target and the protease that cleaves the substrate in the CM in the activatable antibody being tested using these methods are identified as suitable candidates for another form of therapy (*i.e.*, not suitable for treatment with the activatable antibody being tested). In some embodiments, such patients that test negative with respect to a first activatable antibody can be tested with other activatable antibodies comprising different CMs until a suitable activatable antibody for treatment is identified (*e.g.*, an activatable antibody comprising a CM that is cleaved by the patient at the site of disease).

[000814] In some embodiments *in vivo* imaging is used to identify or otherwise refine a patient population suitable for treatment with an activatable antibody of the disclosure. For example, patients that test positive for both the target and a protease that cleaves the substrate in the cleavable moiety (CM) of the activatable antibody being tested (*e.g.*, accumulate activated antibodies at the disease site) are identified as suitable candidates for treatment with such an activatable antibody comprising such a CM. Likewise, patients that test negative are identified as

suitable candidates for another form of therapy (*i.e.*, not suitable for treatment with the activatable antibody being tested). In some embodiments, such patients that test negative with respect to a first activatable antibody can be tested with other activatable antibodies comprising different CMs until a suitable activatable antibody for treatment is identified (e.g., an activatable antibody comprising a CM that is cleaved by the patient at the site of disease).

Pharmaceutical compositions

[000815] The antibodies and/or activatable antibodies of the disclosure (also referred to herein as “active compounds”), and derivatives, fragments, analogs and homologs thereof, can be incorporated into pharmaceutical compositions suitable for administration. Such compositions typically comprise the antibody and/or activatable antibody and a pharmaceutically acceptable carrier. As used herein, the term “pharmaceutically acceptable carrier” is intended to include any and all solvents, dispersion media, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like, compatible with pharmaceutical administration. Suitable carriers are described in the most recent edition of Remington’s Pharmaceutical Sciences, a standard reference text in the field, which is incorporated herein by reference. Suitable examples of such carriers or diluents include, but are not limited to, water, saline, ringer’s solutions, dextrose solution, and 5% human serum albumin. Liposomes and non-aqueous vehicles such as fixed oils can also be used. The use of such media and agents for pharmaceutically active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active compound, use thereof in the compositions is contemplated. Supplementary active compounds can also be incorporated into the compositions.

[000816] A pharmaceutical composition of the disclosure is formulated to be compatible with its intended route of administration. Examples of routes of administration include parenteral, *e.g.*, intravenous, intradermal, subcutaneous, oral (*e.g.*, inhalation), transdermal (*i.e.*, topical), transmucosal, and rectal administration. Solutions or suspensions used for parenteral, intradermal, or subcutaneous application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid (EDTA); buffers such as acetates, citrates or phosphates, and agents for the adjustment of tonicity such as sodium chloride or dextrose. The pH can be

adjusted with acids or bases, such as hydrochloric acid or sodium hydroxide. The parenteral preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

[000817] Pharmaceutical compositions suitable for injectable use include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersion. For intravenous administration, suitable carriers include physiological saline, bacteriostatic water, Cremophor EL™ (BASF, Parsippany, N.J.) or phosphate buffered saline (PBS). In all cases, the composition must be sterile and should be fluid to the extent that easy syringeability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), and suitable mixtures thereof. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of surfactants. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal, and the like. In some embodiments, it will be desirable to include isotonic agents, for example, sugars, polyalcohols such as manitol, sorbitol, sodium chloride in the composition. Prolonged absorption of the injectable compositions can be brought about by including in the composition an agent that delays absorption, for example, aluminum monostearate and gelatin.

[000818] Sterile injectable solutions can be prepared by incorporating the active compound in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the active compound into a sterile vehicle that contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, methods of preparation are vacuum drying and freeze-drying that yields a powder of the active ingredient plus any additional desired ingredient from a previously sterile-filtered solution thereof.

[000819] Oral compositions generally include an inert diluent or an edible carrier. They can be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic

administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules. Oral compositions can also be prepared using a fluid carrier for use as a mouthwash, wherein the compound in the fluid carrier is applied orally and swished and expectorated or swallowed. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition. The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring.

[000820] For administration by inhalation, the compounds are delivered in the form of an aerosol spray from pressured container or dispenser that contains a suitable propellant, *e.g.*, a gas such as carbon dioxide, or a nebulizer.

[000821] Systemic administration can also be by transmucosal or transdermal means. For transmucosal or transdermal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art, and include, for example, for transmucosal administration, detergents, bile salts, and fusidic acid derivatives. Transmucosal administration can be accomplished through the use of nasal sprays or suppositories. For transdermal administration, the active compounds are formulated into ointments, salves, gels, or creams as generally known in the art.

[000822] The compounds can also be prepared in the form of suppositories (*e.g.*, with conventional suppository bases such as cocoa butter and other glycerides) or retention enemas for rectal delivery.

[000823] In one embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters, and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation and Nova Pharmaceuticals, Inc. Liposomal suspensions

(including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) can also be used as pharmaceutically acceptable carriers. These can be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811.

[000824] It is especially advantageous to formulate oral or parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used herein refers to physically discrete units suited as unitary dosages for the subject to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specification for the dosage unit forms of the disclosure are dictated by and directly dependent on the unique characteristics of the active compound and the particular therapeutic effect to be achieved, and the limitations inherent in the art of compounding such an active compound for the treatment of individuals.

[000825] The pharmaceutical compositions can be included in a container, pack, or dispenser together with instructions for administration.

[000826] The invention will be further described in the following examples, which do not limit the scope of the invention described in the claims.

Examples

EXAMPLE 1. Generation of mouse antibodies of the embodiments that bind human PD-1 and block hPD-L1 and hPD-L2 binding to human PD-1

[000827] This example demonstrates mouse antibodies of the disclosure that bind human PD-1 can be isolated from hybridomas derived from mouse immunization with recombinant human PD-1 protein, and that such binding can inhibit PD-1 binding to PDL1 and PDL2.

[000828] Six NZBWF1/J female mice (Jackson Laboratories, Sacramento, CA; cat# 100008) were immunized in the right flank with recombinant human PD-1 (Sino Biological, Beijing, P.R. China; cat# ABIN2181605) on days 0, 7 and 21. Serum was taken from immunized mice on day 28 and binding to HEK293-hPD-1 (cells transfected with an expression vector encoding human PD-1 (Origene, cat# SC117011)) was measured. All six mice showed positive binding. Splenocytes were isolated from mouse 1, 3 and 6 and fused with SP0 mouse B-cells; likewise splenocytes were isolated from mouse 2, 4 and 5 and fused with SP0 mouse B-cells, giving rise to pools of hybridomas, m136 and m245. The hybridoma pools, m136 and m245, arising from the two fusions were plated for single clones, and clonal culture supernatants were assayed for antibodies capable of binding to HEK293-hPD-1, and not to un-transfected HEK293, cells. Hybridoma clones expressing anti-PD-1 antibodies were chosen for further analysis.

[000829] The sequences of the antibodies used in the studies presented herein are shown below:

m136-M13 – MHC723 mIgG1/K

MHC723HC.1 Variable heavy chain region amino acid sequence:

EVKLVESGGGLVKPGGSLKLSCAASGFTFSGYAMSWVRQTPAKRLEWVAYISNSGGNAHYPDSV
KGRFTISRDNKNTLYLQMSSLRSEDTAMYYCTREDYGTSPFVYWGQGLVTVSA (SEQ ID
NO: 1)

MHC723HC.1 Variable heavy chain region nucleic acid sequence:

GAAGTGAAGCTGGTGGAGTCTGGGGGAGGCTTAGTGAAACCTGGAGGGTCCCTGAAACTCTCCT
GTGCAGCCTCTGGATTCACCTTTCAGTGGCTATGCCATGTCTTGGGTTCGCCAGACTCCGGCGAA
GAGGCTGGAGTGGGTTCGCATACATTAGTAATAGTGGTGGTAACGCCCACTATCCAGACAGTGTA

AAGGGCCGATTACCATCTCCAGAGACAATGCCAAGAACACCCTATACCTGCAAATGAGCAGTC
TGAGGTCTGAGGACACGGCCATGTATTACTGTACAAGAGAGGACTACGGTACTAGTCCTTTTGT
TTACTGGGGCCAAGGGACTCTGGTCACTGTCTCTGCA (SEQ ID NO: 2)

MHC723LC.3 Variable light chain region amino acid sequence:

DIVLTQSPASLAVSLGQRTTISCRASESVDNYGISFMNWFQQKPGQPPKLLIYAASNQSGVPA
RFSGSGSGTDFSLNIHPMEEDDTAVYFCQOSKDVPWTFGGGKLEIR (SEQ ID NO: 3)

MHC723LC.3 Variable light chain region nucleic acid sequence:

GACATTGTGCTGACCCAATCTCCAGCTTCTTTGGCTGTGTCTCTTGGGCAGAGGACCACCATCT
CCTGCAGAGCCAGCGAAAGTGTTGATAATTATGGCATTAGTTTTATGAACTGGTTCCAACAGAA
ACCAGGACAGCCACCCAACTCCTCATCTATGCTGCATCCAACCAAGGATCCGGGGTCCCTGCC
AGGTTTAGTGGCAGTGGGTCTGGGACAGACTTCAGCCTCAACATCCATCCTATGGAGGAGGATG
ATACTGCAGTGTATTTCTGTCAGCAAAGTAAGGACGTTCCGTGGACGTTCCGTGGAGGCACCAA
GCTGGAAATCAGAC (SEQ ID NO: 4)

m136-M19 – MHC725 mIgG2b/K

MHC725HC.2 Variable heavy chain region amino acid sequence:

EVQLQQSGPELVKPGDSVKMSCKASGYTFDYYMDWVKQSHGKSLEWIGYIYPKNGGSSYNQKF
KGKATLTVDKSSSTAYMELHSLTSEDSAVYYCARKVVATDYWGQGTTTLTVSS (SEQ ID
NO: 5)

MHC725HC.2 Variable heavy chain region nucleic acid sequence:

GAGGTCCAGCTGCAACAGTCTGGACCTGAGCTGGTGAAGCCTGGGGATTCACTGAAGATGTCCT
GCAAGGCTTCTGGCTACACATTCAGTACTACTACATGGACTGGGTGAAGCAGAGCCATGGAAA
GAGCCTTGAGTGGATTGGATATATTTATCCTAAAAATGGTGGTTCCAGCTACAATCAGAAGTTC
AAGGGCAAGGCCACATTGACTGTAGACAAGTCCTCCAGCACAGCCTACATGGAGCTCCACAGCC
TGACATCTGAGGACTCTGCAGTCTATTACTGTGCAAGAAAGGTCGTAGCTACGGACTACTGGGG
CCAAGGCACCACTCTCACAGTCTCCTCA (SEQ ID NO: 6)

MHC725LC.2 Variable light chain region amino acid sequence:

DIVMSQSPSSLAVSVGEKVTMSCKSSQSLLYSSNQKNYLAWYQQKPGQSPKLLIFWASIRESGV
PDRFTGSGSGTDFTLTISVKAEDRAVYYCQQCDSYPWTFGGGTKLEIK (SEQ ID NO: 7)

MHC725LC.2 Variable light chain region nucleic acid sequence:

GACATTGTGATGTCACAGTCTCCATCCTCCCTAGCTGTGTCAGTTGGAGAGAAGGTTACTATGA
GCTGCAAGTCCAGTCAGAGCCTTTTATATAGTAGCAATCAAAAGAACTACTTGGCCTGGTACCA
GCAGAAACCAGGGCAGTCTCCTAAACTGCTGATTTTCTGGGCATCTATTAGGGAATCTGGGGTC
CCTGATCGCTTCACAGGCAGTGGATCTGGGACAGATTTCACTCTCACCATCAGCAGTGTGAAGG
CTGAAGACCGGGCAGTTTATTACTGTCAGCAATGTGATAGCTATCCGTGGACGTTTCGGTGGAGG
CACCAAACCTGGAAATCAAAC (SEQ ID NO: 8)

m245-M3 – MHC728 mIgG2a/K

MHC728HC.4 Variable heavy chain region amino acid sequence:

EVKLVESGGGLVKPGGSLKLSCAASGFTFSNYAMSWVRQTPAKRLEWVAYISNGGGDTHYPDSL
KGRFTVSRDNAKNTLYLQMSSLKSEDTAMYYCARENYGTSPFVYWGQGLTVTSA (SEQ ID
NO: 9)

MHC728HC.4 Variable heavy chain region nucleic acid sequence:

GAAGTGAAGCTGGTGGAGTCTGGGGGAGGCTTAGTGAAACCTGGAGGGTCCCTGAAACTCTCCT
GTGCAGCCTCTGGATTCACTTTTCTAGTAATGCGCATGTCTTGGGTTCGCCAGACTCCGGCGAA
GAGGCTGGAGTGGGTTCGCATACATTAGTAATGGTGGTGGTGACACCCACTATCCAGACAGTTTA
AAGGGCCGATTACCGTCTCCAGAGACAATGCCAAGAACACCCTGTACCTACAAATGAGCAGTC
TGAAGTCTGAGGACACGGCCATGTATTACTGTGCAAGAGAAAACCTACGGTACTAGTCCCTTTGT
TTACTGGGGCCAAGGGACTCTGGTCACTGTCTCTGCA (SEQ ID NO: 10)

MHC728LC.2 Variable light chain region amino acid sequence:

DIVLTQSPASLAVSLGQRATISCRASESVDNYGISFMNWFQQKPGQPPKLLIYAASNQSGVPA
RFSGSGSGTDFTSLNIHPMEEDDTAMYFCQQSKDVPWTFGGGTKLEIK (SEQ ID NO: 11)

MHC728LC.2 Variable light chain region nucleic acid sequence:

GACATTGTGCTGACCCAATCTCCAGCTTCTTTGGCTGTGTCTCTAGGGCAGAGGGCCACCATCT
 CCTGCAGAGCCAGCGAAAGTGTTGATAATTATGGCATTAGTTTTATGAACTGGTTCCAACAGAA
 ACCAGGACAGCCACCCAAACTCCTCATCTATGCTGCATCCAACCAAGGATCCGGGGTCCCTGCC
 AGGTTTAGTGGCAGTGGGTCTGGGACAGACTTCAGCCTCAACATCCATCCTATGGAGGAGGATG
 ATACTGCAATGTATTTCTGTCAGCAAAGTAAAGATGTTCCGTGGACGTTCCGGTGGAGGCACCAA
 GCTGGAAATCAAAC (SEQ ID NO: 12)

m245-M5 – MHC729 mIgG1/K**MHC729HC.1 Variable heavy chain region amino acid sequence:**

EVQLVESGGGLVKSGGSLKLSCAHSGFSFSSYDMSWVRQTPAKRLEWVATISGGGRYTYYPDSV
 KGRFTISRDNKNTLYLQMSGLRSEDTAMYYCASNYYGFDYWGQGTTLTVSS (SEQ ID
 NO: 13)

MHC729HC.1 Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGGAGTCTGGGGGAGGCTTAGTGAAGTCTGGAGGGTCCCTGAAACTCTCCT
 GTGCGCATTCTGGATTCTAGTTTTAGTAGTTATGACATGTCTTGGGTTCGCCAGACTCCGGCGAA
 GAGGCTGGAGTGGGTGCGAACCATTAGTGGTGGTGGTCGTTACACCTACTATCCAGACAGTGTG
 AAGGGCCGATTACCATCTCCAGAGACAATGCCAAGAACACCCTGTACCTGCAAATGAGCGGTC
 TGAGGTCTGAGGACACAGCCATGTATTACTGTGCAAGTAATTACTACGGTTTTGACTACTGGGG
 CCAAGGCACCACTCTCACAGTCTCTTCA (SEQ ID NO: 14)

MHC729LC.3 Variable light chain region amino acid sequence:

DIVMTQSHKFMSTSVGDRVSI TCKASQDVGTAVAWYQQKPGQSPKLLIYWASTRHTGVDPDRFTG
 SGSGTDFTLTISNVQSEDLADYFCQQYSSYPWTFGGGTKLEIK (SEQ ID NO: 15)

MHC729LC.3 Variable light chain region nucleic acid sequence:

GATATTGTGATGACCCAGTCTCACAAATTCATGTCCACATCAGTAGGAGACAGGGTCAGCATCA
 CCTGCAAGGCCAGTCAGGATGTGGGTACTGCTGTAGCCTGGTATCAACAGAAACCAGGGCAATC
 TCCTAAACTACTGATTTACTGGGCATCCACCCGGCACACTGGAGTCCCTGATCGCTTCACGGGC
 AGTGGATCTGGGACAGATTTCACTCTCACCATTAGCAATGTGCAGTCTGAAGACTTGGCAGATT

ATTTCTGTCAGCAATATAGCAGCTATCCGTGGACGTTTCGGTGGAGGCACCAAGCTGGAAATCAA
AC (SEQ ID NO: 16)

m136-M14 – MHC724 mIgG2a/K

MHC724HC.3 Variable heavy chain region amino acid sequence:

KVMLVESGGDLVKPGGSLKLSCAASSGFTFSSYGMSWVRQTPEKRLEWVATISGGGRDIYYADTV
KGRFTISRDNAKNTLYLQMSLRSEDTALYFCARLYLGF~~FDY~~WGQGTTLTVSS (SEQ ID
NO: 17)

MHC724HC.3 Variable heavy chain region nucleic acid sequence:

AAAGTGATGCTGGTGGAGTCTGGGGGAGACTTAGTGAAGCCTGGAGGGTCCCTGAAACTCTCCT
GTGCAGCCTCTGGATTCACTTTCAGTAGCTATGGCATGTCTTGGGTTGCCAGACTCCGGAGAA
GAGGCTGGAGTGGGTGCGAACCATTAGTGGTGGTGGTAGAGACATCTACTACGCAGACACTGTG
AAGGGCCGATTCACCATCTCCAGAGACAATGCCAAGAACACCCTGTACCTACAAATGAGCAGTC
TGAGGTCTGAGGACACGGCCTTGTATTTCTGTGCAAGGCTCTACCTGGGGTTTGACTACTGGGG
CCAAGGCACCACTCTCACAGTCTCCTCA (SEQ ID NO: 18)

MHC724LC.1 Variable light chain region amino acid sequence:

DIQMTQSPASQSASLGESVTITCLASQTIGTWLAWYQQKPKSPQLLIYAATSLADGVPSRFSG
SGSGTKFSFKISSLQAEDFVSYYCQQLYSIPWTFGGGTKLEIK (SEQ ID NO: 19)

MHC724LC.1 Variable light chain region nucleic acid sequence:

GACATTCAGATGACCCAGTCTCCTGCCTCCCAGTCTGCATCTCTGGGAGAAAGTGTCACCATCA
CATGCCTGGCAAGTCAGACCATTTGGTACATGGTTAGCATGGTATCAGCAGAAACCAGGGAAATC
TCCTCAGCTCCTGATTTATGCTGCAACCAGCTTGGCAGATGGGGTCCCATCAAGGTTTCAAGTGGT
AGTGGATCTGGCACAAAATTTCTTTCAAGATCAGCAGCCTACAGGCTGAAGATTTTGTAAGTT
ATTACTGTCAACAACCTTTACAGTATTCCGTGGACATTCGGTGGAGGCACCAAGCTGGAAATCAA
AC (SEQ ID NO: 20)

[000830] Binding of mouse antibodies m136-M13, m136-M19, m245-M3, m245-M5 and
m136-M14 to human PD-1 was confirmed by ELISA (Figure 1). Briefly, human PD-1-Fc (R and

D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified anti-PD-1 antibodies were applied to the plate in serial dilution and allowed to bind. Plates were washed with PBST (PBS, pH 7.2 + 0.05% Tween-20). Bound antibody was detected with an anti-mouse IgG-HRP conjugate (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prizm (Sigma Plot) and the data were fit to a model of single site saturation binding. Anti-PD-1 antibodies nivolumab (NV1) and/or pembrolizumab (PM1) were used as positive controls in binding assays with anti-human IgG-HRP conjugate, FAb-specific (Sigma, St Louis, MO) for detection. The K_d for each antibody tested is shown in Table 10 below:

Table 10 K_d values for tested antibodies:

Clone	K_d , nM
245-M3	0.22
245-M5	0.34
136-M13	0.19
136-M19	0.23
136-M14	0.19
NV1 nivo	0.28
PM1 pembro	0.35

[000831] Binding of mouse antibodies m136-M13, m136-M19, m245-M3, m245-M5 and m136-M14 to human PD-1 inhibited PD-1 binding to PDL1 and PDL2 in an inhibition ELISA assay (Figures 2-3). Inhibition ELISAs were performed as follows. Human PD-1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified anti-PD-1 antibodies were applied to the plate in serial dilution in the presence of 2nM biotinylated PD-L1 or 2nM biotinylated PDL2. Binding of biotinylated PD-L1 and PD-L2 was detected by Pierce™ Streptavidin-poly HRP conjugate (Thermo Scientific, Rockford, IL) and visualized with TMB. Plots were generated in Prizm (Sigma Plot) and the data were fit to a model of single site competition binding and an IC_{50} was determined. The IC_{50} values for the M13 and M14 antibodies are shown in Table 11 below:

Table 11: IC₅₀ values for antibodies tested

IC50 (nM)	hPDL1/Fc	hPDL2/Fc
M13	3.3	2
M14	4.8	2.8

EXAMPLE 2. Production and testing of humanized anti-PD-1 Antibodies

[000832] This example demonstrates that mouse antibodies of the disclosure that bind human PD-1 can be converted to humanized IgG antibodies that retain PD-1 binding and inhibition of PDL1 and PDL2 binding to PD-1.

[000833] Variable domains of the mouse anti-PD-1 antibodies produced as described in Example 1 were humanized and expressed as full length hIgG4/hKappa antibodies. Fully human IgGs anti-PD-1 antibodies were expressed from transiently transfected HEK-293 cells and purified from the culture supernatant by Protein A chromatography.

[000834] The sequences of the humanized antibodies used in the studies presented herein are shown below:

PD-1 A Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSV
KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGT LVT VSS (SEQ ID
NO: 21)

PD-1 A Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTTAGCGGCTACGCCATGAGCTGGGTGCGCCAGGCTCCTGGCAA
AGGCCTGGAATGGGTGGCCTACATCAGCAACAGCGGCGGCAATGCCCACTACGCCGATAGCGTG
AAGGGCCGGTTACCATCAGCCGGGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTACTGCACCAGAGAGGACTACGGCACCAGCCCCCTTCGT
GTATTGGGGCCAGGGTACCCTCGTGACCGTCTCCTCA (SEQ ID NO: 22)

PD-1 Ab Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVSYISNSGGNAHYADSV
KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCAKEDYGTSPFVYWGQGLTLVTVSS (SEQ ID
NO: 23)

PD-1 Ab Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTTAGCGGCTACGCCATGAGCTGGGTGCGCCAGGCTCCTGGCAA
AGGCCTGGAATGGGTGAGTTACATCAGCAACAGCGGCGGCAATGCCCACTACGCCGATAGCGTG
AAGGGCCGGTTCACCATCAGCCGGGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTACTGCGCCAAGGAGGACTACGGCACCAGCCCCTTCGT
GTATTGGGGCCAGGGTACCCTCGTGACCGTCTCCTCA (SEQ ID NO: 24)

PD-1 Ae Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNTHYADSV
KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCAREDYGTSPFVYWGQGLTLVTVSS (SEQ ID
NO: 25)

PD-1 Ae Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTTAGCGGCTACGCCATGAGCTGGGTGCGCCAGGCTCCTGGCAA
AGGCCTGGAATGGGTGGCCTACATCAGCAACAGCGGCGGCAATACCACTACGCCGATAGCGTG
AAGGGCCGGTTCACCATCAGCCGGGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGAGAGGACTACGGCACCAGCCCCTTCGT
GTATTGGGGCCAGGGTACCCTCGTGACCGTCTCCTCA (SEQ ID NO: 26)

PD-1 Af Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNTHYADSL
KGRFTVSRDN SKNTLYLQMNSLRAEDTAVYYCAREDYGTSPFVYWGQGLTLVTVSS (SEQ ID
NO: 27)

PD-1 Af Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGGAACTCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTTAGCGGCTACGCCATGAGCTGGGTGCGCCAGGCTCCTGGCAA
AGGCCTGGAATGGGTGGCCTACATCAGCAACAGCGGCGGCAATACCCACTACGCCGATAGCCTG
AAGGGCCGGTTACCGTCAGCCGGGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGAGAGGACTACGGCACCAGCCCCCTTCGT
GTATTGGGGCCAGGGTACCCTCGTGACCGTCTCCTCA (SEQ ID NO: 28)

PD-1 Ba Hv Variable heavy chain region amino acid sequence:

QVQLVQSGAEVKKPGASVKMSCKASGYTFTDYMDWVRQAPGQGLEWIGYIYPKNGGSSYAQKF
QGRATLTVDSTSTAYMELSSLRSEDTAVYYCARKVVATDYWGQGTLLTVSS (SEQ ID
NO: 29)

PD-1 Ba Hv Variable heavy chain region nucleic acid sequence:

CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCAGGCGCCAGCGTGAAGATGAGCT
GCAAGGCCAGCGGCTACACCTTCACCGACTACTACATGGACTGGGTGCGCCAGGCCCCCTGGACA
GGGACTGGAATGGATCGGCTACATCTACCCCAAGAACGGCGGCAGCAGCTACGCCCAGAAGTTC
CAGGGCAGAGCCACCCTGACCGTGGACACCAGCACAAGCACCGCCTACATGGAAGTGAAGCAGCC
TGCGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAAAGGTGGTGGCCACAGACTACTGGGG
CCAGGGTACCCTGCTGACCGTGTCTAGT (SEQ ID NO: 30)

PD-1 Bb Hv Variable heavy chain region amino acid sequence:

QVQLVQSGAEVKKPGASVKMSCKASGYTFTDYMDWVRQAPGQGLEWIGYIYPKNGGSSYAQKF
QGRATLTVDKSTSTAYMELSSLRSEDTAVYYCARKVVATDYWGQGTLLTVSS (SEQ ID
NO: 31)

PD-1 Bb Hv Variable heavy chain region nucleic acid sequence:

CAGGTGCAGCTGGTGCAGTCTGGCGCCGAAGTGAAGAAACCAGGCGCCAGCGTGAAGATGAGCT
GCAAGGCCAGCGGCTACACCTTCACCGACTACTACATGGACTGGGTGCGCCAGGCCCCCTGGACA
GGGACTGGAATGGATCGGCTACATCTACCCCAAGAACGGCGGCAGCAGCTACGCCCAGAAGTTC
CAGGGCAGAGCCACCCTGACCGTGGACAAGAGCACCGCCTACATGGAAGTGAAGCAGCC

TGCGGAGCGAGGACACCGCCGTGTACTACTGCGCCAGAAAGGTGGTGGCCACAGACTACTGGGG
CCAGGGTACCCTGCTGACCGTGTCTAGT (SEQ ID NO: 32)

PD-1 C Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSNYAMSWVRQAPGKGLEWVAYISNGGGDTHYADSL
KGRFTVSRDNSKNTLYLQMNSLRAEDTAVYYCARENYGTSPFVYWGQGLVTVSS (SEQ ID
NO: 33)

PD-1 C Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTCAGCAACTACGCCATGAGCTGGGTGCGCCAGGCCCCTGGAAA
AGGCCTGGAATGGGTGGCCTACATCAGCAACGGCGGAGGCGATACCCACTACGCCGATAGCCTG
AAGGGCCGGTTCACCGTGTCCAGAGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTATTGCGCCAGAGAGAACTACGGCACCAGCCCCTTCGT
GTACTGGGGCCAGGGTACCCTCGTGACCGTGTCTCT (SEQ ID NO: 34)

PD-1 Ca Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSNYAMSWVRQAPGKGLEWVAYISNQGGDTHYADSL
KGRFTVSRDNSKNTLYLQMNSLRAEDTAVYYCARENYGTSPFVYWGQGLVTVSS (SEQ ID
NO: 35)

PD-1 Ca Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
GTGCCGCCAGCGGCTTCACCTTCAGCAACTACGCCATGAGCTGGGTGCGCCAGGCCCCTGGAAA
AGGCCTGGAATGGGTGGCCTACATCAGCCAAGGCGGAGGCGATACCCACTACGCCGATAGCCTG
AAGGGCCGGTTCACCGTGTCCAGAGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTATTGCGCCAGAGAGAACTACGGCACCAGCCCCTTCGT
GTACTGGGGCCAGGGTACCCTCGTGACCGTGTCTCT (SEQ ID NO: 36)

PD-1 D Hv Variable heavy chain region amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAHSGFSSYDMSWVRQAPGKGLEWVATISGGGRYTYADSV
 KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCASNYYGFDYWGQGTLTVSS (SEQ ID
 NO: 37)

PD-1 D Hv Variable heavy chain region nucleic acid sequence:

GAAGTGCAGCTGGTGAATCTGGCGGCGGACTGGTGCAGCCTGGCGGATCTCTGAGACTGAGCT
 GTGCCACAGCGGCTTCAGCTTCAGCAGCTACGACATGAGCTGGGTGCGCCAGGCCCTGGCAA
 AGGACTGGAATGGGTGGCCACAATCAGCGGCGGAGGCCGGTACACCTACTACGCCGATAGCGTG
 AAGGGCCGGTTCACCATCAGCCGGGACAACAGCAAGAACACCCTGTACCTGCAGATGAACAGCC
 TGCGGGCCGAGGACACCGCCGTGTACTACTGCGCCAGCAACTACTACGGCTTCGACTACTGGGG
 CCAGGGTACCCTGCTGACCGTGTCTATCT (SEQ ID NO: 38)

PD-1 1.0 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDNYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPS
 RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 39)

PD-1 1.0 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
 CCTGTAGAGCCAGCGAGAGCGTGGACAACCTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
 GCGCGGCAAGGCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
 AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
 ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
 GCTGGAAATCAAG (SEQ ID NO: 40)

PD-1 1.1 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSVSVGDRATITCRASESVDNYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPS
 RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 41)

PD-1 1.1 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCCGTGTCCGTGGGCGACAGAGCCACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACAACCTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 42)

PD-1 1.2 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDQYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 43)

PD-1 1.2 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACCAATACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 44)

PD-1 1.4 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 45)

PD-1 1.4 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 46)

PD-1 1.5 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 47)

PD-1 1.5 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 48)

PD-1 1.6 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDNYGISFMNWFQQKPGKAPKLLIYAASDQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 49)

PD-1 1.6 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACAACTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCGATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 50)

PD-1 1.7 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSVSVGDRATITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKLEIK (SEQ ID NO: 51)

PD-1 1.7 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCCGTGTCCGTGGGCGACAGAGCCACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 52)

PD-1 1.9 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPWTFFGQGTKVEIK (SEQ ID NO: 53)

PD-1 1.9 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAA
GGTGGAAATCAAG (SEQ ID NO: 54)

PD-1 1.10 Lv Variable light chain region amino acid sequence:

DIQLTQSPSSLSASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPS
RFSGSGSGTDFTLTISSMQPEDFATYYCQQSKDVPYTFFGQGTKLEIK (SEQ ID NO: 55)

PD-1 1.10 Lv Variable light chain region nucleic acid sequence:

GACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
CCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAA
GCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGC
AGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGG
ACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTACACCTTTGGCCAGGGTACCAA
GCTGGAAATCAAG (SEQ ID NO: 56)

PD-1 2 Lv Variable light chain region amino acid sequence:

DIQMTQSPSSLSASVGDRVTMTCKSSQSLLYSSNQKNYLAWYQQKPGKAPKLLIFWASIRESGV
 PSRFSGSGSGTDFTLTISVQPEDFATYYCQQSDSYPWTFGQGTKLEIK (SEQ ID
 NO: 57)

PD-1 2 Lv Variable light chain region nucleic acid sequence:

GACATCCAGATGACCCAGAGCCCCAGCAGCCTGTCTGCCAGCGTGGGCGATAGAGTGACCATGA
 CCTGCAAGAGCAGCCAGAGCCTGCTGTACTCCAGCAACCAGAAGAAGTACCTGGCCTGGTATCA
 GCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTTCTGGGCCAGCATCCGGGAAAGCGGCGTG
 CCCAGCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGCGTGCAGC
 CCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCGACAGCTACCCCTGGACCTTTGGCCAGGG
 TACCAAGCTGGAAATCAAG (SEQ ID NO: 58)

PD-1 4 Lv Variable light chain region amino acid sequence:

DIQMTQSPSSLSASVGDRVTITCKASQDVGTAVAWYQQKPGKAPKLLIYWASTRHTGVPSRFSG
 SSGSGTDFTLTISVQPEDFATYYCQQYSSYPWTFGQGTKLEIK (SEQ ID NO: 59)

PD-1 4 Lv Variable light chain region nucleic acid sequence:

GACATCCAGATGACCCAGAGCCCCAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCA
 CATGCAAGGCCAGCCAGGACGTGGGAACAGCCGTGGCCTGGTATCAGCAGAAGCCTGGCAAGGC
 CCCCAGCTGCTGATCTACTGGGCCAGCACACACCGGCGTGCCCAGCAGATTTTCTGGC
 AGCGGCTCCGGCACCGACTTCACCCTGACAATCAGCAGCGTGCAGCCCGAGGACTTCGCCACCT
 ACTACTGCCAGCAGTACAGCAGCTACCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAATCAA
 G (SEQ ID NO: 60)

Kappa constant region amino acid sequence:

RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS
 TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 61)

Kappa constant region nucleic acid sequence:

CGTACGGTGGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGAA
CTGCCTCTGTTGTGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAAGGT
GGATAACGCCCTCCAATCGGGTAACTCCCAGGAGAGTGTACAGAGCAGGACAGCAAGGACAGC
ACCTACAGCCTCAGCAGCACCCCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACG
CCTGCGAAGTCACCCATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTG
T (SEQ ID NO: 62)

hIgG4 S228P amino acid sequence:

ASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVTVTPSSSLGKTYTCNVDHKPSNTKVDKRVESKYGPCCPPCPAPEFLGGPSVFLFPPKPK
DTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQD
WLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSD
IAVEWESNGQPENNYKTTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKS
LSLSLGK (SEQ ID NO: 63)

hIgG4 S228P nucleic acid sequence:

GCTAGCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCCCTGTAGCAGAAGCACCAGCGAGTCTA
CAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGTGGAATC
AGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTCTACAGTCCTCAGGACTCTACTCC
CTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCAAGACCTACACCTGTAACGTGG
ACCACAAGCCCAGCAACACCAAGGTGGACAAGCGGGTGAATCTAAGTACGGCCCTCCCTGCCC
TCCTTGCCCAGCCCCTGAATTTCTGGGGGGACCGTCAGTCTTCCTCTTCCCCCAAACCCAAG
GACACCCTCATGATCTCCCGGACCCCTGAGGTCACATGCGTGGTGGTGGACGTGAGCCAGGAAG
ACCCTGAGGTCCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCC
GCGGGAGGAGCAGTTCAACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGAC
TGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAGGCCTGCCCAGCTCCATCGAGA
AAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGTGTACACCCTGCCCCCATCCCA
GGAGGAGATGACCAAGAACCAGGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGAC
ATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACACTACAAGACCACGCCTCCCGTGC
TGGACTCCGACGGCTCCTTCTTCCTCTACAGCAGACTCACCGTGGACAAGAGCAGGTGGCAGGA

AGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGCAGAAGAGC
CTCTCCCTGTCTCTGGGTAAA (SEQ ID NO: 64)

[000835] These variable heavy chain regions (VH) and variable light chain regions (VL) can be used in a variety of combinations to produce anti-PD-1 antibodies of the disclosure. For example, the antibody referred to herein as A1.0 includes the VH sequence of SEQ ID NO: 21 and the VL sequence of SEQ ID NO: 39, the A1.5 antibody includes the VH sequence of SEQ ID NO: 21 and the VL sequence of SEQ ID NO: 47; the antibody referred to herein as C1.1 includes the VH sequence of SEQ ID NO: 33 and the VL sequence of SEQ ID NO: 41, and so on.

[000836] As shown in Figures 4-5, humanized anti-PD-1 antibodies bound to hPD-1 in a standard ELISA in a manner similar to that of nivolumab and/or pembrolizumab, and binding of the humanized antibodies inhibited the binding of PD-1 to PDL1 and PDL2 (Figures 6-9) in inhibition ELISAs. ELISAs were performed as follows. For hPD-1 binding ELISAs, human PD-1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified anti-PD-1 antibodies were applied to the plate in serial dilution and allowed to bind. Bound antibody was detected with an anti-human IgG-HRP conjugate (FAb-specific) (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prizm (Sigma Plot) and the data were fit to a model of single site saturation binding. For ligand inhibition ELISAs, human PD-1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified anti-PD-1 antibodies were applied to the plate in serial dilution in the presence of 2nM biotinylated hPD-L1 or 2nM biotinylated hPD-L2. Pierce™ Streptavidin-poly HRP conjugate (Thermo Scientific, Rockford, IL) and visualized with TMB. Plots were generated in Prizm (Sigma Plot) and the data were fit to a model of single site competition binding and an IC₅₀ was determined.

EXAMPLE 3: Anti-PD-1 antibodies shows specificity in binding

[000837] Example 3 shows that humanized anti-PD-1 antibody groups A1 and C1 of the disclosure bind specifically to hPD-1 by plate ELISA.

[000838] The binding of anti-PD-1 antibody A1.5 of the disclosure was highly specific to hPD-1-Fc in a standard ELISA against a panel of numerous human and mouse proteins (Figure

10). Binding of anti-PD-1 A1.5 was detected with an anti-human IgG-HRP conjugate (FAb-specific) (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prism (Sigma Plot) and the data were fit to a model of single site saturation binding.

EXAMPLE 4: Anti-PD-1 epitope binding

[000839] This example compares the epitopes bound by humanized anti-PD-1 antibodies of the disclosure, nivolumab, and pembrolizumab.

[000840] Nivolumab and pembrolizumab anti-PD-1 antibodies each bind human PD-1 and such binding by each inhibits the binding of PD-1 to PDL1 and PDL2. To map the epitopes of humanized groups A1 and C1 antibodies of the disclosure, anti-PD-1 inhibition ELISAs were performed, and compared with nivolumab and pembrolizumab inhibition. Dilution series of nivolumab, pembrolizumab, A1 and C1 antibodies were incubated in the presence of biotinylated nivolumab FAb or biotinylated pembrolizumab FAb at a concentration of 0.3nM in a standard plate ELISA format with hPD-1-Fc (R & D systems, Minneapolis, MN). Binding of biotinylated FAb was detected by Pierce™ Streptavidin-poly HRP conjugate (Thermo Scientific, Rockford, IL) and visualized with TMB (Thermo Scientific, Rockford IL) and 1N HCl. Antibodies belonging to the A1 and C1 groups of the disclosure blocked biotinylated Pembrolizumab FAb binding to PD-1 similarly to Pembrolizumab antibody and more completely than nivolumab antibody (Figure 11A). The same A1 and C1 antibodies of the disclosure blocked biotinylated nivolumab binding similarly to nivolumab antibody and more completely than pembrolizumab antibody (Figure 11B). The data in Figures 11A and 11B show that A1 and C1 antibodies of the disclosure completely block both nivolumab and pembrolizumab, whereas nivolumab and pembrolizumab incompletely block each other.

EXAMPLE 5: Anti-PD-1 A1.5 enhances CMV-stimulated cytokine secretion by PBMCs from a CMV-positive donor

[000841] In this example, peripheral blood mononuclear cells from a CMV-positive donor were incubated in the presence of CMV viral lysate and anti-PD-1 antibodies of the disclosure to assess the effect of such anti-PD-1 antibodies on interferon gamma (IFN-gamma, IFN γ , IFN-g, IFN γ , or IFN- γ) cytokine secretion.

[000842] PBMCs from a CMV-positive donor (Hemacare) were plated at 2×10^5 cells/well in the presence of CMV viral lysate (Astarte) and either anti-PD-1 antibody A1.5 of the disclosure or a hIgG4 isotype control antibody. After four days, supernatant was removed from each well, and IFN-gamma levels were assayed using IFN-gamma ELISA kit (Life Technologies, Carlsbad, CA) (Figure 12). Anti-PD-1 antibody A1.5 increased CMV-stimulated IFN-gamma secretion compared with control hIgG4 and with potency similar to anti-PD-1 nivolumab and pembrolizumab antibodies.

EXAMPLE 6: Anti-PD-1 antibodies bind to monomeric hPD-1 with high affinity and slow dissociation kinetics

[000843] In this example, humanized anti-PD-1 antibodies of the disclosure are shown to bind with high affinity and slow dissociation to monomeric PD-1.

[000844] Activatable antibodies can be activated singly, creating a monovalent binding moiety, or doubly, creating a bivalent binding moiety. Single-arm activation of antibodies with differing avidities can favor binding and biological activity of activatable antibodies with higher monovalent affinities. Anti-PD-1 antibodies A1.5 and Bba2 of the disclosure, as well as nivolumab and pembrolizumab were immobilized at equivalent densities on Forte-Bio Octet BioLayer Interferometry (Pall ForteBio, Menlo Park, CA) sensors and allowed to bind to a serial dilution of human PD-1-His (R & D systems, Minneapolis, MN) in solution. Kinetic analysis was performed with ForteBio Data Analysis software. Results (Figure 13) show PD-1 antibodies of the disclosure bind monomeric PD-1 with similar or higher affinities and similar or slower dissociation constants than nivolumab or pembrolizumab.

EXAMPLE 7: Activatable anti-PD-1 M13/A1.4/A1.5 antibody masking moieties

[000845] This example describes identification of masking moieties (MM) that reduce binding of anti-PD-1 antibodies of the disclosure to their target.

[000846] Anti-PD-1 antibodies m136-M13, A1.4 and A1.5 were used to screen libraries using a method similar to that described in PCT International Publication Number WO 2010/081173, published 15 July 2010. The screening consisted of one round of MACS and five rounds of FACS sorting. For the initial MACS, approximately 2×10^{11} cells were incubated with m136-M13 antibody at a concentration of 100nM, and 6×10^6 binders were collected using

Protein-G Dynabeads (Invitrogen). FACS rounds were conducted labeling cells with DyLight 650 (Thermo-Fisher) labeled m136-M19 antibody for FACS rounds 1-4 as follows: 100nM FACS round 1 (F1), 10nM FACS round 2 (F2), 2nM FACS round 3 (F3) and 1nM FACS round 4 (F4), with increasingly small percentages of binders as assessed by fluorescence collected at each round. For FACS round 5 (F5), cells were labeled with 1nM DyLight 650 labeled A1.5 antibody and the brightest 0.2-4% of cells were collected. Individual peptide clones from F3, F4 and F5 were identified by sequence analysis and subsequently verified for their ability to bind DyLight-650 A1.4 or DyLight-650 A1.5.

[000847] The sequences of the anti-PD-1 m136-M13, A1.4 and A1.5 masking moieties are listed in Table 12 (masking moiety PD001 is also referred to in the disclosure as PD01 and/or PD-01; masking moiety PD002 is also referred to herein as PD02 and/or PD-02, and so on):

Table 12: Masking Moieties

Mask	Amino Acid Sequence	SEQ ID NO:
PD001	AMSGCSWSAFCPYLA	66
PD002	DVNCAIWYSVCTTVP	67
PD003	LVCPLYALSSGVCMDG	68
PD004	SVNCRIWSAVCAGYE	69
PD005	MLVCSLQPTAMCERV	70
PD006	APRCYMFASYCKSQY	71
PD007	VGPCELTPKPV CNTY	72
PD008	ETCNQYERSSGLCFA	73
PD009	APRTCYTYQCSSFYT	74
PD010	GLCSWYLSSSGLCVD	75
PD011	VPWCQLTPRVMCMWA	76
PD012	NWLDCQFYSECSVYG	77
PD013	SCPLYVMSSFGGCWD	78
PD014	MSHCWMFSSSCDGVK	79
PD015	VSYCTWLIEVTCLRG	80
PD016	VLCAAYALSSGICGG	81

PD017	TTCNLYQQSSMFCNA	82
PD018	APRCYMFASYCKSQY	83
PD019	PCDQNPYFYYPYVCHA	84
PD020	SVCPMYALSSMLCGA	85
PD021	LSVECYVFSRCSSLP	86
PD022	FYCTYLVS LTCHPQ	87
PD023	SMAGCQWSSFCVQRD	88
PD024	IYSCYMFASRCTSDK	89
PD025	SRC SVYEVSSGLCDW	90
PD026	GMCSAYAYSSKLCTI	91
PD027	MTNTCNLLCQQFLT	92
PD028	FQPCLMFASSCFTSK	93
PD029	WNCHPAGVGPVFCEV	94
PD030	ALCSMYLASSGLCNK	95
PD031	NYLSCQFFQNCYETY	96
PD032	GWCLFSDMWLGLCSA	97
PD033	EFCARDWL PYQCSSF	98
PD034	TSYCSIEHYPCNTHH	99
PD035	PYICSSFPLDCQAGQ	100
PD036	VGCEWYMSSSGMCSR	101
PD037	EVC GGCSMQSVSCWP	102
PD038	FTECQLSPKAICMSN	103
PD039	KYCLFSEYVEGTCLN	104
PD040	SGCPMYAWGWDECWR	105
PD041	VDCP WYASSSAICSR	106
PD042	DMLLCQIRGSCAAWG	107
PD043	ECHPYQASASLWCGY	108
PD044	MMMGC MWSAWCPPSR	109
PD045	NAYFRCSLMCNMIMF	110
PD046	ACCKESVHSVHDCKR	111

PD047	ACIGINSYMSNYCYL	112
PD048	ANCSFLELTNKFCTI	113
PD049	AYCSYLMFASNPCI	114
PD050	CFTSKCPCLCYSLA	115
PD051	CLCRDINCWLGCST	116
PD052	CWCDIYCSPYQCSSF	117
PD053	DCIYYYQQSANLCSY	118
PD054	DCTGVNYYIDKHCTN	119
PD055	DECHGYLRSSGLCGG	120
PD056	DICSAYAASSGFCYY	121
PD057	DIICVLTPTAWCGRT	122
PD058	DNCCMYCSWWIACRD	123
PD059	DSCQWYMLSADLCGT	124
PD060	DSVCFSSSSFLCHKS	125
PD061	DTMCAIWWTVCSGGR	126
PD062	ECTYQTSSFHEACMS	127
PD063	EGCNLYERSSYGCNN	128
PD064	EGCTAFAMSAGICGG	129
PD065	EQSCSLTPIAFCWSE	130
PD066	EWCNAYISSSKLCST	131
PD067	FEVCYMFASACRNGM	132
PD068	FSCSWYAESSSLCDI	133
PD069	FVCQMFEASSGLCGG	134
PD070	FYCPCCMFASSCGSR	135
PD071	FYCSYLPGASHQCSH	136
PD072	FYCSYLYMCEVCCYE	137
PD073	GFCTQHTVLTWCPTS	138
PD074	GSCPSYAVSAGLCYA	139
PD075	GSQCFLTPTAFCTHT	140
PD076	GTCHPYMQSSKICNN	141

PD077	GVECFVFTGGCGGYG	142
PD078	HELCNGHWVPCCWAY	143
PD079	ICDSYYAVSSGLCLL	144
PD080	IGCAWYVSSAGWCSP	145
PD081	INLCWMFASECGEHH	146
PD082	KCWLAEMTNLEHCNM	147
PD083	KHCSDFAVSSRLCDR	148
PD084	KVCSSYASSSGLCGW	149
PD085	LDSCYMFASYCVQAV	150
PD086	LLACHPIFVTVCQTR	151
PD087	LLSCPYNPEHVCHTS	152
PD088	LMCSLYALSSNLCGR	153
PD089	LMWCVLFLWSWCCRI	154
PD090	LPICHLTPTAVCTHI	155
PD091	LSNMCLAFGSCLYAW	156
PD092	LSRCHPIWYTICQNP	157
PD093	LTQCMSVHKECGGYE	158
PD094	LVNCRIWSWVCEEAT	159
PD095	LYCSWYQMSSAVCKE	160
PD096	MECGWYALSARFCEV	161
PD097	MTCSPYAMSAHFCNE	162
PD098	MVCSLYAYSASLCGA	163
PD099	NALCWSTFSWWCDMD	164
PD-100	NFTCMLTPKAYCVQT	165
PD-101	NGACIFTLSWCTNKT	166
PD-102	NGCELYAAASGLCRT	167
PD-103	NIECSVFGRCCCDNY	168
PD-104	PACRPMFWNRSCDNI	169
PD-105	PCRVSNMFFPYNCLD	170
PD-106	PIMCM LLPESYCWIW	171

PD-107	PQSCYMFASLCMPNG	172
PD-108	PRCPQGLPLYQCSSF	173
PD-109	PSVECLVFKRCYALP	174
PD-110	PVCQRSATIYN CNWF	175
PD-111	QCAAYYISSFGGCSN	176
PD-112	QFGCFMLARDFCGTY	177
PD-113	QMMCPYNPEHKCHQK	178
PD-114	QRECWMFASSCNSKN	179
PD-115	QSNMCTTYICSSFNY	180
PD-116	QSRCHSLAPYLCSSF	181
PD-117	RAYCSLLFADSCNNN	182
PD-118	RCIGINQYIDSNCYN	183
PD-119	RLSCFMFASQCALEF	184
PD-120	RQCIILMNHRQCFFK	185
PD-121	RSCTPYMMSSSLCNT	186
PD-122	RYCHYWKMPLYECSSF	187
PD-123	SCVSLSWFDMLKCYE	188
PD-124	SDNCEIWWTVCSAAM	189
PD-125	SFCWSYLVSSGLCGV	190
PD-126	SMCMNNYGTTIMCGN	191
PD-127	SMVGCGWSTFCPSRG	192
PD-128	SSLHCANGHTCPFCL	193
PD-129	SVCSYYEESSGICSP	194
PD-130	SWCGWYAASSGVCAL	195
PD-131	TCISQTIDSYLNCVN	196
PD-132	TFCNLYTKSSNICMS	197
PD-133	TYCVFHEYLDNTCNN	198
PD-134	VATGCPNMLCGSWP	199
PD-135	VEYCSLLLGNRCDYW	200
PD-136	VGCNMYLMSAGLCVD	201

PD-137	VLYCSWDSGTCVGS	202
PD-138	VMFSCYYLETCA	203
PD-139	VRIGLCPESCLV	204
PD-140	VTCTYYATSSSL	205
PD-141	VTGCILLPKAWC	206
PD-142	VWCSIYIYSSNL	207
PD-143	WMLECQYNNTCN	208
PD-144	WPCSPLEYNNIC	209
PD-145	WTYDCHLNQTC	210
PD-146	YCSINMYLIGGN	211
PD-147	YFCSLYANSAGF	212
PD-148	YVSCYMFSSSCP	213

EXAMPLE 8: Activatable anti-PD-1 A1.4 and A1.5 Antibodies

[000848] This example describes examples of activatable anti-PD-1 A1.4 and A1.5 antibodies of the disclosure.

[000849] Activatable anti-PD-1 A1.4 antibodies comprising an anti-PD-1 M13 masking moiety, a cleavable moiety, and an anti-PD-1 A1.4 antibody of the disclosure and activatable anti-PD-1 A1.5 antibodies comprising an anti-PD-1 M13 or an anti-PD-1 A1.5 masking moiety, a cleavable moiety selected from the group consisting of a cleavable moiety, and an anti-PD-1 A1.5 antibody were produced according to methods similar to those described in PCT Publication Nos. WO 2009/025846 and WO 2010/081173, the contents of which are hereby incorporated by reference in their entirety. In some embodiments, the cleavable moiety was selected from the group consisting of a cleavable moiety referred to herein as “2001” and comprising the sequence ISSGLLSGRSDNH (SEQ ID NO: 214) and a cleavable moiety referred to herein as “3001” and comprising the sequence AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318). The amino acid and nucleic acid sequences of several activatable anti-PD-1 antibody variable domains of the disclosure are provided below. Antibodies were produced as hIgG4 containing a single amino acid substitution, S228P (Angal, et al. 1993. Mol Immunol 30:105-8.) HC and hK LC format.

[000850] In some embodiments, the activatable antibody also includes a spacer sequence. In some embodiments, the spacer is joined directly to the MM of the activatable antibody. In some embodiments, the spacer is joined directly to the MM of the activatable antibody in the structural arrangement from N-terminus to C-terminus of spacer-MM-CM-AB. In some embodiments, the spacer joined directly to the N-terminus of MM of the activatable antibody is selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. In some embodiments, the spacer includes at least the amino acid sequence QGQSGQG (SEQ ID NO: 362). In some embodiments, the spacer includes at least the amino acid sequence QGQSGQ (SEQ ID NO: 913). In some embodiments, the spacer includes at least the amino acid sequence QGQSG (SEQ ID NO: 914). In some embodiments, the spacer includes at least the amino acid sequence QGQS (SEQ ID NO: 915). In some embodiments, the spacer includes at least the amino acid sequence QGQ (SEQ ID NO: 916). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 917). In some embodiments, the spacer includes at least the amino acid residue Q. In some embodiments, the activatable antibody does not include a spacer sequence.

[000851] While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art appreciate that the activatable anti-PD-1 antibodies of the disclosure can include any suitable spacer sequence, such as, for example, a spacer sequence selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. . Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG (SEQ ID NO: 2045), QG (SEQ ID NO: 2046), and G. While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art will also appreciate that activatable anti-PD-1 antibodies of the disclosure in some embodiments do not include a spacer sequence.

Activatable Anti-PD-1 variable domains:

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD001 2001 (SEQ ID NO: 919)]

[QQQSGQG] [AMSGCSWSAFCPYLAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFRSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 215)

PD-1 1.4 PD001 2001 amino acid sequence:

AMSGCSWSAFCPYLAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFRSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1041)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD001 2001 (SEQ ID NO: 920)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGATGAGTGGGTGCTCGTGGTCTGCTTTTTGCCCCGTATT
TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 216)

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD002 2001 (SEQ ID NO: 921)]

[QQQSGQG] [DVNCAIWYSVCTTVPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFRSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 217)

PD-1 1.4 PD002 2001 amino acid sequence:

DVNCAIWYSVCTTVPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFRSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1042)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD002 2001 (SEQ ID NO: 922)]

[CAAGGCCAGTCTGGCCAAGGT] [GATGTTAATTGCGCTATTTGGTATTCGGTGTGCACTACTG
 TTCCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 218)

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD003 2001 (SEQ ID NO: 923)]

[QGQSGQG] [LVCPLYALSSGVCMGGGGSSSGS ISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 219)

PD-1 1.4 PD003 2001 amino acid sequence:

LVCPLYALSSGVCMGGGGSSSGS ISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT ISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1043)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD003 2001 (SEQ ID NO: 924)]

[CAAGGCCAGTCTGGCCAAGGT] [TTGGTTTGCCCTTTGTATGCATTGAGTTCTGGGGTGTGCA
 TGGGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 220)

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD008 2001 (SEQ ID NO: 925)]

[QGQSGQG] [ETCNQYERSSGLCFAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVTITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFQGQGTKLEIK] (SEQ ID NO: 221)

PD-1 1.4 PD008 2001 amino acid sequence:

ETCNQYERSSGLCFAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1044)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD008 2001 (SEQ ID NO: 926)]

[CAAGGCCAGTCTGGCCAAGGT] [GAGACTTGCAATCAGTATGAGAGGTCGAGTGGTTTGTGCT
 TTGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 222)

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD009 2001 (SEQ ID NO: 927)]

[QGQSGQG] [APRTCYTYQCSSFYTGGSSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 223)

PD-1 1.4 PD009 2001 amino acid sequence:

APRTCYTYQCSSFYTGGSSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1045)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD009 2001 (SEQ ID NO: 928)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGCCGCGGACGTGCTATACGTATCAGTGCTCTAGTTTTT
 ATACTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 224)

[spacer (SEQ ID NO: 362)] [PD-1 1.4 PD010 2001 (SEQ ID NO: 929)]

[QGQSGQG] [GLCSWYLSSSGLCVDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 225)

PD-1 1.4 PD010 2001 amino acid sequence:

GLCSWYLSSSGLCVDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDSYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1046)

[spacer (SEQ ID NO: 918)] [PD-1 1.4 PD010 2001 (SEQ ID NO: 930)]

[CAAGGCCAGTCTGGCCAAGGT] [GGTCTTTGCAGTTGGTATCTTAGTAGTTCGGGTTTGTGCG
TGGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGACATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACAGTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 226)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD01 2001 (SEQ ID NO: 931)]

[QGQSGQG] [AMSGCSWSAFCPYLAGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 227)

PD-1 1.5 PD01 2001 amino acid sequence:

AMSGCSWSAFCPYLAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTTITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1047)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD001 2001 (SEQ ID NO: 932)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGATGAGTGGGTGCTCGTGGTCTGCTTTTTGCCCCGTATT
 TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 228)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD002 2001 (SEQ ID NO: 933)]

[QGQSGQG] [DVNCAIWYSVCTTVPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVTTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 229)

PD-1 1.5 PD002 2001 amino acid sequence:

DVNCAIWYSVCTTVPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTTITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1048)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD002 2001 (SEQ ID NO: 934)]

[CAAGGCCAGTCTGGCCAAGGT] [GATGTTAATTGCGCTATTTGGTATTCGGTGTGCACTACTG
 TTCTTGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 230)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD003 2001 (SEQ ID NO: 935)]

[QGQSGQG] [LVCPLYALSSGVCMMGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 231)

PD-1 1.5 PD003 2001 amino acid sequence:

LVCPLYALSSGVCMMGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1049)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD003 2001 (SEQ ID NO: 936)]

[CAAGGCCAGTCTGGCCAAGGT] [TTGGTTTGGCCTTTGTATGCATTGAGTTCTGGGGTGTGCA
TGGGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 232)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD004 2001 (SEQ ID NO: 937)]

[QGQSGQG] [SVNCRIWSAVCAGYEGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 233)

PD-1 1.5 PD004 2001 amino acid sequence:

SVNCRISAVCAGYEGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1050)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD004 2001 (SEQ ID NO: 938)]

[CAAGGCCAGTCTGGCCAAGGT] [TCTGTGAATTGCCGGATTTGGTCGGCTGTTTGCGCGGGGT
 ATGAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 234)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD005 2001 (SEQ ID NO: 939)]

[QGQSGQG] [MLVCSLQPTAMCERVGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 235)

PD-1 1.5 PD005 2001 amino acid sequence:

MLVCSLQPTAMCERVGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1051)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD005 2001 (SEQ ID NO: 940)]

[CAAGGCCAGTCTGGCCAAGGT] [ATGCTTGTGTGCTCGTTGCAGCCTACTGCGATGTGCGAGC
 GGGTGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 236)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD006 2001 (SEQ ID NO: 941)]

[QGQSGQG] [APRCYMFASYCKSQYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 237)

PD-1 1.5 PD006 2001 amino acid sequence:

APRCYMFASYCKSQYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 941)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD006 2001 (SEQ ID NO: 942)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGCCTAGGTGCTATATGTTTGCGTCGTATTGCAAGAGTC
AGTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 238)

PD-1 1.5 PD006 2001 nucleic acid sequence:

GCGCCTAGGTGCTATATGTTTGCGTCGTATTGCAAGAGTCAGTATGGAGGTGGCTCGAGCGGCG
GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGATATCCA
GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
AGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC
CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC

ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
TCAAG (SEQ ID NO: 942)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD007 2001 (SEQ ID NO: 943)]

[QGQSGQG] [VGPCELTPKPVNTYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 239)

PD-1 1.5 PD007 2001 amino acid sequence:

VGPCELTPKPVNTYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1052)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD007 2001 (SEQ ID NO: 944)]

[CAAGGCCAGTCTGGCCAAGGT] [GTGGGGCCTTGCGAGTTGACGCCGAAGCCTGTTTGCAATA
CGTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 240)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD008 2001 (SEQ ID NO: 945)]

[QGQSGQG] [ETCNQYERSSGLCFAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 241)

PD-1 1.5 PD008 2001 amino acid sequence:

ETCNQYERSSGLCFAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1053)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD008 2001 (SEQ ID NO: 946)]

[CAAGGCCAGTCTGGCCAAGGT] [GAGACTTGCAATCAGTATGAGAGGTCGAGTGGTTTGTGCT
 TTGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 242)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD009 2001 (SEQ ID NO: 947)]

[QGQSGQG] [APRTCYTYQCSSFYTGGSSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 243)

PD-1 1.5 PD009 2001 amino acid sequence:

APRTCYTYQCSSFYTGGSSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1054)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD009 2001 (SEQ ID NO: 948)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGCCGCGGACGTGCTATACGTATCAGTGCTCTAGTTTTT
 ATACTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 244)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD010 2001 (SEQ ID NO: 949)]

[QGQSGQG] [GLCSWYLSSSGLCVDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 245)

PD-1 1.5 PD010 2001 amino acid sequence:

GLCSWYLSSSGLCVDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1055)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD010 2001 (SEQ ID NO: 950)]

[CAAGGCCAGTCTGGCCAAGGT] [GGTCTTTGCAGTTGGTATCTTAGTAGTTCGGGTTTGTGCG
TGGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 246)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD011 2001 (SEQ ID NO: 951)]

[QGQSGQG] [VPWCQLTPRVMCMWAGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 247)

PD-1 1.5 PD011 2001 amino acid sequence:

VPWCQLTPRVMCMWAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1056)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD011 2001 (SEQ ID NO: 952)]

[CAAGGCCAGTCTGGCCAAGGT] [GTGCCTTGGTGCCAGTTGACGCCGCGGGTTATGTGCATGT
 GGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 248)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD012 2001 (SEQ ID NO: 953)]

[QGQSGQG] [NWLDCQFYSECSVYGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 249)

PD-1 1.5 PD012 2001 amino acid sequence:

NWLDCQFYSECSVYGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 953)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD012 2001 (SEQ ID NO: 954)]

[CAAGGCCAGTCTGGCCAAGGT] [AATTGGTTGGATTGCCAGTTTTATTCTGAGTGCTCTGTTT
 ATGGTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 250)

PD-1 1.5 PD012 2001 nucleic acid sequence:

AATTGGTTGGATTGCCAGTTTTATTCTGAGTGCTCTGTTTATGGTGGAGGTGGCTCGAGCGGCG
GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGATATCCA
GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
AGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC
CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC
ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
TCAAG (SEQ ID NO: 954)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD013 2001 (SEQ ID NO: 955)]

[QGQSGQG] [SCPLYVMSSFGGCWDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 251)

PD-1 1.5 PD013 2001 amino acid sequence:

SCPLYVMSSFGGCWDGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1057)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD013 2001 (SEQ ID NO: 956)]

[CAAGGCCAGTCTGGCCAAGGT] [TCGTGCCCTTTGTATGTGATGTCTAGTTTTTGGTGGGTGCT
GGGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC

AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 252)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD014 2001 (SEQ ID NO: 957)]

[QGQSGQG] [MSHCWMFSSSCDGVKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 253)

PD-1 1.5 PD014 2001 amino acid sequence:

MSHCWMFSSSCDGVKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1058)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD014 2001 (SEQ ID NO: 958)]

[CAAGGCCAGTCTGGCCAAGGT] [ATGAGTCATTGCTGGATGTTTTTCGAGTTCTTGCGATGGGG
TGAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 254)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD015 2001 (SEQ ID NO: 959)]

[QGQSGQG] [VSYCTWLIIEVTCLRGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 255)

PD-1 1.5 PD015 2001 amino acid sequence:

VSYCTWLIIEVTCLRGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTTITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1059)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD015 2001 (SEQ ID NO: 960)]

[CAAGGCCAGTCTGGCCAAGGT] [GTTTCGTATTGCACGTGGTTGATTGAGGTGACTTGCCTGA
 GGGGTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 256)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD016 2001 (SEQ ID NO: 961)]

[QQQSGQG] [VLCAAYALSSGICGGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVTTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 257)

PD-1 1.5 PD016 2001 amino acid sequence:

VLCAAYALSSGICGGGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTTITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1060)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD016 2001 (SEQ ID NO: 962)]

[CAAGGCCAGTCTGGCCAAGGT] [GTTTTGTGCGCTGCTTATGCTTTGAGTTCGGGTATTTGCG
 GTGGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 258)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD017 2001 (SEQ ID NO: 963)]

[QGQSGQG] [TTCNLYQQSSMFCNAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 259)

PD-1 1.5 PD017 2001 amino acid sequence:

TTCNLYQQSSMFCNAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1061)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD017 2001 (SEQ ID NO: 964)]

[CAAGGCCAGTCTGGCCAAGGT] [ACGACTTGCAATCTGTATCAGCAGTCTTCTATGTTTTGCA
ATGCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 260)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD018 2001 (SEQ ID NO: 965)]

[QGQSGQG] [APRCYMFASYCKSQYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 261)

PD-1 1.5 PD018 2001 amino acid sequence:

APRCYMFASYCKSQYGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1062)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD018 2001 (SEQ ID NO: 966)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGCCTAGGTGCTATATGTTTGCGTCGTATTGCAAGAGTC
 AGTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 262)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD019 2001 (SEQ ID NO: 967)]

[QGQSGQG] [PCDQNPYFYYPYVCHAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 263)

PD-1 1.5 PD019 2001 amino acid sequence:

PCDQNPYFYYPYVCHAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 967)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD019 2001 (SEQ ID NO: 968)]

[CAAGGCCAGTCTGGCCAAGGT] [CCTTGCATCAGAATCCGTATTTTATCCGTATGTGTGCC
 ATGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 264)

PD-1 1.5 PD019 2001 nucleic acid sequence:

CCTTGCGATCAGAATCCGTATTTTTATCCGTATGTGTGCCATGCGGGAGGTGGCTCGAGCGGCG
GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGATATCCA
GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
AGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC
CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC
ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
TCAAG (SEQ ID NO: 968)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD020 2001 (SEQ ID NO: 969)]

[QGQSGQG] [SVCPMYALSSMLCGAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFQGQGTKLEIK] (SEQ ID NO: 265)

PD-1 1.5 PD020 2001 amino acid sequence:

SVCPMYALSSMLCGAGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFQGQGTKLEIK (SEQ ID NO: 1063)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD020 2001 (SEQ ID NO: 970)]

[CAAGGCCAGTCTGGCCAAGGT] [TCTGTGTGCCCTATGTATGCGTTGAGTTCTATGTTGTGCG
GTGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC

AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 266)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD021 2001 (SEQ ID NO: 971)]

[QGQSGQG] [LSVECYVFSRCSSLPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 267)

PD-1 1.5 PD021 2001 amino acid sequence:

LSVECYVFSRCSSLPGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1064)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD021 2001 (SEQ ID NO: 972)]

[CAAGGCCAGTCTGGCCAAGGT] [TTGTCTGTGGAGTGCTATGTGTTTTTCGCGGTGCAGTAGTC
TGCCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGCTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 268)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD022 2001 (SEQ ID NO: 973)]

[QGQSGQG] [FYCTYLVS LTCHPQGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASV
GDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 269)

PD-1 1.5 PD022 2001 amino acid sequence:

FYCTYLVSILTCHPQGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTITCRA
 SESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFAT
 YYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1065)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD022 2001 (SEQ ID NO: 974)]

[CAAGGCCAGTCTGGCCAAGGT] [TTTTATTGCACTTATTTGGTGTCTTTGACTTGCCATCCGC
 AGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCA
 CGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGAC
 AGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACT
 GGTTCACAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAG
 CGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGC
 ATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTG
 GCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 270)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD023 2001 (SEQ ID NO: 975)]

[QGQSGQG] [SMAGCQWSSFCVQRDGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 271)

PD-1 1.5 PD023 2001 amino acid sequence:

SMAGCQWSSFCVQRDGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVTITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1066)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD023 2001 (SEQ ID NO: 976)]

[CAAGGCCAGTCTGGCCAAGGT] [TCTATGGCGGGTTGCCAGTGGAGTTCGTTTTGCGTGACGC
 GGGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCACAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 272)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD024 2001 (SEQ ID NO: 977)]

[QGQSGQG] [IYSCYMFASRCTSDKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 273)

PD-1 1.5 PD024 2001 amino acid sequence:

IYSCYMFASRCTSDKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1067)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD024 2001 (SEQ ID NO: 978)]

[CAAGGCCAGTCTGGCCAAGGT] [ATTTATTCGTGCTATATGTTTGCTTCGCGGTGCACGTCTG
ATAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 274)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD002 3001 (SEQ ID NO: 979)]

[QGQSGQG] [DVNCAIWYSVCTTVPGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSL
SASVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDF
TLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 275)

PD-1 1.5 PD002 3001 amino acid sequence:

DVNCAIWYSVCTTVPGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSLSASVGDRVITI
 TCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISMQPE
 DFATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1068)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD002 3001 (SEQ ID NO: 980)]

[CAAGGCCAGTCTGGCCAAGGT] CAAGGCCAGTCTGGCCAAGGTGATGTTAATTGCGCTATTTG
 GTATTCCGGTGTGCACTACTGTTTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTG
 CTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGGAGGTGGCTCGAGCGGCGGCGCTGTGG
 GACTGCTGGCTCCTCCTGGTGGCCTGTCTGGCAGATCTGATAACCACGGAGGATCCGATATCCA
 GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
 GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
 AGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC
 CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC
 ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
 TCAAG] (SEQ ID NO: 276)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD-12 3001 (SEQ ID NO: 981)]

[QGQSGQG] [NWLDCQFYSECSVYGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSL
 SASVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDF
 TLTISMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 277)

PD-1 1.5 PD-12 3001 amino acid sequence:

NWLDCQFYSECSVYGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSLSASVGDRVITI
 TCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISMQPE
 DFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1069)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD-12 3001 (SEQ ID NO: 982)]

[CAAGGCCAGTCTGGCCAAGGT] [AATTGGTTGGATTGCCAGTTTTATTCTGAGTGCTCTGTTT
 ATGGTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGGAGGTGGCTCGAGCGGCGGCGCTGTGGGACTGCTGGCTCCTCCTGGT

GGCCTGTCTGGCAGATCTGATAACCACGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCA
 GCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGC
 TTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATC
 TACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCG
 ACTTCACCCTGACCATCAGCAGCATGCAGCCGAGGACTTCGCCACCTACTACTGCCAGCAGAG
 CAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID
 NO: 278)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD-16 3001 (SEQ ID NO: 983)]

[QGQSGQG] [VLCAAYALSSGICGGGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSL
 SASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDF
 TLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 279)

PD-1 1.5 PD-16 3001 amino acid sequence:

VLCAAYALSSGICGGGGGSSGGAVGLLAPPGGLSGRSDNHGGSDIQLTQSPSSLSASVGDRVTI
 TCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPE
 DFATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1070)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD-16 3001 (SEQ ID NO: 984)]

[CAAGGCCAGTCTGGCCAAGGT] [GTTTTGTGCGCTGCTTATGCTTTGAGTTCGGGTATTTGCG
 GTGGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGGAGGTGGCTCGAGCGGCGGCGCTGTGGGACTGCTGGCTCCTCCTGGT
 GGCCTGTCTGGCAGATCTGATAACCACGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCA
 GCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGC
 TTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATC
 TACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCG
 ACTTCACCCTGACCATCAGCAGCATGCAGCCGAGGACTTCGCCACCTACTACTGCCAGCAGAG
 CAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID
 NO: 280)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD025 2001 (SEQ ID NO: 985)]

[QGQSGQG] [SRCSVYEVSSGLCDWGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 281)

PD-1 1.5 PD025 2001 amino acid sequence:

SRCSVYEVSSGLCDWGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1071)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD025 2001 (SEQ ID NO: 986)]

[CAAGGCCAGTCTGGCCAAGGT] [TCTCGTTGCTCTGTGTATGAGGTTTCGTCGGGGCTGTGCG
ATTGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 282)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD026 2001 (SEQ ID NO: 987)]

[QGQSGQG] [GMCSAYAYSSKLCTIGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 283)

PD-1 1.5 PD026 2001 amino acid sequence:

GMCSAYAYSSKLCTIGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1072)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD026 2001 (SEQ ID NO: 988)]

[CAAGGCCAGTCTGGCCAAGGT] [GGGATGTGCTCGGCGTATGCTTATTCGAGTAAGTTGTGCA
CTATTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 284)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD027 2001 (SEQ ID NO: 989)]

[QGQSGQG] [MTTNTCNLLCQQFLTGGGSSSGS ISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 285)

PD-1 1.5 PD027 2001 amino acid sequence:

MTTNTCNLLCQQFLTGGGSSSGS ISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1073)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD027 2001 (SEQ ID NO: 990)]

[CAAGGCCAGTCTGGCCAAGGT] [ATGACTACGAATACTTGCAATCTGTTGTGCCAGCAGTTTT
TGACGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 286)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD028 2001 (SEQ ID NO: 991)]

[QGQSGQG] [FQPCLMFASSCFTSKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 287)

PD-1 1.5 PD028 2001 amino acid sequence:

FQPCLMFASSCFTSKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 991)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD028 2001 (SEQ ID NO: 992)]

[CAAGGCCAGTCTGGCCAAGGT] [TTTCAGCCGTGCCTGATGTTTGCGAGTAGTTGCTTTACTA
GTAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 288)

PD-1 1.5 PD028 2001 nucleic acid sequence:

TTTCAGCCGTGCCTGATGTTTGCGAGTAGTTGCTTTACTAGTAAGGGAGGTGGCTCGAGCGGCG
GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGATATCCA
GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
AGGCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC
CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC
ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
TCAAG (SEQ ID NO: 992)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD030 2001 (SEQ ID NO: 993)]

[QGQSGQG] [ALCSMYLASSGLCNKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 289)

PD-1 1.5 PD030 2001 amino acid sequence:

ALCSMYLASSGLCNKGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1074)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD030 2001 (SEQ ID NO: 994)]

[CAAGGCCAGTCTGGCCAAGGT] [GCGCTTTGCAGTATGTATCTTGCTAGTTCTGGGCTGTGCA
ATAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 290)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD033 2001 (SEQ ID NO: 995)]

[QGQSGQG] [EFCARDWLPHYQCSSFSGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 291)

PD-1 1.5 PD033 2001 amino acid sequence:

EFCARDWLPHYQCSSFSGGGSSGGSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1075)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD033 2001 (SEQ ID NO: 996)]

[CAAGGCCAGTCTGGCCAAGGT] [GAGTTTTTGCCTCGGGATTGGCTGCCGTATCAGTGCTCGA
GTTTTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA

TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 292)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD034 2001 (SEQ ID NO: 1028)]

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFQGQGTKLEIK] (SEQ ID NO: 1029)

PD-1 1.5 PD034 2001 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFQGQGTKLEIK (SEQ ID NO: 1028)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD034 2001 (SEQ ID NO: 1030)]

[CAAGGCCAGTCTGGCCAAGGT] [ACGTCATACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1031)

PD-1 1.5 PD034 2001 nucleic acid sequence:

ACGTCATACTGCAGTATTGAGCATTACCCCTGCAATACACATCATGGAGGTGGCTCGAGCGGCG
 GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGATCCGATATCCA
 GCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGCGACAGAGTGACCATCACCTGTAGA
 GCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGAACTGGTTCCAGCAGAAGCCCGGCA
 AGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGGCAGCGGCGTGCCAAGCAGATTTTC

CGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCAGCATGCAGCCCGAGGACTTCGCC
ACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCTTTGGCCAGGGTACCAAGCTGGAAA
TCAAG (SEQ ID NO: 1030)

[spacer (SEQ ID NO: 362)] [PD-1 1.5 PD035 2001 (SEQ ID NO: 997)]

[QGQSGQG] [PYICSSFPLDCQAGQGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFQGQTKLEIK] (SEQ ID NO: 293)

PD-1 1.5 PD035 2001 amino acid sequence:

PYICSSFPLDCQAGQGGGSSSGSISGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFQGQTKLEIK] (SEQ ID NO: 1076)

[spacer (SEQ ID NO: 918)] [PD-1 1.5 PD035 2001 (SEQ ID NO: 998)]

[CAAGGCCAGTCTGGCCAAGGT] [CCTTATATTGCTCTAGTTTTCCGTTGGATTGCCAGGCGG
GTCAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTGCCAGCGTGGGC
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 584)

EXAMPLE 9: Anti-PD-1 activatable antibodies of the disclosure

[000852] This example demonstrates that anti-PD-1 activatable antibodies of the disclosure can be made in a variety of combinations of MM, CM, VL, and VH domains, as well as in a variety of distinct Ig isotypes. In addition, this example demonstrates that anti-PD-1 activatable antibodies of the disclosure can be made in a variety of combinations of MM, CM, VLCDR1, VLCDR2, VLCDR3, VHCDR1, VHCDR2, and VHCDR3 domains, as well as in a variety of distinct Ig isotypes.

Table 13A

Mask Sequence	Substrates:	VL SEQ ID NO:	VH SEQ ID NO:	VL CDRs SEQ ID NO: CDR1 CDR2 CDR3	VH CDRs SEQ ID NO: CDR1 CDR2 CDR3
AMGCSWSAFCPYLA (SEQ ID NO: 66)	LSGRSDNH (SEQ ID NO: 294)	1	3	653 658 664	669 678 683
DVNCAIWYSVCTTVP (SEQ ID NO: 67)	TGRGPSWV (SEQ ID NO: 295)	5	7	654 659 665	670 679 684
LVCPLYALSSGVCMG (SEQ ID NO: 68)	PLTGRSGG (SEQ ID NO: 296)	9	11	655 660 666	671 680 685
SVNCRISAVCAGYE (SEQ ID NO: 69)	TARGPSFK (SEQ ID NO: 297)	13	15	656 661 667	672 681 686
MLVCSLQPTAMCERV (SEQ ID NO: 70)	NTLSGRSEHSG (SEQ ID NO: 298)	17	19	657 662 668	673 682 687
APRCYMFASYCKSQY (SEQ ID NO: 71)	NTLSGRSGNHGS (SEQ ID NO: 299)	21	39	663	674
VGPCELTPKPV CNTY (SEQ ID NO: 72)	TSTSGRSANPRG (SEQ ID NO: 300)	23	41		675
ETCNQYERSSGLCFA (SEQ ID NO: 73)	TSGRSANP (SEQ ID NO: 301)	25	43		676
APRTCITYQCSSFYT (SEQ ID NO: 74)	VHMPGLGLGP (SEQ ID NO: 302)	27	45		677
GLCSWYLSSSGLCVD (SEQ ID NO: 75)	AVGLLAPP (SEQ ID NO: 303)	29	47	VL CDRs of SEQ ID NO: 1	VH CDRs of SEQ ID NO: 3
VPWCQLTPRVCMCWA (SEQ ID NO: 76)	AQNLLGMV (SEQ ID NO: 304)	31	49	VL CDRs of SEQ ID NO: 5	VH CDRs of SEQ ID NO: 7
NWLDCCFYSECSVYG	QNQALRMA	33	51	VL CDRs of SEQ ID	VH CDRs of SEQ ID

(SEQ ID NO: 77)	(SEQ ID NO: 305)				NO: 9	NO: 11
SCPLYVMSSFGGCWD	LAAPLGLL	35	53	VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 78)	(SEQ ID NO: 306)			NO: 13	NO: 15	
MSHCWMFSSSCDGVK	STFPFGMF	37	55	VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 79)	(SEQ ID NO: 307)			NO: 17	NO: 19	
VSYCTWLIEVTCIRG	ISSGLLSS		57	VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 80)	(SEQ ID NO: 308)			NO: 21	NO: 39	
VLCAAYALSSGICGG	PAGLWLDP		59	VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 81)	(SEQ ID NO: 309)			NO: 23	NO: 41	
TTCNLVYQQSSMFCNA	VAGRSMRP			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 82)	(SEQ ID NO: 310)			NO: 25	NO: 43	
APRCYMFASYCKSQY	VVPEGRRS			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 83)	(SEQ ID NO: 311)			NO: 27	NO: 45	
PCDQNPYFYPYVCHA	ILPRSPAF			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 84)	(SEQ ID NO: 312)			NO: 29	NO: 47	
SVCPMYALSSMLCGA	MVLGRSLL			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 85)	(SEQ ID NO: 313)			NO: 31	NO: 49	
LSVECYVFSRCSSLP	QGRAITFI			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 86)	(SEQ ID NO: 314)			NO: 33	NO: 51	
FYCTYLVSLTCHPQ	SPRSIMLA			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 87)	(SEQ ID NO: 315)			NO: 35	NO: 53	
SMAGCQWSSFCVQRD	SMLRSMPL			VL CDRs of SEQ ID	VH CDRs of SEQ ID	
(SEQ ID NO: 88)	(SEQ ID NO: 316)			NO: 37	NO: 55	
IYSCYMFASRCTSDK	SARGPSRW				VH CDRs of SEQ ID	
(SEQ ID NO: 89)	(SEQ ID NO: 319)				NO: 57	
SRCSVYEVSSGLCDW	GWHTGRN				VH CDRs of SEQ ID	
(SEQ ID NO: 90)	(SEQ ID NO: 320)				NO: 59	

GC	HT	LC CDRs of SEQ ID	HC CDRs of SEQ ID
GCSEAYAYSSKLCIT (SEQ ID NO: 91)	HTGRSGAL (SEQ ID NO: 321)	NO: 543	NO: 546
MTTNTCNLLCQFLT	AARGPAIH		
(SEQ ID NO: 92)	(SEQ ID NO: 322)		
FQPCILMFASSCFTSK	RGPAFNPM		
(SEQ ID NO: 93)	(SEQ ID NO: 323)		
WNCHPAGVGPVFCEV	SSRGPAYL		
(SEQ ID NO: 94)	(SEQ ID NO: 324)		
ALCSMYLASSGLCNK	RGPATPIM		
(SEQ ID NO: 95)	(SEQ ID NO: 325)		
NYLSCQFFQNCYETY	RGPA		
(SEQ ID NO: 96)	(SEQ ID NO: 326)		
GWCLFSDMWLGLCSA	GGQPSGMWGW		
(SEQ ID NO: 97)	(SEQ ID NO: 327)		
EFCARDWLPLYQCSF	FPRPLGITGL		
(SEQ ID NO: 98)	(SEQ ID NO: 328)		
TSYCSIEHYPCNTHH	SPLTGRSG		
(SEQ ID NO: 99)	(SEQ ID NO: 329)		
PYICSSFPLDCQAGQ	SAGFSLPA		
(SEQ ID NO: 100)	(SEQ ID NO: 330)		
VGCEWYMSSSGMCSR	LAPLGLQRR		
(SEQ ID NO: 101)	(SEQ ID NO: 331)		
EVCGGCSMQSVSCWP	SGGPLGVR		
(SEQ ID NO: 102)	(SEQ ID NO: 332)		
FTECQLSPKAICMSN	PLGL		
(SEQ ID NO: 103)	(SEQ ID NO: 333)		
KYCLFSEYVEGTCLN	ISSGLSS		

(SEQ ID NO: 104)	(SEQ ID NO: 334)
SGCPMYAWGDEWCR	PVGYTSSL
(SEQ ID NO: 105)	(SEQ ID NO: 335)
VDCPWYASSSAICSR	DWLYWPGL
(SEQ ID NO: 106)	(SEQ ID NO: 336)
DMLLCQIRGSCAAWG	LKAAPRWA
(SEQ ID NO: 107)	(SEQ ID NO: 337)
ECHPYQASASLWCGY	GPSHLVLT
(SEQ ID NO: 108)	(SEQ ID NO: 338)
MMMGCMWSAWCPPSR	LPGLSPW
(SEQ ID NO: 109)	(SEQ ID NO: 339)
NAYFRCSLMCNMIMF	MGLFSEAG
(SEQ ID NO: 110)	(SEQ ID NO: 340)
ACCKESVHSVHDKR	SPLPLRVP
(SEQ ID NO: 111)	(SEQ ID NO: 341)
ACIGINSYMSNYCYL	RMHLRSLG
(SEQ ID NO: 112)	(SEQ ID NO: 342)
ANCSFLELTNKFCTI	LLAPSHRA
(SEQ ID NO: 113)	(SEQ ID NO: 343)
AYCSYLMFASNPCI	GPRSFG
(SEQ ID NO: 114)	(SEQ ID NO: 344)
CFTSKCPCLCYSLA	GPRSFG
(SEQ ID NO: 115)	(SEQ ID NO: 345)
CLCRDINCWLGCST	LSGRSGNH
(SEQ ID NO: 116)	(SEQ ID NO: 1157)
CWCDIYCSPYQCSF	SGRSANPRG
(SEQ ID NO: 117)	(SEQ ID NO: 1158)

DCIYYQQSANLCSY (SEQ ID NO: 118)	LSGRSDDH (SEQ ID NO: 1161)
DCTGVNYIDKHCTN (SEQ ID NO: 119)	LSGRSDIH (SEQ ID NO: 1162)
DECHGYLRSSGLCGG (SEQ ID NO: 120)	LSGRSDQH (SEQ ID NO: 1165)
DICSAYAASSGFCYY (SEQ ID NO: 121)	LSGRSDTH (SEQ ID NO: 1166)
DIICVLTPTAWCGRT (SEQ ID NO: 122)	LSGRSDYH (SEQ ID NO: 1169)
DNCCMYCSWWIACRD (SEQ ID NO: 123)	LSGRSDNP (SEQ ID NO: 1520)
DSCQWMLSadLCGT (SEQ ID NO: 124)	LSGRSANP (SEQ ID NO: 1695)
DSVCFSSSSFLCHKs (SEQ ID NO: 125)	LSGRSANI (SEQ ID NO: 1696)
DTMCAIWWTVCSGGR (SEQ ID NO: 126)	LSGRSDNI (SEQ ID NO: 1697)
ECTYQTSSFHEACMS (SEQ ID NO: 127)	MIAPVAYR (SEQ ID NO: 1698)
EGCNLYERSsSYGCNN (SEQ ID NO: 128)	RPSPWAY (SEQ ID NO: 1699)
EGCTAFAMSAGICGG (SEQ ID NO: 129)	WATPRPMR (SEQ ID NO: 1700)
EQSCSLTPIAFcWSE (SEQ ID NO: 130)	FRLLDWQW (SEQ ID NO: 1701)
EWCNAYISSSKLCST	ISSGL

(SEQ ID NO: 131) FEVCYMFASACRNGM	(SEQ ID NO: 1702) ISSGLLS
(SEQ ID NO: 132) FSCSWYAESSLCDI	(SEQ ID NO: 1703) ISSGLL
(SEQ ID NO: 133) FVCQMFEEASSGLCGG	(SEQ ID NO: 1704) ISSGLLSGRSDNH
(SEQ ID NO: 134) FYCPCCMFASSCGSR	(SEQ ID NO: 214) AVGLLAPPGGLSGRSDNH
(SEQ ID NO: 135) FYCSYLPGASHQCCH	(SEQ ID NO: 318) ISSGLLSGGSGGLSGRSDNH
(SEQ ID NO: 136) FYCSYLYMCEVCCYE	(SEQ ID NO: 346) ISSGLLSGGSGGLSGRSDNH
(SEQ ID NO: 137) GFCTQHTVLTWCPTS	(SEQ ID NO: 346) AVGLLAPPGGTSTSGRSANPRG
(SEQ ID NO: 138) GSCPSYAVSAGLCYA	(SEQ ID NO: 347) TSTSGRSANPRGGGAVGLLAPP
(SEQ ID NO: 139) GSQCFLTPTAFCTHT	(SEQ ID NO: 348) VHMPLGLPGGTSTSGRSANPRG
(SEQ ID NO: 140) GTCHPYMQSSKICNN	(SEQ ID NO: 349) TSTSGRSANPRGGGVHMPGLGLGP
(SEQ ID NO: 141) GVECFVFTGGCGGYG	(SEQ ID NO: 350) LSGRSDNHGGAVGLLAPP
(SEQ ID NO: 142) HELNCNHWVPCCWAY	(SEQ ID NO: 351) VHMPLGLPGGLSGRSDNH
(SEQ ID NO: 143) ICDSYYAVSSGLCLL	(SEQ ID NO: 352) LSGRSDNHGGVHMPGLGLGP
(SEQ ID NO: 144)	(SEQ ID NO: 353)

IGCAWYVSSAGWCSP (SEQ ID NO: 145)	LGRSDNHGGSGGSISSGLLSS (SEQ ID NO: 354)
INLCWMFASECGEHH (SEQ ID NO: 146)	LGRSGNHGGSGGSISSGLLSS (SEQ ID NO: 355)
KCWLAEMTNLEHCNM (SEQ ID NO: 147)	ISSGLLSSGGSGGSLSGRSGNH (SEQ ID NO: 356)
KHCSDFAYSRLCDR (SEQ ID NO: 148)	LGRSDNHGGSGGSGNQALRMA (SEQ ID NO: 357)
KVCSSYASSSGLCGW (SEQ ID NO: 149)	QNALRMAGGSGGSLSGRSDNH (SEQ ID NO: 358)
LDSCYMFASYCVQAV (SEQ ID NO: 150)	LGRSGNHGGSGGSGNQALRMA (SEQ ID NO: 359)
LLACHPIFVTVCQTR (SEQ ID NO: 151)	QNALRMAGGSGGSLSGRSGNH (SEQ ID NO: 360)
LLSCPYNPEHVCHTS (SEQ ID NO: 152)	ISSGLLSGRSGNH (SEQ ID NO: 361)
LMCSLYALSSNLCGR (SEQ ID NO: 153)	ISSGLLSSGGSGGSLSGRSGNH (SEQ ID NO: 1091)
LMWCVLFLWSWCCRI (SEQ ID NO: 154)	ISSGLLSGRSANPRG (SEQ ID NO: 1092)
LPICHLTPTAVCTHI (SEQ ID NO: 155)	AVGLLAPPTSGRSANPRG (SEQ ID NO: 1093)
LSNMCLAFGSCLYAW (SEQ ID NO: 156)	AVGLLAPPSGRSANPRG (SEQ ID NO: 1094)
LSRCHPIWYTICQNP (SEQ ID NO: 157)	ISSGLLSGRSDDH (SEQ ID NO: 1095)
LTQCMSVHKECGGYE	ISSGLLSGRSDIH

(SEQ ID NO: 158)	(SEQ ID NO: 1096)
LVNCRISWVCEEAT	ISSGLLSGRSDQH
(SEQ ID NO: 159)	(SEQ ID NO: 1097)
LYCSWYQMSSAVCKE	ISSGLLSGRSDTH
(SEQ ID NO: 160)	(SEQ ID NO: 1098)
MECGWYALSARFCEV	ISSGLLSGRSDYH
(SEQ ID NO: 161)	(SEQ ID NO: 1099)
MTCSPIYAMSAHFCNE	ISSGLLSGRSDNP
(SEQ ID NO: 162)	(SEQ ID NO: 1100)
MVCSLYAYSASLCGA	ISSGLLSGRSANP
(SEQ ID NO: 163)	(SEQ ID NO: 1101)
NALCWSTFSWWCDMD	ISSGLLSGRSANI
(SEQ ID NO: 164)	(SEQ ID NO: 1102)
NFTCMLTPKAYCVQT	AVGLLAPPGGLSGRSDDH
(SEQ ID NO: 165)	(SEQ ID NO: 1103)
NGACIFTLWCNTKT	AVGLLAPPGGLSGRSDIH
(SEQ ID NO: 166)	(SEQ ID NO: 1104)
NGCELYAAAAGLCRT	AVGLLAPPGGLSGRSDQH
(SEQ ID NO: 167)	(SEQ ID NO: 1105)
NIECSVFGRCCCDNY	AVGLLAPPGGLSGRSDTH
(SEQ ID NO: 168)	(SEQ ID NO: 1106)
PACRPMFWNRSCDNI	AVGLLAPPGGLSGRSDYH
(SEQ ID NO: 169)	(SEQ ID NO: 1107)
PCRVSNNMFFPYNCLD	AVGLLAPPGGLSGRSDNP
(SEQ ID NO: 170)	(SEQ ID NO: 1108)
PIMCMLLPESYCIWI	AVGLLAPPGGLSGRSANP
(SEQ ID NO: 171)	(SEQ ID NO: 1109)

PQSCYMFASLCMPNG (SEQ ID NO: 172)	AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110)
PRCPQGLPLYQCSF	ISSGLLSGRSDNI
(SEQ ID NO: 173)	(SEQ ID NO: 1111)
PSVECLVFKRCYALP	AVGLLAPPGGLSGRSDNI
(SEQ ID NO: 174)	(SEQ ID NO: 1112)
PVCQRSATIYNCNWF	
(SEQ ID NO: 175)	
QCAAYYISSFGGCSN	
(SEQ ID NO: 176)	
QFGCFMLARDFCGTY	
(SEQ ID NO: 177)	
QMMCPYNPEHKCHQK	
(SEQ ID NO: 178)	
QRECFMFASSCNSKN	
(SEQ ID NO: 179)	
QSNMCTTYICSSFNY	
(SEQ ID NO: 180)	
QSRCHSLAPYLCSSF	
(SEQ ID NO: 181)	
RAYCSLLFADSCNNN	
(SEQ ID NO: 182)	
RCIGINQYIDSNYN	
(SEQ ID NO: 183)	
RLSCFMFASQCALEF	
(SEQ ID NO: 184)	
RQCIILMNHRCFFK	

(SEQ ID NO: 185)	RSCTPYMMSSSLCNT
(SEQ ID NO: 186)	RYCHYWKMPYECSSF
(SEQ ID NO: 187)	SCVSLSWFDMCLKYE
(SEQ ID NO: 188)	SDNCEIWWTVCSAAM
(SEQ ID NO: 189)	SFCWSYLVSSGLCGV
(SEQ ID NO: 190)	SMCMNNYGTTIMCGN
(SEQ ID NO: 191)	SMVCGGWSTFCPSRG
(SEQ ID NO: 192)	SSLHCANGHTCPFCL
(SEQ ID NO: 193)	SVCSSYYEESGICSP
(SEQ ID NO: 194)	SWCGWYAASSGVCAL
(SEQ ID NO: 195)	TCISQTIDSYLNCVN
(SEQ ID NO: 196)	TFCNLTKSSNICMS
(SEQ ID NO: 197)	TYCVFHEYLDNTCNN
(SEQ ID NO: 198)	

VATGCPNMLCGSWP
(SEQ ID NO: 199)
VEYCSLLLGNRCDYW
(SEQ ID NO: 200)
VGCNMYLMSAGLCVD
(SEQ ID NO: 201)
VLYCSWDSGTCVGS
(SEQ ID NO: 202)
VMFSCYYLETCPGV
(SEQ ID NO: 203)
VRIGLCPESCLVSGF
(SEQ ID NO: 204)
VTCTYYATSSSLCNT
(SEQ ID NO: 205)
VTGCILLPKAWCWGD
(SEQ ID NO: 206)
VMCSIYIYSSNLCSR
(SEQ ID NO: 207)
WMLECYNNNTCNMT
(SEQ ID NO: 208)
WPCSPLEYNNICNV
(SEQ ID NO: 209)
WTYDCHLNQTCPTY
(SEQ ID NO: 210)
YCSINMYLIGNCMY
(SEQ ID NO: 211)
YFCSLYANSAGFCGG

(SEQ ID NO: 212) YVSCYMFSSSCPSTW (SEQ ID NO: 213)					
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Table 13B

Mask Sequence	Substrates:	VL SEQ ID NO:	VH SEQ ID NO:	VL CDRs SEQ ID NO: CDR1 CDR2 CDR3	VH CDRs SEQ ID NO: CDR1 CDR2 CDR3
RYCHAAANPDRFCGIY (SEQ ID NO: 1206) PRVCSTDGGDYCLLP (SEQ ID NO: 1207) PRPQCHHRHNCPDHP (SEQ ID NO: 1208) KCSRPAHQNPDRCSR (SEQ ID NO: 1209) ASYRCPDYKCSHTKH (SEQ ID NO: 1210) LPRCPDHPIKCIETK (SEQ ID NO: 1211) YTFGCPDRYCDRAAT (SEQ ID NO: 1212) RGCPDFNPPSHCYTA (SEQ ID NO: 1213) RDYCGPQSPDYCHEI (SEQ ID NO: 1214) PNKPCPDLQCYVTNY (SEQ ID NO: 1215)	LSGRSDNH (SEQ ID NO: 294) TGRGPSWV (SEQ ID NO: 295) PLTGRSGG (SEQ ID NO: 296) TARGPSFK (SEQ ID NO: 297) NTLSGRSENHSG (SEQ ID NO: 298) NTLSGRSGNHGS (SEQ ID NO: 299) TSTSGRSANPRG (SEQ ID NO: 300) TSGRSANP (SEQ ID NO: 301) VHMPGLGFLGP (SEQ ID NO: 302) AVGLLAPP (SEQ ID NO: 303)	626	1346	VL CDRs of SEQ ID NO: 626 1705 1706 1707	VH CDRs of SEQ ID NO: 1346 1708 1709 1710

PRVACGEPLDLCYSNT (SEQ ID NO: 1216)	AQNLLGMV (SEQ ID NO: 304)
RGCKKHTISTLTCPD (SEQ ID NO: 1217)	QNQALRMA (SEQ ID NO: 305)
PAYRCPPDRPPCKNQM (SEQ ID NO: 1218)	LAAPLGLL (SEQ ID NO: 306)
NARCPYFPGDNCHMN (SEQ ID NO: 1220)	STFFFGMF (SEQ ID NO: 307)
PTLRCPDRWCYDSPR (SEQ ID NO: 1221)	ISSGLLSS (SEQ ID NO: 308)
PSNLCPDKWCQTWRS (SEQ ID NO: 1222)	PAGLWLDP (SEQ ID NO: 309)
TPRYCAASYCPAHGY (SEQ ID NO: 1223)	VAGRSMRP (SEQ ID NO: 310)
RPGCGAVSPRCPDAP (SEQ ID NO: 1224)	VVPEGRRS (SEQ ID NO: 311)
VLRCHKQNPDCNNH (SEQ ID NO: 1225)	ILPRSPAF (SEQ ID NO: 312)
GVKSCREPDFCSRGS (SEQ ID NO: 1226)	MVLGRSLL (SEQ ID NO: 313)
RNNLCPDYSCNNHNS (SEQ ID NO: 1227)	QGRAITFI (SEQ ID NO: 314)
RAACHRLNPDACTNG (SEQ ID NO: 1228)	SPRSIMLA (SEQ ID NO: 315)
VCQSDRIPDYVTCTD (SEQ ID NO: 1229)	SMLRSMPL (SEQ ID NO: 316)
RNCRIASINPDYCNI	SARGPSRW

(SEQ ID NO: 1230)	(SEQ ID NO: 319)
KEWRCPDYKCKPSYH	GWHTGRN
(SEQ ID NO: 1231)	(SEQ ID NO: 320)
NLRICHKSLCPDYIK	HTGRSGAL
(SEQ ID NO: 1232)	(SEQ ID NO: 321)
NTHKCSNTNICPSFN	AARGPAIH
(SEQ ID NO: 1233)	(SEQ ID NO: 322)
STRYCQASQCQMSPY	RGPAFNPM
(SEQ ID NO: 1234)	(SEQ ID NO: 323)
THRFTASLCNKNTS	SSRGPAYL
(SEQ ID NO: 1235)	(SEQ ID NO: 324)
YTLNTRSPDWCPNK	RGPATPIM
(SEQ ID NO: 1236)	(SEQ ID NO: 325)
IRCTTGQSPDYCPQS	RGPA
(SEQ ID NO: 1237)	(SEQ ID NO: 326)
RCNQPDKNDQMLCNI	GGQPSGMWGW
(SEQ ID NO: 1238)	(SEQ ID NO: 327)
GTCRTDHQSPDYCY	FPRPLGITGL
(SEQ ID NO: 1239)	(SEQ ID NO: 328)
RGCFRSGDSLGMCPD	SPLTGRSG
(SEQ ID NO: 1240)	(SEQ ID NO: 329)
SGCFDSNEHRHCSRI	SAGFSLPA
(SEQ ID NO: 1241)	(SEQ ID NO: 330)
NRCMKLWYNPDCVAR	LAPLGLQRR
(SEQ ID NO: 1242)	(SEQ ID NO: 331)
PLCARPHYWSPCDQS	SGGPLGVR
(SEQ ID NO: 1243)	(SEQ ID NO: 332)

DSKCHPNSPDYCFNS (SEQ ID NO: 1244) NGSCRPLGGDFCGNR (SEQ ID NO: 1245) KTRCIEMSGDYCAKS (SEQ ID NO: 1246) IRPCMYNWGDLCNQF (SEQ ID NO: 1247) VKTCMENNPDYCYNN (SEQ ID NO: 1248) LRMCFEASGDYCDQQ (SEQ ID NO: 1249) IRKCQLDGPDQCMLT (SEQ ID NO: 1250) KWKCHKNNPNYCNRR (SEQ ID NO: 1251) RTMCLDTNPDYCQSH (SEQ ID NO: 1252) LAACHSMDSHRCPDY (SEQ ID NO: 1253) RSPCIHNATMCPDYT (SEQ ID NO: 1254) MPRCPDWPPRCSMVI (SEQ ID NO: 1255) VRQLCRLPDYCPGK (SEQ ID NO: 1256) PRPFCQAQSLNCPDRA	PLGL (SEQ ID NO: 333) ISSGLSS (SEQ ID NO: 334) PVGYTSSL (SEQ ID NO: 335) DWLYWPGI (SEQ ID NO: 336) LKAAPRWA (SEQ ID NO: 337) GPSHLVLT (SEQ ID NO: 338) LPGGLSPW (SEQ ID NO: 339) MGLFSEAG (SEQ ID NO: 340) SPLPLRVP (SEQ ID NO: 341) RMHLRSLG (SEQ ID NO: 342) LLAPSHRA (SEQ ID NO: 343) GPRSFGL (SEQ ID NO: 344) GPRSGG (SEQ ID NO: 345) LSGRSGNH
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(SEQ ID NO: 1257)	(SEQ ID NO: 1157)
SFGRCTLVRTCDFM	SGRSANPRG
(SEQ ID NO: 1258)	(SEQ ID NO: 1158)
RDKPCPDFSCATIH	LSGRSDDH
(SEQ ID NO: 1259)	(SEQ ID NO: 1161)
ATKPCPDRWCTMSTL	LSGRSDIH
(SEQ ID NO: 1260)	(SEQ ID NO: 1162)
SSNRCPDLRCTHHNM	LSGRSDQH
(SEQ ID NO: 1261)	(SEQ ID NO: 1165)
RGSMCPDLHC SLSHI	LSGRSDTH
(SEQ ID NO: 1262)	(SEQ ID NO: 1166)
NYQRCPDRTCMHNII	LSGRSDYH
(SEQ ID NO: 1263)	(SEQ ID NO: 1169)
QKRPCPDRKCHAHYN	LSGRSDNP
(SEQ ID NO: 1264)	(SEQ ID NO: 1520)
QNHRCPDRWCNKTTN	LSGRSANP
(SEQ ID NO: 1265)	(SEQ ID NO: 1695)
RLNLCPDKKHCHMTNL	LSGRSANI
(SEQ ID NO: 1266)	(SEQ ID NO: 1696)
PQDRCPDKRCTNPGN	LSGRSDNI
(SEQ ID NO: 1267)	(SEQ ID NO: 1697)
SRWRCPDYKCEHGKY	MIAPVAYR
(SEQ ID NO: 1268)	(SEQ ID NO: 1698)
YENQCPDLYCNRYSM	RPSPMWAY
(SEQ ID NO: 1269)	(SEQ ID NO: 1699)
TARSCPVFNC PDNNS	WATPRPMR
(SEQ ID NO: 1270)	(SEQ ID NO: 1700)

MDQRCPDAWCTSKPK (SEQ ID NO: 1271)	FRLLDWQW (SEQ ID NO: 1701)
GDLRCPDRLCPRHSL (SEQ ID NO: 1272)	ISSGL (SEQ ID NO: 1702)
IQYLCPDYHCKASNN (SEQ ID NO: 1273)	ISSGLLS (SEQ ID NO: 1703)
QHHRCPDRYCNSNN (SEQ ID NO: 1274)	ISSGLL (SEQ ID NO: 1704)
TVALCPDYSCYHINN (SEQ ID NO: 1275)	ISSGLLSGRSDNH (SEQ ID NO: 214)
SPWRCPDRYCLSNHD (SEQ ID NO: 1276)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318)
SSKRCPDRFCNKTHA (SEQ ID NO: 1277)	ISSGLSSGGSGGLSGRSDNH (SEQ ID NO: 346)
HTDRCPDYKCSQNH (SEQ ID NO: 1278)	ISSGLSSGGSGGLSGRSDNH (SEQ ID NO: 346)
SRSNCTPQRCNSDYH (SEQ ID NO: 1279)	AVGLLAPPGGTSTSGRSANPRG (SEQ ID NO: 347)
FAARCPDKYCAIHTN (SEQ ID NO: 1280)	TSTSGRSANPRGGGAVGLLAPP (SEQ ID NO: 348)
GSARCPDILVCQQTQ (SEQ ID NO: 1281)	VHMPILGFLGPGGTSTSGRSANPRG (SEQ ID NO: 349)
RNLMPDKFCNKNTK (SEQ ID NO: 1282)	TSTSGRSANPRGGVHMPILGFLGP (SEQ ID NO: 350)
NIRLCPDKVCTPTWV (SEQ ID NO: 1283)	LSGRSDNHGGAVGLLAPP (SEQ ID NO: 351)
MTDLCPDAHCAKTHM	VHMPILGFLGPGGLSGRSDNH

(SEQ ID NO: 1284)	(SEQ ID NO: 352)
PYSRLCAYPCPDFVG	LSGRSDNHGGVHMPFLGLGP
(SEQ ID NO: 1285)	(SEQ ID NO: 353)
LCGCARSPDYCKCRG	LSGRSDNHGGSGGSISSGLLSS
(SEQ ID NO: 1286)	(SEQ ID NO: 354)
WGRCERVDPCCCPRG	LSGRSGNHGGSGGSISSGLLSS
(SEQ ID NO: 1287)	(SEQ ID NO: 355)
TRNTCHTRICYGMAC	ISSGLLSSGGSGGSLSGRSGNH
(SEQ ID NO: 1288)	(SEQ ID NO: 356)
CVCTSCSSYWTLCPD	LSGRSDNHGGSGGSGNQALRMA
(SEQ ID NO: 1289)	(SEQ ID NO: 357)
LCCSRGSCNCPDRCTW	QNQALRMAGGSGGSLSGRSDNH
(SEQ ID NO: 1290)	(SEQ ID NO: 358)
CCPLCQANMCPDNQS	LSGRSGNHGGSGGSGNQALRMA
(SEQ ID NO: 1291)	(SEQ ID NO: 359)
ECKLCCPDLYCGGTM	QNQALRMAGGSGGSLSGRSGNH
(SEQ ID NO: 1292)	(SEQ ID NO: 360)
CSNPMCAIYCCPDIL	ISSGLLSGRSGNH
(SEQ ID NO: 1293)	(SEQ ID NO: 361)
CPRCNTYSKHDCYHQ	ISSGLLSSGGSGGSLSGRSH
(SEQ ID NO: 1294)	(SEQ ID NO: 1091)
FCCASKMPAPSNCHT	ISSGLLSGRSGNH
(SEQ ID NO: 1295)	(SEQ ID NO: 1092)
	AVGLLAPPTSGRSGNH
	(SEQ ID NO: 1093)
	AVGLLAPPSGRSGNH
	(SEQ ID NO: 1094)

ISSGLLSGRSDDH
(SEQ ID NO: 1095)
ISSGLLSGRSDIH
(SEQ ID NO: 1096)
ISSGLLSGRSDQH
(SEQ ID NO: 1097)
ISSGLLSGRSDTH
(SEQ ID NO: 1098)
ISSGLLSGRSDYH
(SEQ ID NO: 1099)
ISSGLLSGRSDNP
(SEQ ID NO: 1100)
ISSGLLSGRSANP
(SEQ ID NO: 1101)
ISSGLLSGRSANI
(SEQ ID NO: 1102)
AVGLLAPPGGLSGRSDDH
(SEQ ID NO: 1103)
AVGLLAPPGGLSGRSDIH
(SEQ ID NO: 1104)
AVGLLAPPGGLSGRSDQH
(SEQ ID NO: 1105)
AVGLLAPPGGLSGRSDTH
(SEQ ID NO: 1106)
AVGLLAPPGGLSGRSDYH
(SEQ ID NO: 1107)
AVGLLAPPGGLSGRSDNP

	(SEQ ID NO: 1108) AVGLLAPPGGLSGRSANP (SEQ ID NO: 1109) AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110) ISSGLLSGRSDNI (SEQ ID NO: 1111) AVGLLAPPGGLSGRSDNI (SEQ ID NO: 1112)				
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Table 13C

Mask Sequence	Substrates:	VL SEQ ID NO:	VH SEQ ID NO:	VL CDRs SEQ ID NO: CDR1 CDR2 CDR3	VH CDRs SEQ ID NO: CDR1 CDR2 CDR3
GCDFTSAKHNCGSW (SEQ ID NO: 1351) VGSNCWTGPACALTS (SEQ ID NO: 1352) FCAVMFDFLSDRCLH (SEQ ID NO: 1353) FCPPWLDYLGKNCMT (SEQ ID NO: 1354) MSCWDFSSAQGGQH (SEQ ID NO: 1355) LMCADLHYNHYNCKY (SEQ ID NO: 1356) ELCGWQSFSGVCTSE (SEQ ID NO: 1357)	LSGRSDNH (SEQ ID NO: 294) TGRGPSWV (SEQ ID NO: 295) PLTGRSGG (SEQ ID NO: 296) TARGPSFK (SEQ ID NO: 297) NTLSGRSENHSG (SEQ ID NO: 298) NTLSGRSGNHGS (SEQ ID NO: 299) TSTSGRSANPRG (SEQ ID NO: 300)	638	1514	VL CDRs of SEQ ID NO: 638 1711 1712 1713	VH CDRs of SEQ ID NO: 1514 1714 1715 1716

WTYENCWASCQPHLE (SEQ ID NO: 1358) KLTEDFSSAA (SEQ ID NO: 1359) VGQSCFSGLVCDRQL (SEQ ID NO: 1360) ISHYCFSGKSCRD (SEQ ID NO: 1361) HCIPDFTSAAGDCMR (SEQ ID NO: 1362) RLVSAYSFS (SEQ ID NO: 1363) KFHHSHPLVHDF TSA (SEQ ID NO: 1364) ASYPDFSSANGVGLR (SEQ ID NO: 1365) GLATTLSNVDF TSAG (SEQ ID NO: 1366) DFTSANSASFSGDAST (SEQ ID NO: 1367) GRLPGHVVDF TSAW (SEQ ID NO: 1368) SGSFYSSSAFDF TSA (SEQ ID NO: 1369) CDDFTSAQHSRINEC (SEQ ID NO: 1370) CDFTSAQGKKCR TAI	TSGRSANP (SEQ ID NO: 301) VHMPLGFLGP (SEQ ID NO: 302) AVGLLAPP (SEQ ID NO: 303) AQNLIGMV (SEQ ID NO: 304) QNQALRMA (SEQ ID NO: 305) LAAPLGLL (SEQ ID NO: 306) STFFFGMF (SEQ ID NO: 307) ISSGLLSS (SEQ ID NO: 308) PAGLWLDP (SEQ ID NO: 309) VAGRSMRP (SEQ ID NO: 310) VVPEGRRS (SEQ ID NO: 311) ILPRSPAF (SEQ ID NO: 312) MVLGRSLL (SEQ ID NO: 313) QGRAITFI
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(SEQ ID NO: 1371)	(SEQ ID NO: 314)
YYIDKYQSPSYGPVL	SPRSIMLA
(SEQ ID NO: 1372)	(SEQ ID NO: 315)
FSVARARSSADFTSS	SMLRSMPL
(SEQ ID NO: 1373)	(SEQ ID NO: 316)
DSDFTSAGSADRSR	SARGPSRW
(SEQ ID NO: 1374)	(SEQ ID NO: 319)
CDFTSATSISKRCDH	GWHTGRN
(SEQ ID NO: 1375)	(SEQ ID NO: 320)
IESSASSWGLQASRN	HTGRSGAL
(SEQ ID NO: 1376)	(SEQ ID NO: 321)
PRYHNLFNFTTPALSPGS	AARGPAIH
(SEQ ID NO: 1377)	(SEQ ID NO: 322)
DLFARFPIDRDFTSA	RGPAFNPM
(SEQ ID NO: 1378)	(SEQ ID NO: 323)
HCNFTTPPYCSSTLW	SSRGPAYL
(SEQ ID NO: 1379)	(SEQ ID NO: 324)
NVPIILLTDRQLLSG	RGPATPIM
(SEQ ID NO: 1380)	(SEQ ID NO: 325)
NPTACDFTSSMATYC	RGPA
(SEQ ID NO: 1381)	(SEQ ID NO: 326)
FVRTVRFNSMFSVP	GGQPSGMWGW
(SEQ ID NO: 1382)	(SEQ ID NO: 327)
YDFSSASNSPSRQT	FPRPLGITGL
(SEQ ID NO: 1383)	(SEQ ID NO: 328)
AHPDFSSAMRGNLLG	SPLTGRSG
(SEQ ID NO: 1384)	(SEQ ID NO: 329)

SSHVVHKDFTSANSR (SEQ ID NO: 1385) CPDFTSANGGCWQM (SEQ ID NO: 1386) SLGQSYPTDFTCPGC (SEQ ID NO: 1387) ASMRSEQRDFTSAY (SEQ ID NO: 1388) SCQFWFTLCGGVFH (SEQ ID NO: 1389) PYPNNRTGMHDFTSA (SEQ ID NO: 1390) KPPFIDFTSAGTSGT (SEQ ID NO: 1391) SIKSFIPRDDFTSAA (SEQ ID NO: 1392) GIKNPATPPFVDFTSA (SEQ ID NO: 1393) LSHTYPRGSSTIEAS (SEQ ID NO: 1394) PSLDFSSAT (SEQ ID NO: 1395) AFTPRIAPTFDVMKE (SEQ ID NO: 1396) LCGLQIPPPDCERS (SEQ ID NO: 1397) AAKMVSHSERDFTSA	SAGFSLPA (SEQ ID NO: 330) LAPLGLQRR (SEQ ID NO: 331) SGGPLGVR (SEQ ID NO: 332) PLGL (SEQ ID NO: 333) ISSGLSS (SEQ ID NO: 334) PVGYTSSL (SEQ ID NO: 335) DWLYWPGI (SEQ ID NO: 336) LKAAPRWA (SEQ ID NO: 337) GPSHLVLT (SEQ ID NO: 338) LPGGLSPW (SEQ ID NO: 339) MGLFSEAG (SEQ ID NO: 340) SPLPLRVP (SEQ ID NO: 341) RMHLRSLG (SEQ ID NO: 342) LLAPSHRA
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(SEQ ID NO: 1398)	(SEQ ID NO: 343)
VSVECFSGMQCPHY	GPRSFGL
(SEQ ID NO: 1399)	(SEQ ID NO: 344)
ASKCRLPCMASTQIY	GPRSFG
(SEQ ID NO: 1400)	(SEQ ID NO: 345)
GLRSCNIYFSIPCTY	LSGRSGNH
(SEQ ID NO: 1401)	(SEQ ID NO: 1157)
RGTS DGTLDFTTARS	SGRSANPRG
(SEQ ID NO: 1402)	(SEQ ID NO: 1158)
SMYPASRLILHPQYP	LSGRSDDH
(SEQ ID NO: 1403)	(SEQ ID NO: 1161)
HCISCYDFTSAAGSF	LSGRSDIH
(SEQ ID NO: 1404)	(SEQ ID NO: 1162)
SSGRWGDWACARIC	LSGRSDQH
(SEQ ID NO: 1405)	(SEQ ID NO: 1165)
RVFSDFTSASHSFGG	LSGRSDTH
(SEQ ID NO: 1406)	(SEQ ID NO: 1166)
TDRHSASGRDFTSAH	LSGRSDYH
(SEQ ID NO: 1407)	(SEQ ID NO: 1169)
AHCEDFSSAERIATMGC	LSGRSDNP
(SEQ ID NO: 1408)	(SEQ ID NO: 1520)
ACDPYFSFIPCDRL	LSGRSANP
(SEQ ID NO: 1409)	(SEQ ID NO: 1695)
NSPFTLSHIYDR (SEQ	LSGRSANI
ID NO: 1410)	(SEQ ID NO: 1696)
IGTNFTTSPSAFVAFP	LSGRSDNI
(SEQ ID NO: 1411)	(SEQ ID NO: 1697)

RDAPFIYRNADFSTP (SEQ ID NO: 1412)	MIAPVAYR (SEQ ID NO: 1698)
SIPNASSYNFTSSG (SEQ ID NO: 1413)	RPSPMWAY (SEQ ID NO: 1699)
AGIPDKRHTYDFTSA (SEQ ID NO: 1414)	WATPRPMR (SEQ ID NO: 1700)
WPLAHDRDWNFTTP (SEQ ID NO: 1415)	FRLLDWQW (SEQ ID NO: 1701)
RHSPSSGHVDFTSAG (SEQ ID NO: 1416)	ISSGL (SEQ ID NO: 1702)
SCFAWTDVPVWNRCSW (SEQ ID NO: 1417)	ISSGLLS (SEQ ID NO: 1703)
MPCDWTGPGKIWCGG (SEQ ID NO: 1418)	ISSGLL (SEQ ID NO: 1704)
RDCDFSTANFRSCNK (SEQ ID NO: 1419)	ISSGLLSGRSDNH (SEQ ID NO: 214)
LSCVWSPNYLHCNDH (SEQ ID NO: 1420)	AVGLLAPPGGLSGRSDNH (SEQ ID NO: 318)
FVCGLYSFGVCQGV (SEQ ID NO: 1421)	ISSGLLSGGGGSLSGRSDNH (SEQ ID NO: 346)
IGLMCFSGLQCPMLA (SEQ ID NO: 1422)	ISSGLLSGGGGSLSGRSDNH (SEQ ID NO: 346)
PGMNCFSGEICQMST (SEQ ID NO: 1423)	AVGLLAPPGGTSTSGRSANPRG (SEQ ID NO: 347)
GDVGSCWASGGLQGG (SEQ ID NO: 1424)	TSTSGRSANPRGGGAVGLLAPP (SEQ ID NO: 348)
SQFQDCWASCGASFT	VHMPILGLPGGTSTSGRSANPRG

(SEQ ID NO: 1425)	(SEQ ID NO: 349)
VGSLNCWYSCGDIWL	TSTSGRSANPRGGGVHMPGLGFLGP
(SEQ ID NO: 1426)	(SEQ ID NO: 350)
MCESWLNFLGDQCGM	LSGRSDNHGGGAVGLLAPP
(SEQ ID NO: 1427)	(SEQ ID NO: 351)
RCMISQSSFSFGMCGM	VHMPGLGFLGPGGLSGRSDNH
(SEQ ID NO: 1428)	(SEQ ID NO: 352)
NCAPWTSNMSNHCLK	LSGRSDNHGGGVHMPGLGFLGP
(SEQ ID NO: 1429)	(SEQ ID NO: 353)
LCGVGSATGLELCGV	LSGRSDNHGGGGSISSGLLSS
(SEQ ID NO: 1430)	(SEQ ID NO: 354)
GCDFFSLGGRQPCTP	LSGRSGNHGGGGSISSGLLSS
(SEQ ID NO: 1431)	(SEQ ID NO: 355)
MGCNFTTYPYHTCNT	ISSGLLSSGGGSLSGRSGNH
(SEQ ID NO: 1432)	(SEQ ID NO: 356)
GSCDFTSGAGKKCGS	LSGRSDNHGGGGSQNQALRMA
(SEQ ID NO: 1433)	(SEQ ID NO: 357)
VSCDFTSSHARMCSR	QNQALRMAGGGGSLSGRSDNH
(SEQ ID NO: 1434)	(SEQ ID NO: 358)
MRCTDFYNNHTNCIG	LSGRSGNHGGGGSQNQALRMA
(SEQ ID NO: 1435)	(SEQ ID NO: 359)
RSCDFTSAANKYCAT	QNQALRMAGGGGSLSGRSGNH
(SEQ ID NO: 1436)	(SEQ ID NO: 360)
LYCDSFVPRPNCAP	ISSGLLSGRSGNH
(SEQ ID NO: 1437)	(SEQ ID NO: 361)
NSCDFTSARVSKCST	ISSGLLSSGGGSLSGRSGNH
(SEQ ID NO: 1438)	(SEQ ID NO: 1091)

STCSDNFTTPMPCNT	ISSGLLSGRSANPRG
(SEQ ID NO: 1439)	(SEQ ID NO: 1092)
DICNDRPNLTHCHYF	AVGLLAPPTSGRSANPRG
(SEQ ID NO: 1440)	(SEQ ID NO: 1093)
LRCDDFTSAIGCRGY	AVGLLAPPSGRSANPRG
(SEQ ID NO: 1441)	(SEQ ID NO: 1094)
EGCDFTSALHSCNNY	ISSGLLSGRSDDH
(SEQ ID NO: 1442)	(SEQ ID NO: 1095)
RKGGDFTSASCFW	ISSGLLSGRSDIH
(SEQ ID NO: 1443)	(SEQ ID NO: 1096)
GMLCAGSSFGLCESM	ISSGLLSGRSDQH
(SEQ ID NO: 1444)	(SEQ ID NO: 1097)
RESCFGSSIGLCTNK	ISSGLLSGRSDTH
(SEQ ID NO: 1445)	(SEQ ID NO: 1098)
ILRCYDIPTNCMNFN	ISSGLLSGRSDYH
(SEQ ID NO: 1446)	(SEQ ID NO: 1099)
NSECTFGAMYCRNKP	ISSGLLSGRSDNP
(SEQ ID NO: 1447)	(SEQ ID NO: 1100)
ASGCFDEDIRCSGGA	ISSGLLSGRSANP
(SEQ ID NO: 1448)	(SEQ ID NO: 1101)
HYFCNQSNPSCQTAP	ISSGLLSGRSANI
(SEQ ID NO: 1449)	(SEQ ID NO: 1102)
AMGCELGGAGCIGSP	AVGLLAPPGGLSGRSDDH
(SEQ ID NO: 1450)	(SEQ ID NO: 1103)
TLKCHMPRKL CandP	AVGLLAPPGGLSGRSDIH
(SEQ ID NO: 1451)	(SEQ ID NO: 1104)
RPACRDLPHNCITST	AVGLLAPPGGLSGRSDQH

(SEQ ID NO: 1452) QMSCHGNFTTCHSNP (SEQ ID NO: 1453) LTGCARGARPCRLRV (SEQ ID NO: 1454) WSELCLAGPSCGWVG (SEQ ID NO: 1455) VTKSCWQLPHCITAP (SEQ ID NO: 1456) KAASCPHNQICNMTA (SEQ ID NO: 1457) VSKNCFSGMMCPVFA (SEQ ID NO: 1458) NRSSCWGTGPTCHVLH (SEQ ID NO: 1459) ARTGCSGPVCLNDVS (SEQ ID NO: 1460) STRTCLAFTCINGNT (SEQ ID NO: 1461) MLDGNCWYACSYKNT (SEQ ID NO: 1462) FSRSDCW\$ACAPWRV (SEQ ID NO: 1463) GGRMDCWASCQPLSR (SEQ ID NO: 1464) NSPHSCMTNCDFTSA (SEQ ID NO: 1465)	(SEQ ID NO: 1105) AVGLLAPPGGLSGRSDTH (SEQ ID NO: 1106) AVGLLAPPGGLSGRSDYH (SEQ ID NO: 1107) AVGLLAPPGGLSGRSDNP (SEQ ID NO: 1108) AVGLLAPPGGLSGRSANP (SEQ ID NO: 1109) AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110) ISSGLLSGRSDNI (SEQ ID NO: 1111) AVGLLAPPGGLSGRSDNI (SEQ ID NO: 1112)
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Table 13D

Mask Sequence	Substrates:	LC SEQ ID NO:	HC SEQ ID NO:	LC CDRs SEQ ID NO: CDR1 CDR2 CDR3	HC CDRs SEQ ID NO: CDR1 CDR2 CDR3
DYTYCRWVNWCLSGV (SEQ ID NO: 384)	LSGRSDNH (SEQ ID NO: 294)	543	546	LC CDRs of SEQ ID NO: 543	HC CDRs of SEQ ID NO: 546
ILCPEDPWGHKCKLP (SEQ ID NO: 385)	TGRGPSWV (SEQ ID NO: 295)				
TNIWSCQTYCDHKHK (SEQ ID NO: 386)	PLTGRSGG (SEQ ID NO: 296)				
SDHKCKLQNCMTKV (SEQ ID NO: 387)	TARGPSFK (SEQ ID NO: 297)				
PGNCHPMQKEMCQFI (SEQ ID NO: 388)	NTLSGRSENHSG (SEQ ID NO: 298)				
VEHLCTHKNCKHPD (SEQ ID NO: 389)	NTLSGRSGNHGS (SEQ ID NO: 299)				
TIPRCGQHPKCKDTL (SEQ ID NO: 390)	TSTSGRSANPRG (SEQ ID NO: 300)				
ACRICQDHPKTKWNS (SEQ ID NO: 391)	TSGRSANP (SEQ ID NO: 301)				
LIQCTGNLDHKCKHY (SEQ ID NO: 392)	VHMPGLGLGP (SEQ ID NO: 302)				
IPCHHSADHKHKCTS (SEQ ID NO: 393)	AVGLLAPP (SEQ ID NO: 303)				
SRQICADYNCHNKYK (SEQ ID NO: 394)	AQNLLGMV (SEQ ID NO: 304)				
QPCNPQIDHKIKCIY	QNQALRMA				

(SEQ ID NO: 395) HYTICMTHNKCKDMA	(SEQ ID NO: 305) LAAPLGLL
(SEQ ID NO: 396) ANSLAVEHKCKHNY	(SEQ ID NO: 306) STFPFGMF
(SEQ ID NO: 397) AALHCTEHKCKNHIK	(SEQ ID NO: 307) ISSGLLS
(SEQ ID NO: 398) APCIINTVDWKSCEI	(SEQ ID NO: 308) PAGLWLD
(SEQ ID NO: 399) ATNWC TH KQCKQDM	(SEQ ID NO: 309) VAGRMRP
(SEQ ID NO: 400) DCYNEHKLKTRVCNN	(SEQ ID NO: 310) VVPEGRRS
(SEQ ID NO: 401) DEM Q CSHKQCKTNSK	(SEQ ID NO: 311) ILPRSPAF
(SEQ ID NO: 402) DVGICS Q HNKCKPTK	(SEQ ID NO: 312) MVLGRSLL
(SEQ ID NO: 403) EKYC S DDHKCKITL	(SEQ ID NO: 313) QGRAITFI
(SEQ ID NO: 404) ELEC S HNKVNCIQI	(SEQ ID NO: 314) SPRSIMLA
(SEQ ID NO: 405) ELHPCNTHKCKPIVN	(SEQ ID NO: 315) SMLRSMPL
(SEQ ID NO: 406) EVGSCNHPKCKSN NY	(SEQ ID NO: 316) SARGPSRW
(SEQ ID NO: 407) EYSPSLAHPK L KDNA	(SEQ ID NO: 319) GWH T GRN
(SEQ ID NO: 408)	(SEQ ID NO: 320)

FESLHPKGKHPEDLG (SEQ ID NO: 409)	HTGRSGAL (SEQ ID NO: 321)
FPLCVRADRVCGDAQ (SEQ ID NO: 410)	AARGPAIH (SEQ ID NO: 322)
FQAPPASHNKLKPSL (SEQ ID NO: 411)	RGPAFNPM (SEQ ID NO: 323)
GAIDSCHHKCKSPHY (SEQ ID NO: 412)	SSRGPAYL (SEQ ID NO: 324)
GKIYTCEHNCTFGYS (SEQ ID NO: 413)	RGPATPIM (SEQ ID NO: 325)
HCTVNNHSSDHKCKI (SEQ ID NO: 414)	RGPA (SEQ ID NO: 326)
HGTQCTHNKCKPILS (SEQ ID NO: 415)	GGQPSGMWGW (SEQ ID NO: 327)
HIGWCLHPKCKTTTT (SEQ ID NO: 416)	FPRPLGITGL (SEQ ID NO: 328)
HLRTCIQKWCEHKMK (SEQ ID NO: 417)	SPLTGRSG (SEQ ID NO: 329)
HTDCTMMSNHKCKIN (SEQ ID NO: 418)	SAGFSLPA (SEQ ID NO: 330)
IRQQCTALACLLKVH (SEQ ID NO: 419)	LAPLGLQRR (SEQ ID NO: 331)
KGCSTHKMRAYCNQM (SEQ ID NO: 420)	SGGPLGVR (SEQ ID NO: 332)
KMFTPCKIWCNNSYN (SEQ ID NO: 421)	PLGL (SEQ ID NO: 333)
KTMCSGHKQKCNSSS	ISSGLSS

(SEQ ID NO: 422)	(SEQ ID NO: 334)
LACHSASLVDPKCKL	PVGYTSSL
(SEQ ID NO: 423)	(SEQ ID NO: 335)
LCNVSMDDHKHKPCYL	DWLYWPGL
(SEQ ID NO: 424)	(SEQ ID NO: 336)
LGLNCFSEHKCKEHM	LKAAPRWA
(SEQ ID NO: 425)	(SEQ ID NO: 337)
LGTCTHKHKNCNYTL	GPSHLVLT
(SEQ ID NO: 426)	(SEQ ID NO: 338)
LHEGCTTHNKCKPIA	LPGGLSPW
(SEQ ID NO: 427)	(SEQ ID NO: 339)
LKRCTGHWTCYTNW	MGLFSEAG
(SEQ ID NO: 428)	(SEQ ID NO: 340)
LQRCETHKEKYCHAIH	SPLPLRVP
(SEQ ID NO: 429)	(SEQ ID NO: 341)
LSHCYDHHKRCYSYIV	RMHLRSLG
(SEQ ID NO: 430)	(SEQ ID NO: 342)
LSKCHNKEKNCNNN	LLAPSHRA
(SEQ ID NO: 431)	(SEQ ID NO: 343)
MDTCEMHKQKCRPSF	GPRSFGL
(SEQ ID NO: 432)	(SEQ ID NO: 344)
MHNECLTHKCKVPIT	GPRSFGL
(SEQ ID NO: 433)	(SEQ ID NO: 345)
MLTLCNTNACHKEKN	LSGRSGNH
(SEQ ID NO: 434)	(SEQ ID NO: 1157)
MRPCLNNLEHKCKHY	SGRSANPRG
(SEQ ID NO: 435)	(SEQ ID NO: 1158)

MSRCPTRHKMCSLNI (SEQ ID NO: 436)	LSGRSDDH (SEQ ID NO: 1161)
MWICQEHKCLKMTDT (SEQ ID NO: 437)	LSGRSDIH (SEQ ID NO: 1162)
MYICKRRSAFYCTLN (SEQ ID NO: 438)	LSGRSDQH (SEQ ID NO: 1165)
NDCQHDKQMHKCKMH (SEQ ID NO: 439)	LSGRSDTH (SEQ ID NO: 1166)
NFGPCPMLLGCFGFR (SEQ ID NO: 440)	LSGRSDYH (SEQ ID NO: 1169)
NHTDCSHPKCKSHDS (SEQ ID NO: 441)	LSGRSDNP (SEQ ID NO: 1520)
NLNCPHKQKNCDKYH (SEQ ID NO: 442)	LSGRSANP (SEQ ID NO: 1695)
NPQCTPIDHKCKTHH (SEQ ID NO: 443)	LSGRSANI (SEQ ID NO: 1696)
NTTSCTHPKCKHQGK (SEQ ID NO: 444)	LSGRSDNI (SEQ ID NO: 1697)
NVGGCDNYGCHKLKN (SEQ ID NO: 445)	MIAPVAYR (SEQ ID NO: 1698)
PCSPGNLTWDHKCKY (SEQ ID NO: 446)	RPSPWAY (SEQ ID NO: 1699)
PFTKCHGFNKCKEHT (SEQ ID NO: 447)	WATPRPMR (SEQ ID NO: 1700)
PGDKCTHKEKCYNN (SEQ ID NO: 448)	FRLLDWQW (SEQ ID NO: 1701)
PNICNLDHKRCRIN	ISSGL

(SEQ ID NO: 449)	(SEQ ID NO: 1702)
PQLACKHPKCKDAGN	ISSGLLS
(SEQ ID NO: 450)	(SEQ ID NO: 1703)
PSCTMWTHGGVCKHA	ISSGLL
(SEQ ID NO: 451)	(SEQ ID NO: 1704)
PSHRHPLAKPGRVE	ISSGLLSGRSDNH
(SEQ ID NO: 452)	(SEQ ID NO: 214)
PTCFKTHNKSCKNRV	AVGLLAPPGGLSGRSDNH
(SEQ ID NO: 453)	(SEQ ID NO: 318)
PTPVCHHNFHCFGYD	ISSGLLSGGSGGLSGRSDNH
(SEQ ID NO: 454)	(SEQ ID NO: 346)
QATCQWKRSKCHNK	ISSGLLSGGSGGLSGRSDNH
(SEQ ID NO: 455)	(SEQ ID NO: 346)
QHSWCQHKAKCNYGN	AVGLLAPPGGTSTSGRSANPRG
(SEQ ID NO: 456)	(SEQ ID NO: 347)
QNCSPYTTTHKCKLT	TSTSGRSANPRGGGAVGLLAPP
(SEQ ID NO: 457)	(SEQ ID NO: 348)
QSSNCEHKRKCISIS	VHMPGLGFLPGGTSTSGRSANPRG
(SEQ ID NO: 458)	(SEQ ID NO: 349)
RPCLLGLVPDHKCKL	TSTSGRSANPRGGGVHMPGLGFLGP
(SEQ ID NO: 459)	(SEQ ID NO: 350)
RRSCMRSINTCKQKY	LSGRSDNHGGAVGLLAPP
(SEQ ID NO: 460)	(SEQ ID NO: 351)
RSSCPTVTPQNCENQ	VHMPGLGFLPGGLSGRSDNH
(SEQ ID NO: 461)	(SEQ ID NO: 352)
RTMCLDLNHHKCKPSN	LSGRSDNHGGGVHMPGLGFLGP
(SEQ ID NO: 462)	(SEQ ID NO: 353)

RTYWCNTNHNKCKHNM (SEQ ID NO: 463)	LSGRSDNHGGSGGSISSGLLSS (SEQ ID NO: 354)
RVENCEHNQYCHKWK (SEQ ID NO: 464)	LSGRSGNHGGSGGSISSGLLSS (SEQ ID NO: 355)
SCHEDDHKNKNICSL (SEQ ID NO: 465)	ISSGILLSGGSGGSLSGRSGNH (SEQ ID NO: 356)
SDTCVMNHPKCKRDN (SEQ ID NO: 466)	LSGRSDNHGGSGGSQLQALRMA (SEQ ID NO: 357)
SSTCFHPNQKECMTK (SEQ ID NO: 467)	QNALRMAGGSGGSLSGRSDNH (SEQ ID NO: 358)
SSYCGGITMRCRRAM (SEQ ID NO: 468)	LSGRSGNHGGSGGSQLQALRMA (SEQ ID NO: 359)
STGYCTYVNWNCYTN (SEQ ID NO: 469)	QNALRMAGGSGGSLSGRSGNH (SEQ ID NO: 360)
THKCKLHLQVCTQTT (SEQ ID NO: 470)	ISSGILLSGRSGNH (SEQ ID NO: 361)
TMNCTHPKQKCQHTN (SEQ ID NO: 471)	ISSGILLSGGSGGSLSGRSGNH (SEQ ID NO: 1091)
TNVLCESHNCDHKNK (SEQ ID NO: 472)	ISSGILLSGRSANPRG (SEQ ID NO: 1092)
TQHAASLGVEHKSKI (SEQ ID NO: 473)	AVGLLAPPTSGRSANPRG (SEQ ID NO: 1093)
TQLPCFDDHKCKNTN (SEQ ID NO: 474)	AVGLLAPPSGRSANPRG (SEQ ID NO: 1094)
TSDSCMRQKCEHKEK (SEQ ID NO: 475)	ISSGILLSGRSDDH (SEQ ID NO: 1095)
TTCDDDHKYKHKCAQL	ISSGILLSGRSDIH

(SEQ ID NO: 476)	(SEQ ID NO: 1096)
TTCGAHKEKQHCIYT	ISSGLLSGRSDQH
(SEQ ID NO: 477)	(SEQ ID NO: 1097)
TTYCAYWHNKCKFET	ISSGLLSGRSDTH
(SEQ ID NO: 478)	(SEQ ID NO: 1098)
VGPTCGHAKCKQSEV	ISSGLLSGRSDYH
(SEQ ID NO: 479)	(SEQ ID NO: 1099)
VSHPCNTHKCKTNIV	ISSGLLSGRSDNP
(SEQ ID NO: 480)	(SEQ ID NO: 1100)
WDCRNTSHPKLKCHN	ISSGLLSGRSANP
(SEQ ID NO: 481)	(SEQ ID NO: 1101)
WSPCNSDHRKRCNNG	ISSGLLSGRSANT
(SEQ ID NO: 482)	(SEQ ID NO: 1102)
YANMSCEYDCHKMKY	AVGLLAPPGGLSGRSDDH
(SEQ ID NO: 483)	(SEQ ID NO: 1103)
YANPCTHKEKCHFN	AVGLLAPPGGLSGRSDIH
(SEQ ID NO: 484)	(SEQ ID NO: 1104)
YDCSPSWTHPKCKHK	AVGLLAPPGGLSGRSDQH
(SEQ ID NO: 485)	(SEQ ID NO: 1105)
YGTCTTTHPKCKTTN	AVGLLAPPGGLSGRSDTH
(SEQ ID NO: 486)	(SEQ ID NO: 1106)
YQKCHPKAKDCGNNT	AVGLLAPPGGLSGRSDYH
(SEQ ID NO: 487)	(SEQ ID NO: 1107)
YWECPNMEHNKCKNN	AVGLLAPPGGLSGRSDNP
(SEQ ID NO: 488)	(SEQ ID NO: 1108)
PMGNRYCVLDHPKLIK	AVGLLAPPGGLSGRSANP
(SEQ ID NO: 489)	(SEQ ID NO: 1109)

GHKSCCQKHCEYTQT (SEQ ID NO: 490)	AVGLLAPPGGLSGRSANI (SEQ ID NO: 1110)
LYLEMCSCCCWESIT (SEQ ID NO: 491)	ISSGLLSGRSDNI (SEQ ID NO: 1111)
ACQAQHCYKTYACKP (SEQ ID NO: 492)	AVGLLAPPGGLSGRSDNI (SEQ ID NO: 1112)
CCYTCSVHPKCKNQL (SEQ ID NO: 493)	
CKHRCSHKKEKCPANH (SEQ ID NO: 494)	
CHVLFCLMQCCRWSL (SEQ ID NO: 495)	
LNSSLVFDHPKAKPN (SEQ ID NO: 496)	
MCLLCRSKFGCKVKG (SEQ ID NO: 497)	
IICNDHKCKQNQCNN (SEQ ID NO: 498)	
IRCSLRDSLGCGERM (SEQ ID NO: 499)	
TSCQPPKHKCTCNHG (SEQ ID NO: 500)	
TQCPHRCVKPNCWLH (SEQ ID NO: 501)	
KCCETKRNHHKCTYK (SEQ ID NO: 502)	
LPHCCHKAKHCNHTS	

(SEQ ID NO: 503)
PAMCAAIHEKCCIKV
(SEQ ID NO: 504)
PRSCGNQLCPCHYYK
(SEQ ID NO: 505)
TNKCSCNHNMKCINY
(SEQ ID NO: 506)
VETCCQHNKCKYPFI
(SEQ ID NO: 507)
IFCCSNHEDHKCKTN
(SEQ ID NO: 508)
VCRLLICPLTCVIGVG
(SEQ ID NO: 509)
FHGCCSVYSCLTNPG
(SEQ ID NO: 510)
ALACHPKQKPLEGQL
(SEQ ID NO: 511)
SIICCATSSCPLKHL
(SEQ ID NO: 512)
APCCRPHEKPIDSR
(SEQ ID NO: 513)
WELCCPSADCRVAMG
(SEQ ID NO: 514)
QDHPKTKWNS (SEQ
ID NO: 548)
ACRICQDHPATKWNS
(SEQ ID NO: 549)

ACRICQDHPKTAWNS
(SEQ ID NO: 550)
ACRICQDAPKTKWNS
(SEQ ID NO: 551)
ACRICQDHAKTKWNS
(SEQ ID NO: 552)
DHPATKWNS
(SEQ ID NO: 553)
DHPKTAWNS
(SEQ ID NO: 554)
DAPKTKWNS
(SEQ ID NO: 555)
DAPATKWNS
(SEQ ID NO: 556)
ACRICQDHP
(SEQ ID NO: 557)
HPQSKDTL
(SEQ ID NO: 558)
HPKSQDTL
(SEQ ID NO: 559)
TIPRCGQHPLCLDTL
(SEQ ID NO: 560)
HPLSLDTL
(SEQ ID NO: 561)
HPASKDTL
(SEQ ID NO: 562)
HPKSADTL

(SEQ ID NO: 563) PGNCHPLQKELCQFI (SEQ ID NO: 564) HPLQKELAQFI (SEQ ID NO: 565) HPLALELAQFI (SEQ ID NO: 566) PGNCHPLQLELCQFI (SEQ ID NO: 567) TNIWSCQTYCDHHAHA (SEQ ID NO: 568) TNIWSCQTYCDHAHL (SEQ ID NO: 569) TNIWSCQTYCDHLHA (SEQ ID NO: 570) TNIWSCQTYCDHKHA (SEQ ID NO: 571)					
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[000853] Any of the combinations described in Tables 13A, 13B, 13C, and/or 13D can be combined with human immunoglobulin constant regions to result in fully humanized IgGs including IgG1, IgG2, IgG4 or mutated constant regions to result in human IgGs with altered functions such as IgG1 N297A, IgG1 N297Q, or IgG4 S228P. Additional examples are known to those skilled in the art. Examples of Ig heavy chain constant region amino acids in which mutations in at least one amino acid leads to reduced Fc function include, but are not limited to, mutations in amino acid 228, 233, 234, 235, 236, 237, 239, 252, 254, 256, 265, 270, 297, 318, 320, 322, 327, 329, 330, and 331 of the heavy constant region. Examples of combinations of mutated amino acids are also known in the art, such as, but not limited to a combination of mutations in amino acids 234, 235, and 331, such as L234F, L235E, and P331S or a combination of amino acids 318, 320, and 322, such as E318A, K320A, and K322A.

[000854] As an example a masking moiety comprising SEQ ID NO: 66 can be combined with a substrate comprising SEQ ID NO: 294, and a VL region comprising SEQ ID NO: 39 in combination with a human kappa constant domain comprising SEQ ID NO: 61 to produce a light chain comprising SEQ ID NO: 380. In some embodiments, for example, a VH region comprising SEQ ID NO: 21 can be combined with human immunoglobulin heavy chain constant domain to give an IgG1 heavy chain (SEQ ID NO: 2048), an IgG4 heavy chain (SEQ ID NO: 2051), an IgG4 S228P heavy chain (SEQ ID NO: 2047), a mutated IgG1 N297A heavy chain (SEQ ID NO: 2049) or a mutated IgG1 N297Q heavy chain (SEQ ID NO: 2050). Co-expression of SEQ ID NO: 380 with SEQ ID NO: 2048 will yield an IgG1 anti-PD-1 activatable antibody. Co-expression of SEQ ID NO: 380 with SEQ ID NO: 2051 will yield an IgG4 anti-PD-1 activatable antibody. Co-expression of SEQ ID NO: 380 with SEQ ID NO: 2047 will yield an IgG4S228P anti-PD-1 activatable antibody. Co-expression of SEQ ID NO: 380 with SEQ ID NO: 2049 will yield an IgG1 N297A anti-PD-1 activatable antibody. Co-expression of SEQ ID NO: 380 with SEQ ID NO: 2050 will yield an IgG1 N297Q anti-PD-1 activatable antibody.

[000855] In some embodiments, a masking moiety comprising SEQ ID NO: 99 is combined with a substrate comprising SEQ ID NO: 214 and a VL region comprising SEQ ID NO: 47 in combination with a human kappa constant domain comprising SEQ ID NO: 61 to produce a light chain comprising SEQ ID NO: 2055. In some embodiments, spacer QGQSGQG (SEQ ID NO: 362) is added to the N-terminus of SEQ ID NO: 2055 to form SEQ ID NO: 2054. In some embodiments, a masking moiety comprising SEQ ID NO: 99 is combined with a substrate

comprising SEQ ID NO: 1100 and a VL region comprising SEQ ID NO: 47 in combination with a human kappa constant domain comprising SEQ ID NO: 61 to produce a light chain comprising SEQ ID NO: 2057. In some embodiments, spacer QGQSGQG (SEQ ID NO: 362) is added to the N-terminus of SEQ ID NO: 2057 to form SEQ ID NO: 2056. In some embodiments, a masking moiety comprising SEQ ID NO: 99 is combined with a substrate comprising SEQ ID NO: 1101, and a VL region comprising SEQ ID NO: 47 in combination with a human kappa constant domain comprising SEQ ID NO: 61 to produce a light chain comprising SEQ ID NO: 2059. In some embodiments, spacer QGQSGQG (SEQ ID NO: 362) is added to the N-terminus of SEQ ID NO: 2059 to form SEQ ID NO: 2058. In some embodiments, a VH region comprising SEQ ID NO: 21 can be combined with an IgG4 S228P human immunoglobulin heavy chain constant domain comprising SEQ ID NO: 63 to produce a heavy chain comprising SEQ ID NO: 2052. In some embodiments, the C-terminal lysine of SEQ ID NO: 2052 is missing to form an amino acid sequence having SEQ ID NO: 2053. Co-expression of any one of these light chains with any one of these heavy chains will lead to production of an activatable antibody of the embodiments.

[000856] Constant region amino acid sequences are shown below in SEQ ID NO: 381, 382, 383, and 1807.

IgG1 Hc CH1-end amino acid sequence:

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP
KPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY**N**STYRVVSVLTVL
HQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFY
PSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVVFSCVMHEALHNHYT
QKSLSLSPG (SEQ ID NO: 381)

IgG1NA CH1-end amino acid sequence:

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP
KPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY**A**STYRVVSVLTVL
HQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFY

PSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPG (SEQ ID NO: 382)

IgG1NQ CH1-end amino acid sequence:

ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP
KPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY[Q]STYRVVSVLTVL
HQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFY
PSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
QKSLSLSPG (SEQ ID NO: 1807)

IgG4 Hc CH1-end amino acid sequence:

ASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVVTVPSSSLGTQTYTCNVDHKPSNTKVDKRVESKYGP[PCP]CPAPEFLGGPSVFLFPPKPK
DTLMISRTPEVTCVVDVVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQD
WLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSD
IAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKS
LSLSLG (SEQ ID NO: 383)

Light Chain (Lc) amino acid sequence:

AMSGCSWSAFCPYLA[X1]LSGRSDNH[X2]DIQLTQSPSSLSASVGDRVITITCRASESVDNYG
ISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISMQPEDFATYYCQQSKD
VPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNFFYPREAKVQWKVDNALQSG
NSQESVTEQDSKDSTYLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ
ID NO: 380)

where each of X1 and X2 independently can be a linking peptide or absent.

IgG4 S228P Heavy Chain (Hc) amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSV
KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGLTVTVSSASTKGPSVF

PLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPS
 SSLGTKTYTCNVDHKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVDSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKSLSLSPG
 (SEQ ID NO: 2047)

IgG1 Heavy Chain (Hc) amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLVTVSS
 ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYNSTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG (SEQ ID NO: 2048)

IgG1NA Hc amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLVTVSS
 ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYASTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG (SEQ ID NO: 2049)

IgG1NQ Hc amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLVTVSS
 ASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPPKPKDTLMISRTPEVTCVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQYASTYRVVSVLTVLHQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYTQKSLSLSPG

LSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSCDKTHTCPPCPAPELLGGPSVFLFPP
 KPKDTLMISRTPEVTCVVVDVSHEDPEVKFNWYVDGVEVHNAKTKPREEQY[Q]STYRVVSVLTVL
 HQDWLNGKEYKCKVSNKALPAPIEKTISKAKGQPREPQVYTLPPSREEMTKNQVSLTCLVKGFY
 PSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSKLTVDKSRWQQGNVFSCSVMHEALHNHYT
 QKSLSLSPG (SEQ ID NO: 2050)

IgG4 Hc amino acid sequence:

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNAHYADSV
 KGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLLTVSS
 ASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
 LSSVVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGP[PCP]CPAPEFLGGPSVFLFPPKPK
 DTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQD
 WLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSD
 IAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMHEALHNHYTQKS
 LSLSLG (SEQ ID NO: 2051)

Heavy Chain 1 Amino Acid Sequence (SEQ ID NO: 2052):

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNA
 HYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLLT
 VSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVL
 QSSGLYSLSSVVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGP[PCP]CPAPEFLGG
 PSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQF
 NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQ
 EEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDK
 SRWQEGNVFSCSVMHEALHNHYTQKSLSLSLGK

Heavy Chain 2 Amino Acid Sequence (SEQ ID NO: 2053):

EVQLVESGGGLVQPGGSLRLSCAASGFTFSGYAMSWVRQAPGKGLEWVAYISNSGGNA
 HYADSVKGRFTISRDN SKNTLYLQMNSLRAEDTAVYYCTREDYGTSPFVYWGQGTLLT
 VSSASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVL
 QSSGLYSLSSVVTVPSSSLGKTYTCNV DHKPSNTKVDKRVESKYGP[PCP]CPAPEFLGG

PSVFLFPPKPKDTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQF
NSTYRVVSVLTVLHQDWLNGKEYKCKVSNKGLPSSIEKTISKAKGQPREPQVYTLPPSQ
EEMTKNQVSLTCLVKGFYPSDIAVEWESNGQPENNYKTTPPVLDSDGSFFLYSRLTVDK
SRWQEGNVFSCSVMHEALHNHYTQKSLSLGLG

Light Chain 1 Amino Acid Sequence with spacer (SEQ ID NO: 2054):

[QGQSGQG][TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLS
ASVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGS
GTDFTLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKS
GTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSSSTLTLSKAD
YEKHKVYACEVTHQGLSSPVTKSFNRGEC]

Light Chain 1 Amino Acid Sequence without spacer (SEQ ID NO: 2055):

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVIT
ITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISS
MQPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLL
NNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSSSTLTLSKADYEKHKVYAC
EVTHQGLSSPVTKSFNRGEC

Light Chain 2 Amino Acid Sequence with spacer (SEQ ID NO: 2056):

[QGQSGQG][TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNPGGGSDIQLTQSPSSLS
ASVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGS
GTDFTLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKS
GTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSSSTLTLSKAD
YEKHKVYACEVTHQGLSSPVTKSFNRGEC]

Light Chain 2 Amino Acid Sequence with spacer (SEQ ID NO: 2057):

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNPGGGSDIQLTQSPSSLSASVGDRVIT
ITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISS
MQPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN

FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEV
THQGLSSPVTKSFNRGEC

Light Chain 3 Amino Acid Sequence with spacer (SEQ ID NO: 2058):

[QGQSGQG][TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLS
ASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGGSGVPSRFSGSGS
GTDFTLTISMQPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKS
GTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKAD
YEKHKVYACEVTHQGLSSPVTKSFNRGEC]

Light Chain 3 Amino Acid Sequence with spacer (SEQ ID NO: 2059):

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLSASVGDRVTIT
CRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGGSGVPSRFSGSGSGTDFTLTISMQ
QPEDFATYYCQQSKDVPWTFGQGTKLEIKRTVAAPSVFIFPPSDEQLKSGTASVVCLLNN
FYPREAKVQWKVDNALQSGNSQESVTEQDSKDYSLSTLTLSKADYEKHKVYACEV
THQGLSSPVTKSFNRGEC

EXAMPLE 10: Evaluation of efficiency of masking moieties

[000857] This example describes activatable anti-PD-1 antibodies of the disclosure with reduced binding to hPD-1.

[000858] Masking the ability of an antibody to bind to its antigen is an example of inhibition of binding. The extent of inhibition is dependent on the affinity of the antibody for its antigen, the affinity of the masking moiety (also referred to herein as the mask) for the antibody and the concentration of all reactants. Local concentrations of the tethered peptide mask (inhibitor) is very high in the activatable antibody context, on the order of 10 mM, therefore moderate affinity peptides would effectively mask activatable antibody antigen binding.

[000859] Masking efficiencies were evaluated by standard plate ELISA. Briefly, human PD-1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified anti-PD-1 antibodies and activatable antibodies were applied to the plate in serial dilution and allowed to bind. Bound antibody and activatable antibodies were detected with anti-human IgG-HRP conjugate, FAb-specific (Sigma, St Louis, MO) and visualized with the

chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Shown are plots of binding isotherms for anti-PD-1 A1.4 and anti-PD-1 A1.5 activatable antibodies (Figures 14-18). Plots were generated in Prizm (Sigma Plot); the data were fit to a model of single site saturation binding, and the equilibrium dissociation constant, K_d , was determined. Masking efficiency was calculated by dividing the K_d for binding of the activatable antibodies by the K_d of the parental antibody. The masking efficiency (ME) values for the tested antibodies are shown in Table 14 and Table 15 below (masking moiety PD001 is also referred to herein as PD01 and/or PD-01; masking moiety PD002 is also referred to herein as PD02 and/or PD-02, and so on), and the apparent K_d (nM) and masking efficiency (ME) values for the tested activatable antibodies are shown in Table 26:

Table 14: Masking efficiency values for activatable antibodies tested in Figure 14

Mask	ME
PD01	25
PD02	18
PD03	5
PD08	13
PD09	2
PD-10	13

Table 15: Masking efficiency values for activatable antibodies tested in Figure 18

Mask	ME
None	1
PD06	30
PD09	2
PD-12	10
PD-17	20
PD-19	30
PD25	30
PD26	20

PD28	25
PD34	42

Table 26: Apparent K_d and Masking efficiency values for activatable antibodies tested in Figure 15

Molecule	Apparent K _D , nM	Masking Efficiency
A1.5 (n=5 avg)	1.8	
PD01	348	193
PD02	21	12
PD03	109	61
PD05	68	38
PD06	71	39
PD08	55	31
PD09	10	6
PD10	54	30
PD11	67	37
PD12	25	14
PD13	151	84
PD14	100	56
PD15	117	65
PD16	57	32
PD17	22	12
PD18	75	42
PD19	93	52
PD20	77	43
PD21	49	27
PD22	41	23
PD23	31	17
PD24	92	51

EXAMPLE 11: Anti-PD-1 activatable antibodies of the embodiments are functionally masked in a human T-cell restimulation assay

[000860] This example describes the effect of masking moieties on the biological function of the anti-PD-1 antibody. (Figure 19)

[000861] PBMCs from a CMV-positive donor (Hemacare) were plated at 2×10^5 cells/well in the presence of CMV viral lysate (Astarte) and either anti-PD-1 antibody A1.5, an anti-PD-1 activatable antibody of the disclosure, or hIgG4 isotype control antibody. After four days, supernatant was removed from each well and IFN-gamma levels were assayed using IFN-gamma ELISA kit (Life Technologies, Carlsbad, CA). Anti-PD-1 A1.5 activatable antibodies increased CMV-stimulated IFN-gamma secretion compared with control hIgG4 but with decreased potency relative to anti-PD-1 A1.5 parental antibody (Figure 19).

EXAMPLE 12: Anti mouse PD-1 J43v2 mIgG2a antibody binds mouse PD-1 and blocks mouse PD-L1 and mouse PD-L2 binding

[000862] This example demonstrates that Armenian hamster anti mouse PD-1 J43 antibody (US 7,858,746; Agata et al, 1996, International Immunology, Vol. 8 No. 5 pp765-77) can be functionally expressed as a mouse IgG2a antibody.

[000863] Anti-mouse PD-1 antibody J43 was reformatted as a hamster LC (SEQ ID NO: 543 amino acid; SEQ ID NO: 544 or SEQ ID NO: 545 nucleotide), mouse IgG2a antibody by fusion of the heavy chain variable domain to mIgG2a, resulting in amino acid SEQ ID NO 546, nucleotide sequence SEQ ID NO 547. Full length J43v2 mIgG2a (also referred to as J43v2, J43 m2a and J43v2 m2a) was expressed by transient transfection of HEK293 cells and purified from the culture supernatant by Protein G chromatography. One embodiment of this disclosure is an antibody comprising a light chain comprising amino acid sequence SEQ ID NO: 543. One embodiment of this disclosure is an antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 546. One embodiment of this disclosure is an antibody comprising a light chain comprising amino acid sequence SEQ ID NO: 543 and a heavy chain comprising amino acid sequence SEQ ID NO: 546. One embodiment of this disclosure is an antibody comprising the CDRs of an antibody comprising a light chain comprising amino acid sequence SEQ ID NO: 543 and a heavy chain comprising amino acid sequence SEQ ID NO: 546.

[000864] Functionality was assayed by binding of J43v2 m2a to HEK293 cells expressing mouse PD-1 and confirmation that binding of J43v2 m2a to HEK293-mPD-1 inhibited binding of biotinylated mPD-L1 and biotinylated mPD-L2 to HEK293-mPD-1 (Figure 20). About 100,000 mPD-1 HEK293 cells were transferred to a U-bottom 96-well plate. For the binding experiment, a 1:4 titration of J43v2 m2a antibody starting at 150 nM was added to cells. For blocking experiments, 20nM of biotinylated-mPD-L1-Fc or 20 nM of biotinylated-mPD-L2-Fc were premixed with a 1:4 titration of J43v2 m2a starting at 150 nM and added to cells. Both were incubated for 1 hour on ice and cells were washed 3 times. J43v2 m2a antibody binding was detected using an anti-mouse secondary antibody (Jackson ImmunoResearch, West Grove, PA). For the blocking experiment, biotinylated-PD-L1 or biotinylated-PD-L2 was detected using streptavidin-PE (Life Technologies, Carlsbad, CA). Both binding and blocking were incubated for 30 minutes on ice and were washed and read on a flow cytometer (MACSQuant).

EXAMPLE 13: J43v2 m2a induces diabetes in NOD mouse model

[000865] In this Example, anti-PD-1 J43v2 m2a antibody was confirmed to induce diabetes in the NOD mouse model.

[000866] Antibody J43v2 m2a was confirmed to induce diabetes in NOD mice as follows. The NOD mice, substrain NOD/ShiLtJ were obtained from Jackson Laboratory at 8 weeks and acclimated on site for 2 weeks. At 10 weeks, mice were checked for diabetes prior to enrollment, grouped, and dosed as set forth in Table 16. Figure 21 shows the dose-dependent induction of diabetes in NOD mice by J43v2 m2a. At day eight after a single dose, 100% of mice in the mIgG2a isotype control group remained non-diabetic, while 0%, 14% and 86% of the 20mg/kg, 3 mg/kg and 1mg/kg groups, respectively, remained non-diabetic.

Table 16: Study design

Group	Count	Gender	Treatment	Dose (mg/kg)	Dose volume (mL/kg)	Schedule	Route
1	7	F	mIgG2a (C1.18.4)	20	10	q7dx1	IP
2	7	F	Anti-PD-1 (J43v2)	20	10	q7dx1	IP
3	7	F	Anti-PD-1 (J43v2)	5	10	q7dx1	IP
4	7	F	Anti-PD-1 (J43v2)	1	10	q7dx1	IP

EXAMPLE 14: Activatable anti-mouse PD-1 J43 masking moieties

[000867] This example describes identification of masking moieties (MM) that reduce binding of anti-PD-1 J43 antibody to its target.

[000868] Anti-PD-1 J43v2 mIgG2a was used to screen libraries using a method similar to that described in PCT International Publication Number WO 2010/081173, published 15 July 2010. The screening consisted of one round of MACS and three rounds of FACS sorting. For the initial MACS, approximately 1×10^{12} cells were incubated with J43v2 m2a antibody at a concentration of 200nM, and approximately 9×10^7 binders were collected using Protein-G Dynabeads (Life Technologies, Carlsbad, CA). FACS rounds were conducted labeling cells with DyLight 650 (Thermo-Fisher) labeled J43v2 m2a antibody and collecting cells with strongest fluorescence as follows: 200nM J43-650 collecting brightest 3% for FACS round 1 (F1), 20nM J43-650 collecting brightest 8% for FACS round 2 (F2) and 1nM J43-650 collecting two populations, the brightest 20% and the brightest 0.5% for FACS round 3 (F3-20 and F3-0.5). Individual peptide clones from F2, F3-20 and F3-0.5 were identified by sequence analysis (Table 17), and selected peptide clones MP001-MP014 were subsequently verified for their ability to bind J43-650 (masking moiety MP001 is also referred to herein as MP01 and/or MP-01; masking moiety MP002 is also referred to herein as MP02 and/or MP-02, and so on).

Table 17: Masking moiety sequences

Mask	Amino Acid Sequence	SEQ ID NO:
MP001	DYTYCRWVNWCLSGV	384
MP002	ILCPEDPWGHKCKLP	385
MP003	TNIWSCQTYCDHKHK	386
MP004	SDHKCKLQNCMNTKV	387
MP005	PGNCHPMQKEMCQFI	388
MP006	VEHLCYTHNKCKHPD	389
MP007	TIPRCGQHPKCKDTL	390
MP008	ACRICQDHPKTKWNS	391
MP009	LIQCTGNLDHKCKHY	392
MP010	IPCHHSADHKHKCTS	393

MP011	SRQICADYNCHNKYK	394
MP012	QPCNPQIDHKIKCIY	395
MP013	HYTICMTHNKCKDMA	396
MP014	ANSCLAVEHKCKHNY	397
MP015	AALHCTEHKCKNHIK	398
MP016	APCIINTVDWKSCEI	399
MP017	ATNWCTHKQKCKQDM	400
MP018	DCYNEHKLKTRVCNN	401
MP019	DEMQCSHKQKCTNSK	402
MP020	DVGICSQHNKCKPTK	403
MP021	EKYCSSDDHKCKITL	404
MP022	ELECSHNKVKNCIQI	405
MP023	ELHPCNTHKCKPIVN	406
MP024	EVGSCNHPKCKSNNY	407
MP025	EYSPSLAHPKLKDNA	408
MP026	FESLHPKGKHPEDLG	409
MP027	FPLCVRADRVCGDAQ	410
MP028	FQAPPASHNKLKPSL	411
MP029	GAIDSCHHKCKSPHY	412
MP030	GKIYTCEHNCTFGYS	413
MP031	HCTVNNHSSDHKCKI	414
MP032	HGTQCTHNKCKPILS	415
MP033	HIGWCLHPKCKTTTT	416
MP034	HLRTCIRQKWCEHKMK	417
MP035	HTDCTMMSNHKCKIN	418
MP036	IRQQCTALACLLKVH	419
MP037	KGCSTHKMRAYCNQM	420
MP038	KMFTPCKIWCNNSYN	421
MP039	KTMCSGHKQKCNSS	422
MP040	LACHSASLVDHKCKL	423

MP041	LCNVSMDHKHKPCYL	424
MP042	LGLNCFSEHKCKEHM	425
MP043	LGTCTHKHKNCNYTL	426
MP044	LHEGCTTHNKCKPIA	427
MP045	LKRSC TGHWTCTYNW	428
MP046	LQRCTHKEKYCHAIH	429
MP047	LSHCYDHKRKCSYIV	430
MP048	LSKCHNKEKNCSNNN	431
MP049	MDTCEMHKQKCRPSF	432
MP050	MHNECLTHKCKVPIT	433
MP051	MLTLCNTNACHKEKN	434
MP052	MRPCLNNLEHKCKHY	435
MP053	MSRCPTHKMKCSLNI	436
MP054	MWICQEHKLKCM TDT	437
MP055	MYECKRRSAFYCTLN	438
MP056	NDCQHDKQMHKCKMH	439
MP057	NFGPCPMLLGCFGR	440
MP058	NHTDCSHPKCKSHDS	441
MP059	NLNCPHKQKNCDKYH	442
MP060	NPQCTPIDHKCKTHH	443
MP061	NTTSCTHPKCKHQGK	444
MP062	NVGGCDNYGCHKLKN	445
MP063	PCSPGNLTWDHKCKY	446
MP064	PFTKCHGFNKCKEHT	447
MP065	PGDKCTHKEKCYNN	448
MP066	PNICNLDHKRKCRIN	449
MP067	PQLACKHPKCKDAGN	450
MP068	PSCTMWTHGGVCKHA	451
MP069	PSHRHPLAKPGFRVE	452
MP070	PTCFKTHNKSCKNRV	453

MP071	PTPVCHHNFHCFGYD	454
MP072	QATCQWKKRSKCHNK	455
MP073	QHSWCQHKAKCNYGN	456
MP074	QNCSPTYTTHKCKLT	457
MP075	QSSNCEHKRKCISIS	458
MP076	RPCLLGLVPDHKCKL	459
MP077	RRSCMRSINTCKQKY	460
MP078	RSSCPTVTPQNCENQ	461
MP079	RTMCLDLNHKCKPSN	462
MP080	RTYWCTNHNKCKHNM	463
MP081	RVENCEHNQYCHKWK	464
MP082	SCHEDDHKNKNICSL	465
MP083	SDTCVMNHPKCKRDN	466
MP084	SSTCFHPNQKECMTK	467
MP085	SSYCGGITMRCRRAM	468
MP086	STGYCTYVNWNYTN	469
MP087	THKCKLHLQVCTQTT	470
MP088	TMNCTHPKQKCQHTN	471
MP089	TNVLCESHNCDHKNK	472
MP090	TQHAASLGVEHKSki	473
MP091	TQLPCFDDHKCKNTN	474
MP092	TSDSCMRQKCEHKEK	475
MP093	TTCDDHKYKHKCAQL	476
MP094	TTCGAHKEKQHCIYT	477
MP095	TTYCAYWHNKCKFET	478
MP096	VGPTCGHAKCKQSEV	479
MP097	VSHPCNTHKCKTNIV	480
MP098	WDCRNTSHPKLKCHN	481
MP099	WSPCNSDHKRKCNG	482
MP100	YANMSCEYDCHKMKY	483

MP101	YANPCTHKEKCHFKN	484
MP102	YDCSPSWTHPKCKHK	485
MP103	YGWTCTTHPKCKTTN	486
MP104	YQKCHPKAKDCGNNT	487
MP105	YWECPNMEHNKCKNN	488
MP106	PMGNRYCVLDHPKLLK	489
MP107	GHKSCCQKHCEYTQT	490
MP108	LYLEMCSCCCWESIT	491
MP109	ACQAQHCYKTYACKP	492
MP110	CCYTCSVHPKCKNQL	493
MP111	CKHRCSHKEKCPANH	494
MP112	CHVLFCLMQCCRWSL	495
MP113	LNSSLVFDHPKAKPN	496
MP114	MCLLCRSKFGCKVKG	497
MP115	IICNDHKCKQNQCNN	498
MP116	IRCSLRDSLGCERM	499
MP117	TSCQPPKHKCTCNHG	500
MP118	TQCPHRCVKPNCWLH	501
MP119	KCCETKRNNHKHCTYK	502
MP120	LPHCCHKAKHCNHTS	503
MP121	PAMCAAIHEKCCIKV	504
MP122	PRSCGNQLCPCHYYK	505
MP123	TNKCSCNHNMKCINY	506
MP124	VETCCQHNKCKYPFI	507
MP125	IFCCSNHEDHKCKTN	508
MP126	VCRLICPLTCVIGVG	509
MP127	FHGCCSVYSCLTNPG	510
MP128	ALACHPKQKPLEGQL	511
MP129	SIICCATSSCPLKHL	512
MP130	APCCRPHKEKPIDSR	513

MP131	WELCCPSADCRVAMG	514
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EXAMPLE 15: Characterization of activatable anti-mouse PD-1 antibodies

[000869] This example describes activatable anti-PD-1 antibodies of the disclosure with reduced binding to hPD-1 compared to binding by parental antibody.

[000870] Masking efficiencies were evaluated by standard plate ELISA. Briefly, mouse PD-1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified J43v2 and activatable J43v2 antibodies comprising the indicated masks, cleavable moiety 2001, and antibody J43v2 were applied to the plate in serial dilution and allowed to bind. Bound antibody and activatable antibodies were detected with anti-mouse IgG-HRP conjugate (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prizm (Sigma Plot). All activatable J43 antibodies showed strongly decreased binding compared with the parental J43 (Figure 22).

[000871] In some embodiments, the activatable antibody also includes a spacer sequence. In some embodiments, the spacer is joined directly to the MM of the activatable antibody. In some embodiments, the spacer is joined directly to the MM of the activatable antibody in the structural arrangement from N-terminus to C-terminus of spacer-MM-CM-AB. In some embodiments, the spacer joined directly to the N-terminus of MM of the activatable antibody is selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. In some embodiments, the spacer includes at least the amino acid sequence QGQSGQG (SEQ ID NO: 362). In some embodiments, the spacer includes at least the amino acid sequence QGQSGQ (SEQ ID NO: 913). In some embodiments, the spacer includes at least the amino acid sequence QGQSG (SEQ ID NO: 914). In some embodiments, the spacer includes at least the amino acid sequence QGQS (SEQ ID NO: 915). In some embodiments, the spacer includes at least the amino acid sequence QGQ (SEQ ID NO: 916). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 917). In some embodiments, the spacer includes at least the amino acid residue Q. In some embodiments, the activatable antibody does not include a spacer sequence.

[000872] Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG(SEQ ID NO: 2045), QG(SEQ ID NO: 2046), and G.

[000873] While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art appreciate that the activatable anti-PD-1 antibodies of the disclosure can include any suitable spacer sequence, such as, for example, a spacer sequence selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. . Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG(SEQ ID NO: 2045), QG(SEQ ID NO: 2046), and G. While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art will also appreciate that activatable anti-PD-1 antibodies of the disclosure in some embodiments do not include a spacer sequence.

Activatable Anti-mouse PD-1 light chain variable domains

[spacer (SEQ ID NO: 362)] [J43 MP001 2001 (SEQ ID NO: 999)] amino acid sequence:

[QGQSGQG] [DYTYCRWVNWCLSGVGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVL] (SEQ ID NO: 515)

J43 MP001 2001 amino acid sequence:

DYTYCRWVNWCLSGVGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVL (SEQ ID NO: 1077)

[spacer (SEQ ID NO: 918)] [J43 MP001 2001 (SEQ ID NO: 1000)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GATTATACGTATTGCCGTTGGGTTAATTGGTGCTTGTCTGGGTGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAA TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACTGTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT

CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
 AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
 GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
 GCGGAACCCAGCTCACCGTCCT (SEQ ID NO: 516)

[spacer (SEQ ID NO: 362)] [J43 MP002 2001 (SEQ ID NO: 1001)] amino acid sequence:

[QGQSGQG] [ILCPEDPWGHKCKLPGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 517)

J43 MP002 2001 amino acid sequence:

ILCPEDPWGHKCKLPGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
 QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
 GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1078)

[spacer (SEQ ID NO: 918)] [J43 MP002 2001 (SEQ ID NO: 1002)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [ATTCTGTGCCCCGAGGATCCGTGGGGGCATAAGTGCAAGC
 TGCTTGAGAGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
 GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
 CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
 AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
 GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
 GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 518)

[spacer (SEQ ID NO: 362)] [J43 MP003 2001 (SEQ ID NO: 1003)] amino acid sequence:

[QGQSGQG] [TNIWSCQTYCDHKHKGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 519)

J43 MP003 2001 amino acid sequence:

TNIWSCQTYCDHKHKGGGSSSGSISGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1079)

[spacer (SEQ ID NO: 918)] [J43 MP003 2001 (SEQ ID NO: 1004)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [ACGAATATTTGGAGTTGCCAGACTTATTGCGATCATAAGC
ATAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 520)

[spacer (SEQ ID NO: 362)] [J43 MP004 2001 (SEQ ID NO: 1005)] amino acid sequence:

[QGQSGQG] [SDHKCKLQNCMNTKVGGGSSSGSISGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 521)

J43 MP004 2001 amino acid sequence:

SDHKCKLQNCMNKTVGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1080)

[spacer (SEQ ID NO: 918)] [J43 MP004 2001 (SEQ ID NO: 1006)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [AGTGATCATAAGTGCAAGCTTCAGAATTGCATGAATACTA
AGGTTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 522)

[spacer (SEQ ID NO: 362)] [J43 MP005 2001 (SEQ ID NO: 1007)] amino acid sequence:

[QGQSGQG] [PGNCHPMQKEMCQFIGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 523)

J43 MP005 2001 amino acid sequence:

PGNCHPMQKEMCQFIGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1081)

[spacer (SEQ ID NO: 918)] [J43 MP005 2001 (SEQ ID NO: 1008)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CCTGGTAATTGCCATCCTATGCAGAAGGAGATGTGCCAGT
TTATTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG

AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
 GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
 GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 524)

[spacer (SEQ ID NO: 362)] [J43 MP006 2001 (SEQ ID NO: 1009)] amino acid sequence:

[QGQSGQG] [VEHLCYTHNKCKHPDGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 525)

J43 MP006 2001 amino acid sequence:

VEHLCYTHNKCKHPDGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
 QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
 GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1082)

[spacer (SEQ ID NO: 918)] [J43 MP006 2001 (SEQ ID NO: 1010)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GTTGAGCATTTGTGCTATACGCATAATAAGTGCAAGCATC
 CTGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
 GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
 CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
 AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
 GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
 GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 526)

[spacer (SEQ ID NO: 362)] [J43 MP007 2001 (SEQ ID NO: 1011)] amino acid sequence:

[QGQSGQG] [TIPRCGQHPKCKDTLGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 527)

J43 MP007 2001 amino acid sequence:

TIPRCGQHPKCKDTLGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSDDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1083)

[spacer (SEQ ID NO: 918)] [J43 MP007 2001 (SEQ ID NO: 1012)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [ACTATTCCGAGGTGCGGTCAGCATCCGAAGTGCAAGGATA
CTTTGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 528)

[spacer (SEQ ID NO: 362)] [J43 MP008 2001 (SEQ ID NO: 1013)] amino acid sequence:

[QGQSGQG] [ACRICQDHPKTKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSDDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 529)

J43 MP008 2001 amino acid sequence:

ACRICQDHPKTKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSDDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1084)

[spacer (SEQ ID NO: 918)] [J43 MP008 2001 (SEQ ID NO: 1014)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTAAGACGAAGTGA
ATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCCTCGGGTATCCCTGAAAG

AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 530)

[spacer (SEQ ID NO: 362)] [J43 MP009 2001 (SEQ ID NO: 1015)] amino acid sequence:

[QGQSGQG] [LIQCTGNLDHKCKHYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 531)

J43 MP009 2001 amino acid sequence:

LIQCTGNLDHKCKHYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1085)

[spacer (SEQ ID NO: 918)] [J43 MP009 2001 (SEQ ID NO: 1016)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CTTATTCAGTGCACTGGTAATCTTGATCATAAGTGCAAGC
ATTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCT (SEQ ID NO: 532)

[spacer (SEQ ID NO: 362)] [J43 MP010 2001 (SEQ ID NO: 1017)] amino acid sequence:

[QGQSGQG] [IPCHHSADHKHKCTSGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 533)

J43 MP010 2001 amino acid sequence:

IPCHHSADHKHKCTSGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDS DSKLYVFGSGTQLTVL (SEQ ID NO: 1086)

[spacer (SEQ ID NO: 918)] [J43 MP010 2001 (SEQ ID NO: 1018)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [ATTCCTTGCCATCATAGTGCTGATCATAAGCATAAGTGCA
CGAGTGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 534)

[spacer (SEQ ID NO: 362)] [J43 MP011 2001 (SEQ ID NO: 1019)] amino acid sequence:

[QGQSGQG] [SRQICADYNCHNKYKGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDS DSKLYVFGSGTQLTVL] (SEQ ID NO: 535)

J43 MP011 2001 amino acid sequence:

SRQICADYNCHNKYKGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDS DSKLYVFGSGTQLTVL (SEQ ID NO: 1087)

[spacer (SEQ ID NO: 918)] [J43 MP011 2001 (SEQ ID NO: 1020)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TCGCGGCAGATTTGCGCTGATTATAATTGCCATAATAAGT
ATAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCCTCGGGTATCCCTGAAAG

AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 536)

[spacer (SEQ ID NO: 362)] [J43 MP012 2001 (SEQ ID NO: 1021)] amino acid sequence:

[QGQSGQG] [QPCNPQIDHKIKCIYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 537)

J43 MP012 2001 amino acid sequence:

QPCNPQIDHKIKCIYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1088)

[spacer (SEQ ID NO: 918)] [J43 MP012 2001 (SEQ ID NO: 1022)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CAGCCTTGCAATCCGCAGATTGATCATAAGATTAAGTGCA
TTTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 538)

[spacer (SEQ ID NO: 362)] [J43 MP013 2001 (SEQ ID NO: 1023)] amino acid sequence:

[QGQSGQG] [HYTICMTHNKCKDMAGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 539)

J43 MP013 2001 amino acid sequence:

HYTICMTHNKCKDMAGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1089)

[spacer (SEQ ID NO: 918)] [J43 MP013 2001 (SEQ ID NO: 1024)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CATTATACTATTTGCATGACGCATAATAAGTGCAAGGATA
TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG
AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 540)

[spacer (SEQ ID NO: 362)] [J43 MP014 2001 (SEQ ID NO: 1025)] amino acid sequence:

[QGQSGQG] [ANSCLAVEHKCKHNYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 541)

J43 MP014 2001 amino acid sequence:

ANSCLAVEHKCKHNYGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1090)

[spacer (SEQ ID NO: 918)] [J43 MP014 2001 (SEQ ID NO: 1026)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCTAATAGTTGCCTTGCTGTTGAGCATAAGTGCAAGCATA
ATTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTTATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACT
GTCAAAATCACCTGCTCTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGT
CAGACCAGACCATTTTGCAAGTGATATATGATGATAATAAGCGCCCCTCGGGTATCCCTGAAAG

AATCTCTGGGTCCAGCTCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGAT
GAAGGTGACTATTACTGTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCA
GCGGAACCCAGCTCACCGTCCTA (SEQ ID NO: 542)

Reformatted anti-mouse PD-1 heavy and light chains:

J43v2 light chain (LC) amino acid sequence:

YELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSS
SGTTATLTIRDVRAEDEGDYYCFSGYVDSKLYVFGSGTQLTVLGGPKSSPKVTVFPSPPEEL
RTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNR
VSCQVTHEGETVEKSLSPAEL (SEQ ID NO: 543)

J43v2 LC nucleotide sequence 1:

TATGAGCTGACTCAGCCACCTTCAGCATCAGTCAATGTAGGAGAGACTGTCAAATCACCTGCT
CTGGGGACCAATTGCCGAAATATTTTGCAGATTGGTTTCATCAAAGGTCAGACCAGACCATTTT
GCAAGTGATATATGATGATAATAAGCGCCCCCTCGGGTATCCCTGAAAGAATCTCTGGGTCCAGC
TCAGGGACAACAGCCACCTTGACCATCAGAGATGTCCGGGCTGAGGATGAAGGTGACTATTACT
GTTTCTCAGGATATGTTGATAGTGATAGCAAATTGTATGTTTTTGGCAGCGGAACCCAGCTCAC
CGTCCTAGGTGGACCCAAGTCTTCTCCCAAAGTCACAGTGTTTCCACCTTCACCTGAGGAGCTC
CGGACAAACAAAGCCACACTGGTGTGTCTGGTTAATGACTTCTACCCGGGTTCTGCAACAGTGA
CCTGGAAGGCAAATGGAGCAACTATCAATGATGGGGTGAAGACTACAAAGCCTTCCAAACAGGG
CCAAAACACTACATGACCAGCAGCTACCTAAGTTTGACAGCAGACCAGTGGAATCTCACAACAGG
GTTTCTGCCAAGTTACCCATGAAGGGGAACTGTGGAGAAGAGTTTGTCCCCTGCAGAATGTC
TT (SEQ ID NO: 544)

J43v2 LC nucleotide sequence 2:

TACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACCTGTA
GCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAATCCT
GCAAGTGATCTACGACGACAACAAGCGGGCCAGCGGCATCCCCGAGAGAATCAGCGGAAGCAGC
AGCGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTACTACT
GCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGTCCGGTACCCAGCTGAC
AGTGCTGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCAAGCCCCGAGGAACTG
AGGACCAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACCGTGA

CCTGGAAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCACCAAGCCCAGCAAGCAGGG
CCAGAACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCCACAACAGA
GTGTCCTGCCAAGTGACCCACGAGGGGGAAACCGTGAAAAGAGCCTGAGCCCTGCCGAGTGCC
TG (SEQ ID NO: 545)

J43v2 mIgG2a heavy chain (HC) amino acid sequence:

EVRLLESGLLVKPEGSLKLSCVASGFTFSDYFMSWVRQAPGKGLEWVAHIYTKSYNYATYYSG
SVKGRFTISRDDSRSMVYLQMNLRTEDTATYYCTRDGSGYPSLDFWGQGTQVTVSSAKTTAPS
VYPLAPVCGDTTGSSVTLGCLVKGYFPEPVTLTWNSGSLSSGVHTFPAVLQSDLYTLSSSVTVT
SSTWPSQSITCNVAHPASSTKVDDKIEPRGPTIKPCPPCKCPAPNLLGGPSVFIFFPKIKDVL
ISLSPIVTCVVVDVSEDDPDVQISWVFNVEVHTAQQTQTHREDYNSTLRVVSALPIQHQQDWMSG
KEFKCKVNNKDLPAPIERTISKPKGSVRAPQVYVLPPEEEMTKKQVTLTCMVTDFMPEDIYVE
WTNNGKTELNYKNTEPVLDSGYSYFMYSKLRVEKKNWVERNSYSCSVHEGLHNHHTTKSFSRT
PGK (SEQ ID NO: 546)

J43v2 mIgG2a HC nucleic acid sequence:

GAGGTGCGGCTTCTGGAGTCTGGTGGAGGATTAGTGAAGCCTGAGGGGTCACTGAAACTCTCCT
GTGTGGCCTCTGGATTACCTTCAGTGACTATTTTCATGAGCTGGGTCCGCCAGGCTCCAGGGAA
GGGGCTGGAGTGGGTTGCTCACATATACACGAAAAGTTATAATTATGCAACTTATTACTCGGGT
TCGGTGAAAGGCAGATTCACCATCTCCAGAGATGATTCCCGAAGCATGGTCTACCTGCAAATGA
ACAACCTGAGAACTGAGGACACGGCCACTTATTACTGTACAAGAGATGGAAGCGGATATCCCTC
TCTGGATTTCTGGGGTCAAGGGACCCAAGTCACTGTCTCCTCAGCTAAAACAACAGCCCCATCG
GTCTATCCACTGGCCCCTGTGTGTGGAGATACAACTGGCTCCTCGGTGACTCTAGGATGCCTGG
TCAAGGGTTATTTCCCTGAGCCAGTGACCTTGACCTGGAACCTCTGGATCCCTGTCCAGTGGTGT
GCACACCTTCCAGCTGTCTTGCAGTCTGACCTCTACACCCTCAGCAGCTCAGTGACTGTAACC
TCGAGCACCTGGCCCAGCCAGTCCATCACCTGCAATGTGGCCCACCCGGCAAGCAGCACCAAGG
TGGACAAGAAAATTGAGCCCAGAGGGCCACAATCAAGCCCTGTCTCCATGCAATGCCCAGC
ACCTAACCTCTTGGGTGGACCATCCGTCTTCATCTTCCCTCCAAAGATCAAGGATGTACTCATG
ATCTCCCTGAGCCCCATAGTCACATGTGTGGTGGTGGATGTGAGCGAGGATGACCCAGATGTCC
AGATCAGCTGGTTTGTGAACAACGTGGAAGTACACACAGCTCAGACACAAACCCATAGAGAGGA
TTACAACAGTACTCTCCGGGTGGTCAGTGCCCTCCCCATCCAGCACCAGGACTGGATGAGTGGC
AAGGAGTTCAAATGCAAGGTCAACAACAAAGACCTCCAGCGCCCATCGAGAGAACCATCTCAA

AACCCAAAGGGTCAGTAAGAGCTCCACAGGTATATGTCTTGCCTCCACCAGAAGAAGAGATGAC
 TAAGAAACAGGTCCTCTGACCTGCATGGTCACAGACTTCATGCCTGAAGACATTTACGTGGAG
 TGGACCAACAACGGGAAAACAGAGCTAACTACAAGAACACTGAACCAGTCCTGGACTCTGATG
 GTTCTTACTTCATGTACAGCAAGCTGAGAGTGGAAAAGAAGAACTGGGTGGAAAGAAATAGCTA
 CTCCTGTTTCAGTGGTCCACGAGGGTCTGCACAATCACCACACGACTAAGAGCTTCTCCCGGACT
 CCGGGTAAA (SEQ ID NO: 547)

EXAMPLE 16: Activatable anti-mPD-1 antibodies with a range of masking efficiencies.

[000874] This example describes the modulation of anti-mPD-1 antibody masking efficiencies by masking moiety truncation and single amino-acid substitution.

[000875] Four J43 activatable antibodies were selected to generate families of activatable anti-mPD-1 antibodies by masking moiety truncation or amino acid substitution of one or more MM residues. Table 18 lists the designed MMs. Activatable J43v2 antibodies with the designed MM and 2001 substrate were produced by transfection of 30mL HEK293 cells and selected activatable antibodies (sequences MPtrunc ELISA activatable antibody LCs) were evaluated for masking by ELISA as described in Example 15 (Figure 23).

[000876] In some embodiments, the activatable antibody also includes a spacer sequence. In some embodiments, the spacer is joined directly to the MM of the activatable antibody. In some embodiments, the spacer is joined directly to the MM of the activatable antibody in the structural arrangement from N-terminus to C-terminus of spacer-MM-CM-AB. In some embodiments, the spacer joined directly to the N-terminus of MM of the activatable antibody is selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. In some embodiments, the spacer includes at least the amino acid sequence QGQSGQG (SEQ ID NO: 362). In some embodiments, the spacer includes at least the amino acid sequence QGQSGQ (SEQ ID NO: 913). In some embodiments, the spacer includes at least the amino acid sequence QGQSG (SEQ ID NO: 914). In some embodiments, the spacer includes at least the amino acid sequence QGQS (SEQ ID NO: 915). In some embodiments, the spacer includes at least the amino acid sequence QGQ (SEQ ID NO: 916). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 917). In

some embodiments, the spacer includes at least the amino acid residue Q. In some embodiments, the activatable antibody does not include a spacer sequence.

[000877] While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art appreciate that the activatable anti-PD-1 antibodies of the disclosure can include any suitable spacer sequence, such as, for example, a spacer sequence selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art will also appreciate that activatable anti-PD-1 antibodies of the disclosure in some embodiments do not include a spacer sequence.

Table 18: Masking moiety sequences

Mask	Amino Acid Sequence	SEQ ID NO:
MP8-1	QDHPKTKWNS	548
MP8-2	ACRICQDHPATKWNS	549
MP8-3	ACRICQDHPKTAWNS	550
MP8-4	ACRICQDAPKTKWNS	551
MP8-5	ACRICQDHAKTKWNS	552
MP8-6	DHPATKWNS	553
MP8-7	DHPKTAWNS	554
MP8-8	DAPKTKWNS	555
MP8-9	DAPATKWNS	556
MP8-10	ACRICQDHP	557
MP7-1	HPQSKDTL	558
MP7-2	HPKSQDTL	559
MP7-3	TIPRCGQHPLCLDTL	560
MP7-4	HPLSLDTL	561
MP7-5	HPASKDTL	562
MP7-6	HPKSADTL	563
MP5-1	PGNCHPLQKELCQFI	564

MP5-2	HPLQKELAQFI	565
MP5-3	HPLALELAQFI	566
MP5-4	PGNCHPLQLELCQFI	567
MP3-1	TNIWSCQTYCDHAHA	568
MP3-2	TNIWSCQTYCDHAHL	569
MP3-3	TNIWSCQTYCDHLHA	570
MP3-4	TNIWSCQTYCDHKHA	571

[spacer (SEQ ID NO: 362)] [J43 MP5-2 2001 (SEQ ID NO: 1027)] amino acid sequence:

[QGQSGQG] [HPLQKELAQFIGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVK
ITCSGDQLPKYFADWFHQRSDDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEG
DYFCFSGYVDSKLYVFGSGTQLTVL] (SEQ ID NO: 572)

J43 MP5-2 2001 amino acid sequence:

HPLQKELAQFIGGGSSGGSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPK
YFADWFHQRSDDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYFCFSGYVD
SDSKLYVFGSGTQLTVL (SEQ ID NO: 1113)

[spacer (SEQ ID NO: 918)] [J43 MP5-2 2001 (SEQ ID NO: 1038)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CATCCTCTGCAGAAGGAGCTGGCCCAGTTTATTGGAGGTG
GCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGG
CTCTTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACC
TGTAGCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAA
TCCTGCAAGTGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGAGAGAATCAGCGGAAG
CAGCAGCGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTAC
TACTGCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCGGCTCCGGTACCCAGC
TGACAGTGCTGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCACCCCAAGCCCCGAGGA
ACTGAGGACCAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACC
GTGACCTGGAAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCAAGCCCAGCAAGC
AGGGCCAGAACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCACAA

CAGAGTGTCTGCGCAAGTGACCCACGAGGGGGAAACCGTGGAAGAGCCTGAGCCCTGCCGAG
TGCCTG (SEQ ID NO: 573)

[spacer (SEQ ID NO: 362)] [J43 MP7-1 2001 (SEQ ID NO: 1039)] amino acid sequence:

[QGQSGQG] [HPQSKDTLGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITC
SGDQLPKYFADWFHQRSQDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYY
CFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 574)

J43 MP7-1 2001 amino acid sequence:

HPQSKDTLGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFA
DWFHQRSQDQTLQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDSDS
KLYVFGSGTQLTVL (SEQ ID NO: 1114)

[spacer (SEQ ID NO: 918)] [J43 MP7-1 2001 (SEQ ID NO: 1029)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CATCCGCAGTCTAAGGATACTTTGGGAGGTGGCTCGAGCG
GCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGCTCTTACGA
GCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACCTGTAGCGGC
GACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAATCCTGCAAG
TGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGAGAGAATCAGCGGAAGCAGCAGCGG
CACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTACTACTGCTTT
AGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGCTCCGGTACCCAGCTGACAGTGC
TGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCAAGCCCCGAGGAACTGAGGAC
CAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACCCTGACCTGG
AAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCACCAAGCCCAGCAAGCAGGGCCAGA
ACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCCACAACAGAGTGTC
CTGCCAAGTGACCCACGAGGGGGAAACCGTGGAAGAGCCTGAGCCCTGCCGAGTGCCTG
(SEQ ID NO: 575)

MPtrunc ELISA Activatable Antibody LCs:

[spacer (SEQ ID NO: 362)] [J43 MP7-5 2001 (SEQ ID NO: 1115)] amino acid sequence:

[QGQSGQG] [HPASKDTLGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITC
SGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYY
CFSGYVDS DSKLYVFGSGTQLTVL] (SEQ ID NO: 576)

J43 MP7-5 2001 amino acid sequence:

HPASKDTLGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFA
DWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS D
SKLYVFGSGTQLTVL (SEQ ID NO: 1115)

[spacer (SEQ ID NO: 918)] [J43 MP7-5 2001 (SEQ ID NO: 1031)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CATCCGGCGTCTAAGGATACTTTGGGAGGTGGCTCGAGCG
GCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGCTCTTACGA
GCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACCTGTAGCGGC
GACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAATCCTGCAAG
TGATCTACGACGACAACAAGCGGCCCAGCGGCATCCCCGAGAGAATCAGCGGAAGCAGCAGCGG
CACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTACTACTGCTTT
AGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGCTCCGGTACCCAGCTGACAGTGC
TGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCAAGCCCCGAGGAACTGAGGAC
CAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACCGTGACCTGG
AAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCAAGCCCAGCAAGCAGGGCCAGA
ACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCCACAACAGAGTGTC
CTGCCAAGTGACCCACGAGGGGGAAACCGTGGAAGAGCCTGAGCCCTGCCGAGTGCCTG
(SEQ ID NO: 577)

[spacer (SEQ ID NO: 362)] [J43 MP8-2 2001 (SEQ ID NO: 1032)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDS DSKLYVFGSGTQLTVL] (SEQ ID NO: 578)

J43 MP8-2 2001 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1116)

[spacer (SEQ ID NO: 918)] J43 MP8-2 2001 (SEQ ID NO: 1033)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTGCGACGAAGTGGA
ATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGCTCTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAG
TGAAGATCACCTGTAGCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAG
CGACCAGACAATCCTGCAAGTGATCTACGACGACAACAAGCGGCCCCAGCGGCATCCCCGAGAGA
ATCAGCGGAAGCAGCAGCGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACG
AGGGCGACTACTACTGCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGCTC
CGGTACCCAGCTGACAGTGCTGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCCA
AGCCCCGAGGAAGT GAGGACCAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTG
GCAGCGCCACCGTGACCTGGAAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCACCAA
GCCCAGCAAGCAGGGCCAGAACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGG
AAGTCCCAACAACAGAGTGTCTTGCCAAGTGACCCACGAGGGGAAACCGTGGAAGAGCCTGA
GCCCTGCCGAGTGCCTG (SEQ ID NO: 579)

[spacer (SEQ ID NO: 362)] [J43 MP8-8 2001 (SEQ ID NO: 1034)] amino acid sequence:

[QGQSGQG] [DAPKTKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKIT
CSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDY
YCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 580)

J43 MP8-8 2001 amino acid sequence:

DAPKTKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYF
ADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS
SKLYVFGSGTQLTVL (SEQ ID NO: 1117)

[spacer (SEQ ID NO: 918)] J43 MP8-8 2001 (SEQ ID NO: 1035)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GATGCTCCTAAGACGAAGTGGAATTCTGGAGGTGGCTCGA
GCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGCTCTTA
CGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACCTGTAGC
GGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAATCCTGC
AAGTGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGAGAGAATCAGCGGAAGCAGCAG
CGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTACTACTGC
TTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGCTCCGGTACCCAGCTGACAG
TGCTGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCAAGCCCCGAGGAACTGAG
GACCAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACCGTGACC
TGGAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCACCAAGCCCAGCAAGCAGGGCC
AGAACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCCACAACAGAGT
GTCCTGCCAAGTGACCCACGAGGGGGAAACCGTGAAAAGAGCCTGAGCCCTGCCGAGTGCCTG
(SEQ ID NO: 581)

[spacer (SEQ ID NO: 362)] [J43 MP8-9 2001 (SEQ ID NO: 1036)] amino acid sequence:

[QGQSGQG] [DAPATKWNSSGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKIT
CSGDQLPKYFADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDY
YCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 582)

J43 MP8-9 2001 amino acid sequence:

DAPATKWNSSGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYF
ADWFHQRS DQTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDSD
SKLYVFGSGTQLTVL (SEQ ID NO: 1118)

[spacer (SEQ ID NO: 918)] [J43 MP8-9 2001 (SEQ ID NO: 1037)] nucleic acid sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GATGCTCCTGCGACGAAGTGGAATTCTGGAGGTGGCTCGA
GCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGAGGCTCTTA
CGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACAGTGAAGATCACCTGTAGC
GGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGAGCGACCAGACAATCCTGC
AAGTGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGAGAGAATCAGCGGAAGCAGCAG
CGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGACGAGGGCGACTACTACTGC

TTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTTCGGCTCCGGTACCCAGCTGACAG
 TGCTGGGCGGACCTAAGAGCAGCCCCAAAGTGACCGTGTTCCCCCAAGCCCCGAGGAACTGAG
 GACCAACAAGGCCACCCTCGTGTGCCTCGTGAACGACTTCTACCCTGGCAGCGCCACCGTGACC
 TGGAAAGCCAATGGCGCCACCATCAACGACGGCGTGAAAACCACCAAGCCCAGCAAGCAGGGCC
 AGAACTACATGACCAGCAGCTACCTGAGCCTGACCGCCGACCAGTGGAAGTCCCACAACAGAGT
 GTCCTGCCAAGTGACCCACGAGGGGGAAACCGTGGAAAAGAGCCTGAGCCCTGCCGAGTGCCTG
 (SEQ ID NO: 583)

[spacer (SEQ ID NO: 362)][J43 MP8-2 2003 (SEQ ID NO: 1119) amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSISSGLLSGRSANPRGGGSYELTQPPSASVNV
 GETVKITCSGDQLPKYFADWFHQRSDDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVR
 AEDEGDYYCFSGYVDSKLYVFGSGTQLTVL] (SEQ ID NO: 1120)

J43 MP8-2 2003 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSISSGLLSGRSANPRGGGSYELTQPPSASVNVGETVKITCSG
 DQLPKYFADWFHQRSDDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCF
 SGYVDSKLYVFGSGTQLTVL (SEQ ID NO: 1119)

[spacer (SEQ ID NO: 918)][J43 MP8-2 2003 (SEQ ID NO: 1121) nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTGCGACGAAGTGGA
 ATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGCTGCTCCGGCAGATCCGCCAA
 TCCTCGTGGCGGAGGATCCTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAG
 ACAGTGAAGATCACCTGTAGCGGCGACCAGCTGCCCCAAGTACTTCGCCGACTGGTTCCACCAGC
 GGAGCGACCAGACAATCCTGCAAGTGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGA
 GAGAATCAGCGGAAGCAGCAGCGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAG
 GACGAGGGCGACTACTACTGCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTTCG
 GCTCCGGTACCCAGCTGACAGTGCTG] (SEQ ID NO: 1122)

[spacer (SEQ ID NO: 362)][J43 MP8-2 2005 (SEQ ID NO: 1123) amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGAVGLLAPPSGRSANPRGGGSYELTQPPSASVN
 VGETVKITCSGDQLPKYFADWFHQRSDDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDV
 RAEDEGDYYCFSGYVDSKLYVFGSGTQLTVL] (SEQ ID NO: 1124)

J43 MP8-2 2005 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGAVGLLAPPSGRSANPRGGGSYELTQPPSASVNVGETVKITCS
GDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYC
FSGYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1123)

[spacer (SEQ ID NO: 362)][J43 MP8-2 2008 (SEQ ID NO: 1127) amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSISSGLLSGRSDGHGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 1128)

J43 MP8-2 2008 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSISSGLLSGRSDGHGGGSYELTQPPSASVNVGETVKITCSGD
QLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFS
GYVDSDSKLYVFGSGTQLTVL (SEQ ID NO: 1127)

[spacer (SEQ ID NO: 918)][J43 MP8-2 2008 (SEQ ID NO: 1129) nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTGCGACGAAGTGGA
ATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACCA
GCACGGCGGAGGATCCTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTGGGCGAGACA
GTGAAGATCACCTGTAGCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGTTCCACCAGCGGA
GCGACCAGACAATCCTGCAAGTGATCTACGACGACAACAAGCGGCCAGCGGCATCCCCGAGAG
AATCAGCGGAAGCAGCAGCGGCACCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGAC
GAGGGCGACTACTACTGCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTCCGGCT
CCGGTACCCAGCTGACAGTGCTG] (SEQ ID NO: 1130)

[spacer (SEQ ID NO: 362)][J43 MP8-2 2012 (SEQ ID NO: 1131) amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSISSGLLSGRSANPGGGSYELTQPPSASVNVG
ETVKITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
EDEGDYYCFSGYVDSDSKLYVFGSGTQLTVL] (SEQ ID NO: 1132)

J43 MP8-2 2012 amino acid sequence:

ACRICQDHPATKWNSGGSSGGSISSGLLSGRSANPGGGSYELTQPPSASVNVGETVKITC
SGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEG
DYCYFSGYVDSKLYVFGSGTQLTVL (SEQ ID NO: 1131)

[spacer (SEQ ID NO: 918)][J43 MP8-2 2012 (SEQ ID NO: 616) nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTGCGACGAAGT
GGAATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATC
CGCTAATCCCGGCGGAGGATCCTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTG
GGCGAGACAGTGAAGATCACCTGTAGCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGT
TCCACCAGCGGAGCGACCAGACAATCCTGCAAGTGATCTACGACGACAACAAGCGGCCAG
CGGCATCCCCGAGAGAATCAGCGGAAGCAGCAGCGGCACCACCGCCACCCTGACCATTAGA
GATGTGCGGGCCGAGGACGAGGGCGACTACTACTGCTTTAGCGGCTACGTGGACAGCGACA
GCAAGCTGTACGTGTTCCGGCTCCGGTACCCAGCTGACAGTGCTG] (SEQ ID
NO: 1133)

[spacer (SEQ ID NO: 362)][J43 MP8-2 2011 (SEQ ID NO: 1134) amino acid sequence:

[QQQSGQG] [ACRICQDHPATKWNSGGSSGGSISSGLLSGRSDNPGGGSYELTQPPSASV
NVGETVKITCSGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLT
IRDVRAEDEGDYCYFSGYVDSKLYVFGSGTQLTVL] (SEQ ID NO: 1135)

J43 MP8-2 2011 amino acid sequence:

ACRICQDHPATKWNSGGSSGGSISSGLLSGRSDNPGGGSYELTQPPSASVNVGETVKITC
SGDQLPKYFADWFHQRSQDQILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEG
DYCYFSGYVDSKLYVFGSGTQLTVL (SEQ ID NO: 1134)

[spacer (SEQ ID NO: 918)][J43 MP8-2 2011 (SEQ ID NO: 1136) nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGTGCCGTATTTGTCAGGATCATCCTGCGACGAAGT
GGAATTCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTGTCCGGCAGATC
CGACAATCCCGGCGGAGGATCCTACGAGCTGACCCAGCCTCCTAGCGCCTCCGTGAATGTG
GGCGAGACAGTGAAGATCACCTGTAGCGGCGACCAGCTGCCCAAGTACTTCGCCGACTGGT
TCCACCAGCGGAGCGACCAGACAATCCTGCAAGTGATCTACGACGACAACAAGCGGCCAG
CGGCATCCCCGAGAG

AATCAGCGGAAGCAGCAGCGGCACCGCCACCCTGACCATTAGAGATGTGCGGGCCGAGGAC
 GAGGGCGACTACTACTGCTTTAGCGGCTACGTGGACAGCGACAGCAAGCTGTACGTGTTTCGGCT
 CCGGTACCCAGCTGACAGTGCTG] (SEQ ID NO: 1137)

EXAMPLE 17: Activatable anti-mouse PD-1 J43 antibodies reduce incidence of diabetes in NOD mice

[000878] In this Example, anti-PD-1 J43 activatable antibodies were analyzed for the ability to protect from anti-PD-1 induction of diabetes in NOD mice. The NOD mice, substrain NOD/ShiLtJ, were obtained from Jackson Laboratory at 8 weeks and acclimated on site for 2 weeks. At 10 weeks, mice were checked for diabetes prior to enrollment, grouped, and dosed as set forth in Table 4.

Table 4. Dosing regimen

Group	Count	Gender	Treatment	Dose (mg/kg)	Dose volume (mL/kg)	Schedule	Route
1	7	F	mIgG2a (C1.18.4)	10	10	q7dx1	IP
2	7	F	Anti-PD-1 J43 m2a	10	10	q7dx1	IP
3	7	F	Anti-PD-1 J43 m2a	3	10	q7dx1	IP
4	7	F	Anti-PD-1 J43 m2a	1	10	q7dx1	IP
5	7	F	J43 MP7-1 2001 m2a	10	10	q7dx1	IP
6	7	F	J43 MP7-1 2001 m2a	3	10	q7dx1	IP
7	7	F	J43 MP8-2 2001 m2a	10	10	q7dx1	IP
8	7	F	J43 MP8-2 2001 m2a	3	10	q7dx1	IP

[000879] Figure 24 which plots % non-diabetic versus number of days post dose, shows that anti-PD-1 J43 antibody induced diabetes in NOD mice with increased frequency as dosage increased. At day fourteen post dose, the percentage of non-diabetic mice for the antibody-treated groups was 14%, 43% and 71% for the 20 mg/kg, 3 mg/kg and 1 mg/kg dose groups, respectively. Activatable antibodies J43 MP7-1 2001 m2a and J43 MP8-2 2001 m2a required increased doses to induce diabetes at frequencies comparable to the parental antibody. At day fourteen post dose with J43 MP7-1 2001 m2a, 57% of the 20 mg/kg group remained non-diabetic and all of the 3 mg/kg were non-diabetic. At day fourteen post dose with J43 MP8-2 2001 m2a, 71% of the 20 mg/kg group and 86% of the 3 mg/kg group remained non-diabetic.

EXAMPLE 18: Evaluation of efficiency of masking moieties

[000880] This example describes additional activatable anti-PD-1 antibodies that exhibit reduced binding to hPD-1.

[000881] Examples of additional activatable antibodies of the disclosure comprising anti-PD1 antibody A1.5 and a variety of mask and substrate combinations were produced using techniques as described herein. The amino acid and nucleic acid sequences of these activatable anti-PD-1 antibody variable domains of the disclosure are provided below. Antibodies were produced as hIgG4 containing a single amino acid substitution, S228P (Angal, et al. 1993. Mol Immunol 30:105-8.) HC and hK LC format.

[000882] Masking efficiencies of several of these activatable antibodies were determined as described herein. The results are shown in Figures 25A and 25B.

[000883] In some embodiments, the activatable antibody also includes a spacer sequence. In some embodiments, the spacer is joined directly to the MM of the activatable antibody. In some embodiments, the spacer is joined directly to the MM of the activatable antibody in the structural arrangement from N-terminus to C-terminus of spacer-MM-CM-AB. In some embodiments, the spacer joined directly to the N-terminus of MM of the activatable antibody is selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. In some embodiments, the spacer includes at least the amino acid sequence QGQSGQG (SEQ ID NO: 362). In some embodiments, the spacer includes at least the amino acid sequence QGQSGQ (SEQ ID NO: 913). In some embodiments, the spacer includes at least the amino acid sequence QGQSG (SEQ ID NO: 914). In some embodiments, the spacer includes at least the amino acid sequence QGQS (SEQ ID NO: 915). In some embodiments, the spacer includes at least the amino acid sequence QGQ (SEQ ID NO: 916). In some embodiments, the spacer includes at least the amino acid sequence QG (SEQ ID NO: 917). In some embodiments, the spacer includes at least the amino acid residue Q. In some embodiments, the activatable antibody does not include a spacer sequence. . Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG (SEQ ID NO: 2045), QG (SEQ ID NO: 2046), and G.

[000884] While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art appreciate that the activatable anti-PD-1 antibodies of the disclosure can include any suitable spacer sequence, such as, for example, a spacer sequence selected from the group consisting of QGQSGQG (SEQ ID NO: 362), QGQSGQ (SEQ ID NO: 913), QGQSG (SEQ ID NO: 914), QGQS (SEQ ID NO: 915), QGQ (SEQ ID NO: 916), QG (SEQ ID NO: 917), and Q. . Additional examples of spacers include GQSGQG (SEQ ID NO: 2042), QSGQG (SEQ ID NO: 2043), SGQG (SEQ ID NO: 2044), GQG (SEQ ID NO: 2045), QG (SEQ ID NO: 2046), and G. While the sequences shown below include the spacer sequence of SEQ ID NO: 362, those of ordinary skill in the art will also appreciate that activatable anti-PD-1 antibodies of the disclosure in some embodiments do not include a spacer sequence.

[spacer (SEQ ID NO: 362)][PD1 1.5 PD01 2003 (SEQ ID NO: 1138) amino acid sequence:
[QGQSGQG] [AMSGCSWSAFCPYLAGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSA
SVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTL
TISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1139)

PD1 1.5 PD01 2003 amino acid sequence:

AMSGCSWSAFCPYLAGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSASVGDRVTITC
RASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDF
ATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1138)

[spacer (SEQ ID NO: 918)] [PD1 1.5 PD001 2003 (SEQ ID NO: 1141)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGATGAGTGGGTGCTCGTGGTCTGCTTTTTGCCCGTATT
TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGCCAA
TCCTCGTGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTG
GGCGACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCA
TGA ACTGGTTCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCA
GGGCAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCTGACCATC
AGCAGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGA
CCTTTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1142)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD01 2012 (SEQ ID NO: 1144) amino acid sequence:

[QGQSGQG] [AMSGCSWSAFCPYLGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLSASV
GDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTII
SSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1143)

PD1 1.5 PD01 2012 amino acid sequence:

AMSGCSWSAFCPYLGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLSASVGDRVITITCRA
SESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTIISSMQPEDFAT
YYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1144)

[spacer (SEQ ID NO: 918)] [PD1 1.5 PD01 2012 (SEQ ID NO: 1145)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGATGAGTGGGTGCTCGTGGTCTGCTTTTTGCCCGTATT
TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGCTAA
TCCCGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1146)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD01 2011 (SEQ ID NO: 1148) amino acid sequence:

[QGQSGQG] [AMSGCSWSAFCPYLGGGSSGGSISSGLLSGRSDNPGGGSDIQLTQSPSSLSASV
GDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTII
SSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1147)

PD1 1.5 PD01 2011 amino acid sequence:

AMSGCSWSAFCPYLGGGSSGGSISSGLLSGRSDNPGGGSDIQLTQSPSSLSASVGDRVITITCRA
SESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTIISSMQPEDFAT
YYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1148)

[spacer (SEQ ID NO: 918)] [PD1 1.5 PD01 2011 (SEQ ID NO: 1149)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGATGAGTGGGTGCTCGTGGTCTGCTTTTTTGCCCGTATT
TGGCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCCCGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1150)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2001 (SEQ ID NO: 1152)] amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1151)

PD1 1.5 PD34 2001 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1152)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2001 (SEQ ID NO: 1153)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1154)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 1004/GG/0001 (SEQ ID NO: 1156) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGS AVGLLAPPGGLSGRSDNHGGGSDIQLTQSPS
SLSASVGDRVTITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSGVPSRFSGSGSGT
DFTLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1155)

PD1 1.5 PD34 1004/GG/0001 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSAVGLLAPPGGLSGRSDNHGGGSDIQLTQSPSSLSASVGDRV
TITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQ
PEDFATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1156)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2005 (SEQ ID NO: 1160) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSAVGLLAPPSGRSANPRGGGGSDIQLTQSPSS
LSASVGDRV TITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTD
FTLTISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1159)

PD1 1.5 PD34 2005 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSAVGLLAPPSGRSANPRGGGGSDIQLTQSPSSLSASVGDRV
TITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQP
EDFATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1160)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2010 (SEQ ID NO: 1164) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDYHGGGSDIQLTQSPSSLSAS
VGDRV TITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1163)

PD1 1.5 PD34 2010 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDYHGGGSDIQLTQSPSSLSASVGDRV TITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1164)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2014 (SEQ ID NO: 1168) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDNIGGGSDIQLTQSPSSLSAS
VGDRV TITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1167)

PD1 1.5 PD34 2014 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNIGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1168)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2003 (SEQ ID NO: 1170) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSA
 SVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
 TISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1171)

PD1 1.5 PD34 2003 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSASVGDRVITITC
 RASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDF
 ATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1170)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2003 (SEQ ID NO: 1172)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGCCAA
 TCCTCGTGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTG
 GGCGACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCA
 TGA ACTGGTTCCAGCAGAAGCCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCA
 GGGCAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATC
 AGCAGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGA
 CCTTTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1173)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2006 (SEQ ID NO: 1174) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDDHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1175)

PD1 1.5 PD34 2006 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDDHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1174)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2006 (SEQ ID NO: 1176)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACGA
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1177)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2007 (SEQ ID NO: 1178)] amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDIHGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1179)

PD1 1.5 PD34 2007 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDIHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1178)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2007 (SEQ ID NO: 1180)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAT
 TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1181)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2009 (SEQ ID NO: 1182) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDTHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1183)

[PD1 1.5 PD34 2009 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDTHGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1182)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2009 (SEQ ID NO: 1184)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAC
TCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1185)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2008 (SEQ ID NO: 1186) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDQHGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1187)

PD1 1.5 PD34 2008 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSDQHGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1186)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2008 (SEQ ID NO: 1188)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACCA
 GCACGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1189)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2012 (SEQ ID NO: 1190) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLSAS
 VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
 ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1191)

PD1 1.5 PD34 2012 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSISSGLLSGRSANPGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1190)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2012 (SEQ ID NO: 1192)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGCTAA
 TCCCGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG

CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1193)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2013 (SEQ ID NO: 1194) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSANIGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1195)

PD1 1.5 PD34 2013 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSANIGGGSDIQLTQSPSSLSASVGDRVITITCR
ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLTISSMQPEDFA
TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1194)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2013 (SEQ ID NO: 1196)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGCTAA
TATTGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1197)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD34 2011 (SEQ ID NO: 1198) amino acid sequence:

[QGQSGQG] [TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNPGGGSDIQLTQSPSSLSAS
VGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQGSVPSRFSGSGSGTDFTLT
ISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1199)

PD1 1.5 PD34 2011 amino acid sequence:

TSYCSIEHYPCNTHHGGGSSGGSSISSGLLSGRSDNPGGGSDIQLTQSPSSLSASVGDRVITITCR
 ASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDFA
 TYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1198)

[spacer (SEQ ID NO: 1125)] [PD1 1.5 PD034 2011 (SEQ ID NO: 1200)] nucleotide sequence:

[CAGGGGCAATCTGGCCAGGGG] [ACGTCTTACTGCAGTATTGAGCATTACCCCTGCAATACAC
 ATCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTGCTCCGGCAGATCCGACAA
 TCCCGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTGGGC
 GACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCATGA
 ACTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCAGGG
 CAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGC
 AGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGACCT
 TTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1201)

[spacer (SEQ ID NO: 362)][PD1 1.5 PD006 2003 (SEQ ID NO: 1202) amino acid sequence:

[QGQSGQG] [APRCYMFASYCKSQYGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSA
 SVGDRVITITCRASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLT
 TISSMQPEDFATYYCQQSKDVPWTFGQGTKLEIK] (SEQ ID NO: 1203)

PD1 1.5 PD006 2003 amino acid sequence:

APRCYMFASYCKSQYGGGSSGGSSISSGLLSGRSANPRGGGSDIQLTQSPSSLSASVGDRVITITC
 RASESVDAYGISFMNWFQQKPGKAPKLLIYAASNQSGVPSRFSGSGSGTDFTLTISSMQPEDF
 ATYYCQQSKDVPWTFGQGTKLEIK (SEQ ID NO: 1202)

[spacer (SEQ ID NO: 918)] [PD1 1.5 PD006 2003 (SEQ ID NO: 1204)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCGCCTAGGTGCTATATGTTTTCGTCGTATTGCAAGAGTC
 AGTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTGCTCCGGCAGATCCGCCAA
 TCCTCGTGGCGGAGGATCCGATATCCAGCTGACCCAGAGCCCTAGCAGCCTGTCTGCCAGCGTG
 GGCGACAGAGTGACCATCACCTGTAGAGCCAGCGAGAGCGTGGACGCTTACGGCATCAGCTTCA
 TGAAGTGGTTCCAGCAGAAGCCCGGCAAGGCCCCCAAGCTGCTGATCTACGCCGCCAGCAATCA

GGGCAGCGGCGTGCCAAGCAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATC
 AGCAGCATGCAGCCCGAGGACTTCGCCACCTACTACTGCCAGCAGAGCAAGGACGTGCCCTGGA
 CCTTTGGCCAGGGTACCAAGCTGGAAATCAAG] (SEQ ID NO: 1205)

EXAMPLE 19: Activatable anti-PD1 Nivolumab masking moieties

[000885] This example describes identification of masking moieties (MM) that reduce binding of anti-PD1 Nivolumab antibody to its target.

[000886] Anti-PD1 Nivolumab (NV1) (see, e.g., US 8,008,449) was used to screen libraries using a method similar to that described in PCT International Publication Number WO 2010/081173, published 15 July 2010. The screening consisted of one round of MACS and four rounds of FACS sorting. For the initial MACS, approximately 6×10^{11} cells were incubated with NV1 antibody at a concentration of 100nM, and approximately 3×10^7 binders were collected using Protein-A Dynabeads (Life Technologies, Carlsbad, CA). FACS rounds were conducted labeling cells with DyLight 650 (Thermo-Fisher) labeled NV1 antibody and collecting cells with strongest fluorescence as follows: 20nM NV1-650 collecting brightest 13% for FACS round 1 (F1), 2nM NV1-650 collecting brightest 2% for FACS round 2 (F2-1), 1nM NV1-650 collecting brightest 1% for FACS round 3 (F3-1b) and 1nM NV1-650 collecting two populations, the brightest 88% and the brightest 1% for FACS round 4 (F4-1b.1 and F4-1b.2). Individual peptide clones from F4-1b.1 and F4-1b.2 were identified by sequence analysis (Table 19), and selected peptide clones NV01-NV12 were subsequently verified for their ability to bind NV1 (masking moiety NV001 is also referred to herein as NV01 and/or NV-01; masking moiety NV002 is also referred to herein as NV02 and/or NV-02, and so on).

Table 19

Mask	Amino Acid Sequence	SEQ ID NO:
NV01	RYCHAANPDRFCGIY	1206
NV02	PRVCSTDGGDYCLLP	1207
NV03	PRPQCHHRHNCPDHP	1208
NV04	KCSRPAHQNPDRCSR	1209
NV05	ASYRCPDYKCSHTKH	1210
NV06	LPRCPDHPIKCIETK	1211

NV07	YTFGCPDRYCDRAAT	1212
NV08	RGCPDFNPPSHCYTA	1213
NV09	RDYCGPQSPDYCHEI	1214
NV10	PNKPCPDLQCYVTNY	1215
NV11	PRVACGEPDLCYSNT	1216
NV12	RGCKKHTISTLTCPD	1217
NV13	PAYRCPDRPPCKNQM	1218
NV14	NARCYPYFGDNCHMN	1220
NV15	PTLRCPDRWCYDSPR	1221
NV16	PSNLCPDKWCQTWRS	1222
NV17	TPRYCAASYCPAHGY	1223
NV18	RPGCGAVSPRCPDAP	1224
NV19	VLRCHKQNPDCNNH	1225
NV20	GVKSCREPDFCSRGS	1226
NV21	RNNLCPDYSCNNHNS	1227
NV22	RAACHRLNPDACTNG	1228
NV23	VCQSDRIPDYVTCTD	1229
NV24	RNCRIASINPDYCNI	1230
NV25	KEWRCPDYKCKPSYH	1231
NV26	NLRICHKSLCPDYIK	1232
NV27	NTHKCSNTNICPSFN	1233
NV28	STRYCQASQCQMSPY	1234
NV29	THRFCTASLCNKNTS	1235
NV30	YTLCNTRSPDWCPNK	1236
NV31	IRCTTGQSPDYCPQS	1237
NV32	RCNQPDKNDQMLCNI	1238
NV33	GTCRTDHQSPDYCY Y	1239
NV34	RGCFRSGDSLGMCPD	1240
NV35	SGCFDSNEHRHCSRI	1241
NV36	NRCMKLWYNPDCVAR	1242

NV37	PLCARPHYWSPCDQS	1243
NV38	DSKCHPNSPDYCFNS	1244
NV39	NGSCRPLGGDFCGNR	1245
NV40	KTRCIEMSGDYCAKS	1246
NV41	IRPCMYNWGDLCNQF	1247
NV42	VKTCMENNPDYCYNN	1248
NV43	LRMCFEASGDYCDQQ	1249
NV44	IRKCQLDGPDQCMLT	1250
NV45	KWKCHKNNPNYCNR	1251
NV46	RTMCLDTNPDYCQSH	1252
NV47	LAACHSMDSHRCPDY	1253
NV48	RSPCIHNATMCPDYT	1254
NV49	MPRCPDWPPRCSMVI	1255
NV50	VRQLCRLPDYCPSGK	1256
NV51	PRPPCAQSLNCPDRA	1257
NV52	SFGRCTLVRTCPDFM	1258
NV53	RDKPCPDFSCATIHY	1259
NV54	ATKPCPDRWCTMSTL	1260
NV55	SSNRCPDLRCTHHNM	1261
NV56	RGSMCPDLHCSLSHI	1262
NV57	NYQRCPDRTCMHNII	1263
NV58	QKRPCPDRKCHAHYN	1264
NV59	QNHRCPDRWCNKTTN	1265
NV60	RLNLCPDKHCHMTNL	1266
NV61	PQDRCPDKRCTNPGN	1267
NV62	SRWRCPDYKCEHGKY	1268
NV63	YENQCPDLYCNRYSM	1269
NV64	TARSCPVFNCPDNNS	1270
NV65	MDQRCPDWCTSKPK	1271
NV66	GDLRCPDRLCPRHSL	1272

NV67	IQYLCPDYHCKASNN	1273
NV68	QHHRCPDRYCNSNNN	1274
NV69	TVALCPDYSCYHINN	1275
NV70	SPWRCPDRYCLSNDH	1276
NV71	SSKRCPDRFCNKTHA	1277
NV72	HTDRCPDYKCSQNH	1278
NV73	SRSNCTPQRCNSDYH	1279
NV74	FAARCPDKYCAIHTN	1280
NV75	GSARCPDLVCQQTKQ	1281
NV76	RNLMCPDKFCNKNTK	1282
NV77	NIRLCPDKVCTPTWV	1283
NV78	MTDLCPDAHCAKTHM	1284
NV79	PYSRLCAYPCPDFVG	1285
NV80	LCGCARSPDYCKCRG	1286
NV81	WGRCERVPDCCCPRG	1287
NV82	TRNTCHTRICYGMAC	1288
NV83	CVCTSCSSYWTLCPD	1289
NV84	LCCSRGSNCPDRCTW	1290
NV85	CCPLCQANMCPDNQS	1291
NV86	ECKLCCPDLYCGGTM	1292
NV87	CSNPMCAYCCPDLIL	1293
NV88	CPRCNTYSKHDCYHQ	1294
NV89	FCCASKMPAPSNCHT	1295

EXAMPLE 20: Activatable anti-PD1 Nivolumab Antibodies

[000887] This example describes examples of activatable anti-PD1 Nivolumab antibodies of the disclosure.

[000888] Activatable anti-PD1 NV1 antibodies comprising an anti-PD1 NV1 masking moiety, a cleavable moiety, and an anti-PD1 NV1 antibody of the disclosure were produced according to methods similar to those described in PCT Publication Nos. WO 2009/025846 and WO 2010/081173. One embodiment of the disclosure is an anti-PD-1 activatable antibody

comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1346. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising a light chain comprising amino acid sequence SEQ ID NO: 626. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1346 and a light chain comprising amino acid sequence SEQ ID NO: 626. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising the CDRs of an anti-PD-1 activatable antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1346 and a light chain comprising amino acid sequence SEQ ID NO: 626.

[000889] The amino acid and nucleic acid sequences of several activatable anti-PD1 antibody variable domains of the disclosure are provided below. Antibodies were produced as hIgG4 containing a single amino acid substitution, S228P (Angal, et al. 1993. Mol Immunol 30:105-8.) HC and hK LC format.

NV1 light chain sequence:

EIVLTQSPATLSLSPGERATLSCRASQSVSSYLA^{WY}QQKPGQAPRLLIYDASN^{RAT}GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQSSNWPRTFGQGTKVEIK (SEQ ID NO: 626)

[spacer (SEQ ID NO: 362)][NV1 NV01 2001 (SEQ ID NO: 1296) amino acid sequence:

[QGQSGQG] [RYCHAANPDRFCGIYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRASQSVSSYLA^{WY}QQKPGQAPRLLIYDASN^{RAT}GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1297)

NV1 NV01 2001 amino acid sequence:

RYCHAANPDRFCGIYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRASQSVSSYLA^{WY}QQKPGQAPRLLIYDASN^{RAT}GIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQSSNWPRTFGQGTKVEIK (SEQ ID NO: 1296)

[spacer (SEQ ID NO: 918)] [NV1 NV01 2001 (SEQ ID NO: 1298)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CGGTATTGCCATGCTGCGAATCCTGATCGGTTTTGCGGTA
TTTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA

AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
 AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
 CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
 GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
 AGGTGGAAATCAAG] (SEQ ID NO: 1299)

[spacer (SEQ ID NO: 362)][NV1 NV02 2001 (SEQ ID NO: 1300) amino acid sequence:

[QGQSGQG] [PRVCSTDGGDYCLLPGGGSSSGS ISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
 GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
 PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1301)

NV1 NV02 2001 amino acid sequence:

PRVCSTDGGDYCLLPGGGSSSGS ISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
 SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSELEPEDFAVYYCQ
 QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1300)

[spacer (SEQ ID NO: 918)] [NV1 NV02 2001 (SEQ ID NO: 1302)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CCTAGGGTTTGCTCTACTGATGGTGGTGATTATTGCTTGC
 TTCCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
 AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
 AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
 CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
 GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
 AGGTGGAAATCAAG] (SEQ ID NO: 1303)

[spacer (SEQ ID NO: 362)] [NV1 NV03 2001 (SEQ ID NO: 1304) amino acid sequence:

[QGQSGQG] [PRPQCHHRHNC PDHPGGGSSSGS ISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
 GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
 PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1305)

NV1 NV03 2001 amino acid sequence:

PRPQCHHRHNCPDHPGGGSSGGSSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1304)

[spacer (SEQ ID NO: 918)] [NV1 NV03 2001 (SEQ ID NO: 1306)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CCTCGTCCGCAGTGCCATCATCGGCATAATTGTCCTGATC
ATCCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1307)

[spacer (SEQ ID NO: 362)] [NV1 NV04 2001 (SEQ ID NO: 1308)] amino acid sequence:

[QGQSGQG] [KCSRPAHQNPDRCSRGGGSSGGSSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLE
PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1309)

NV1 NV04 2001 amino acid sequence:

KCSRPAHQNPDRCSRGGGSSGGSSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1308)

[spacer (SEQ ID NO: 918)] [NV1 NV04 2001 (SEQ ID NO: 1310)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [AAGTGCTCGCGGCCTGCTCATCAGAATCCGGATCGTTGCT
CGCGAGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC

CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1311)

[spacer (SEQ ID NO: 362)][NV1 NV05 2001 (SEQ ID NO: 1312) amino acid sequence:

[QGQSGQG] [ASYRCPDYKCSHTKHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
PEDFAVYYCQQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1313)

NV1 NV05 2001 amino acid sequence:

ASYRCPDYKCSHTKHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1312)

[spacer (SEQ ID NO: 918)][NV1 NV05 2001 (SEQ ID NO: 1314)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GCTTCGTATCGGTGCCCTGATTATAAGTGCAGTCATACTA
AGCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1315)

[spacer (SEQ ID NO: 362)][NV1 NV06 2001 (SEQ ID NO: 1316) amino acid sequence:

[QGQSGQG] [LPRCPDHPIKCIETKGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
PEDFAVYYCQQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1317)

NV1 NV06 2001 amino acid sequence:

LPRCPDHPIKCIETKGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1316)

[spacer (SEQ ID NO: 918)] [NV1 NV06 2001 (SEQ ID NO: 1318)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TTGCCGAGGTGCCCCGGATCATCCGATTAAGTGCATTGAGA
CTAAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAATGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1319)

[spacer (SEQ ID NO: 362)] [NV1 NV07 2001 (SEQ ID NO: 1320)] amino acid sequence:

[QGQSGQG] [YTFGCPDRYCDRAATGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSELP
EDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1321)

NV1 NV07 2001 amino acid sequence:

YTFGCPDRYCDRAATGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1320)

[spacer (SEQ ID NO: 918)] [NV1 NV07 2001 (SEQ ID NO: 1322)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TATACGTTTGGTTGCCCTGATAGGTATTGCGATCGTGCGG
CGACGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAATGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC

CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1323)

[spacer (SEQ ID NO: 362)][NV1 NV08 2001 (SEQ ID NO: 1324) amino acid sequence:

[QGQSGQG] [RGCPDFNPPSHCYTAGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1325)

NV1 NV08 2001 amino acid sequence:

RGCPDFNPPSHCYTAGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1324)

[spacer (SEQ ID NO: 918)][NV1 NV08 2001 (SEQ ID NO: 1326)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CGTGGTTGTCCGGATTTAATCCTCCTTCTCATTGCTATA
CTGCTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1327)

[spacer (SEQ ID NO: 362)][NV1 NV09 2001 (SEQ ID NO: 1328) amino acid sequence:

[QGQSGQG] [RDYCGPQSPDYCHEIGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSE
PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1329)

NV1 NV09 2001 amino acid sequence:

RDYCGPQSPDYCHEIGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1328)

[spacer (SEQ ID NO: 918)] [NV1 NV09 2001 (SEQ ID NO: 1330)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [AGGGATTATTGCGGGCCTCAGAGTCCTGATTATTGCCATG
AGATTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAATGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1331)

[spacer (SEQ ID NO: 362)] [NV1 NV10 2001 (SEQ ID NO: 1332)] amino acid sequence:

[QGQSGQG] [PNKPCPDLQCYVTNYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSE
LPEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1333)

NV1 NV10 2001 amino acid sequence:

PNKPCPDLQCYVTNYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRAITGIPARFSGSGSGTDFTLTISSELPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1332)

[spacer (SEQ ID NO: 918)] [NV1 NV10 2001 (SEQ ID NO: 1334)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CCGAATAAGCCTTGCCCGGATCTGCAGTGCTATGTGACGA
ATTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAATGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC

CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1335)

[spacer (SEQ ID NO: 362)][NV1 NV11 2001 (SEQ ID NO: 1336) amino acid sequence:

[QGQSGQG] [PRVACGEPDLCYSNTGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLE
PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1337)

NV1 NV11 2001 amino acid sequence:

PRVACGEPDLCYSNTGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQ
QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1336)

[spacer (SEQ ID NO: 918)][NV1 NV11 2001 (SEQ ID NO: 1338)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CCGCGGGTTGCTTGCGGTGAGCCTGATCTTTGCTATTCTA
ATACTGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCAGCTACCTGGCCTGGTATCAGCAGA
AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
AGGTGGAAATCAAG] (SEQ ID NO: 1339)

[spacer (SEQ ID NO: 362)][NV1 NV12 2001 (SEQ ID NO: 1340) amino acid sequence:

[QGQSGQG] [RGCKKHTISTLTCPDGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLE
PEDFAVYYCQSSNWPRTFGQGTKVEIK] (SEQ ID NO: 1341)

NV1 NV12 2001 amino acid sequence:

RGCKKHTISTLTCPDGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
 SQSVSSYLAWYQQKPGQAPRLLIYDASNRATGIPARFSGSGSGTDFTLTISSLEPEDFAVYYCQ
 QSSNWPRTFGQGTKVEIK (SEQ ID NO: 1340)

[spacer (SEQ ID NO: 918)] [NV1 NV12 2001 (SEQ ID NO: 1342)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CGGGGGTGCAAGAAGCATACTATTTGACGCTTACGTGCC
 CTGATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGAAGTGTCTCCGGCAGATCCGACAA
 TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
 AGAGCCACCCTGAGCTGTAGAGCCAGCCAGAGCGTGTCCAGCTACCTGGCCTGGTATCAGCAGA
 AGCCCGGCCAGGCTCCCCGGCTGCTGATCTACGATGCCAGCAATAGAGCCACCGGCATCCCCGC
 CAGATTTTCCGGCTCTGGCAGCGGCACCGACTTCACCCTGACCATCAGCTCCCTGGAACCCGAG
 GACTTCGCCGTGTACTACTGCCAGCAGAGCAGCAACTGGCCCCGGACATTTGGCCAGGGCACCA
 AGGTGGAAATCAAG] (SEQ ID NO: 1343)

human kappa constant

RTVAAPSVFIFPPSDEQLKSGTASVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDS
 TYSLSSTLTLSKADYEKHKVYACEVTHQGLSSPVTKSFNRGEC (SEQ ID NO: 1344)

human kappa constant

CGTACGGTGGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGAA
 CTGCCTCTGTTGTGTGCCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAGGT
 GGATAACGCCCTCCAATCGGGTAACTCCCAGGAGAGTGTACAGAGCAGGACAGCAAGGACAGC
 ACCTACAGCCTCAGCAGCACCTGACGCTGAGCAAAGCAGACTACGAGAAACACAAAGTCTACG
 CCTGCGAAGTCACCCATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCAACAGGGGAGAGTG
 T (SEQ ID NO: 1345)

NV1 HC variable

QVQLVESGGGVVQPGRSLRLDCKASGITFSNSGMHWVRQAPGKGLEWVAVIWIYDGSKRYYADSV
 KGRFTISRDN SKNTLFLQMNSLRAEDTAVYYCATNDDDYWGQGLVTVSS (SEQ ID
 NO: 1346)

NV1 HC variable

CAGGTGCAGCTGGTGGAACTCTGGCGGCGGAGTGGTGCAGCCTGGCAGAAGCCTGAGACTGGACT
GCAAGGCCAGCGGCATCACCTTCAGCAACAGCGGCATGCACTGGGTGCGCCAGGCCCTGGAAA
AGGCCTGGAATGGGTGGCCGTGATTTGGTACGACGGCAGCAAGCGGTACTACGCCGACAGCGTG
AAGGGCCGGTTCACCATCAGCCGGGACAACAGCAAGAATACCCTGTTCTGCAGATGAACAGCC
TGCGGGCCGAGGACACCGCCGTGTACTACTGCGCCACCAACGACGACTATTGGGGCCAGGGCAC
ACTCGTGACCGTGTCTCT (SEQ ID NO: 1347)

hIgG4 S228P constant

ASTKGPSVFPLAPCSRSTSESTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVLQSSGLYS
LSSVTVTPSSSLGKTYTCNVDPKPSNTKVDKRVESKYGPPCPPCPAPEFLGGPSVFLFPPKPK
DTLMISRTPEVTCVVVDVSQEDPEVQFNWYVDGVEVHNAKTKPREEQFNSTYRVVSVLTVLHQD
WLNGKEYKCKVSNKGLPSSIEKTIKAKGQPREPQVYTLPPSQEEMTKNQVSLTCLVKGFYPSD
IAVEWESNGQPENNYKTTTPVLDSDGSFFLYSRLTVDKSRWQEGNVFSCSVMEALHNHYTQKS
LSLSLGK (SEQ ID NO: 1348)

hIgG4 S228P constant

GCTAGCACCAAGGGCCCATCGGTCTTCCCCCTGGCACCTGTAGCAGAAGCACCAGCGAGTCTA
CAGCGGCCCTGGGCTGCCTGGTCAAGGACTACTTCCCCGAACCGGTGACGGTGTCTGTGGAATC
AGGCGCCCTGACCAGCGGCGTGCACACCTTCCCGGCTGTCTACAGTCCTCAGGACTCTACTCC
CTCAGCAGCGTGGTGACCGTGCCCTCCAGCAGCTTGGGCACCAAGACCTACACCTGTAACGTGG
ACCACAAGCCCAGCAACACCAAGGTGGACAAGCGGGTGGAATCTAAGTACGGCCCTCCCTGCCC
TCCTTGCCCAGCCCCTGAATTTCTGGGGGGACCGTCAGTCTTCTCTTCCCCCAAAACCCAAG
GACACCCTCATGATCTCCCGGACCCCTGAGGTACATGCGTGGTGGTGGACGTGAGCCAGGAAG
ACCCTGAGGTCCAGTTCAACTGGTACGTGGACGGCGTGGAGGTGCATAATGCCAAGACAAAGCC
GCGGGAGGAGCAGTTCAACAGCACGTACCGTGTGGTCAGCGTCCTCACCGTCCTGCACCAGGAC
TGGCTGAATGGCAAGGAGTACAAGTGCAAGGTCTCCAACAAAGGCCTGCCCAGCTCCATCGAGA
AAACCATCTCCAAAGCCAAAGGGCAGCCCCGAGAACCACAGGTGTACACCCTGCCCCCATCCCA
GGAGGAGATGACCAAGAACCAGGTCAGCCTGACCTGCCTGGTCAAAGGCTTCTATCCCAGCGAC
ATCGCCGTGGAGTGGGAGAGCAATGGGCAGCCGGAGAACAACTACAAGACCACGCCTCCCGTGC
TGGACTCCGACGGCTCCTTCTTCTCTACAGCAGACTCACCGTGGACAAGAGCAGGTGGCAGGA

AGGGAACGTCTTCTCATGCTCCGTGATGCATGAGGCTCTGCACAACCACTACACGCAGAAGAGC
CTCTCCCTGTCTCTGGGTAAA (SEQ ID NO: 1349)

EXAMPLE 21: Characterization of activatable anti-PD1 NV1 antibodies

[000890] This example describes activatable anti-PD1 NV1 antibodies with reduced binding to hPD1.

[000891] Masking efficiencies were evaluated by standard plate ELISA. Briefly, human PD1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified NV1 and activatable NV1 antibodies were applied to the plate in serial dilution and allowed to bind. Bound antibody and activatable antibodies were detected with anti-hu Fab-HRP conjugate (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prizm (Sigma Plot). All activatable NV1 antibodies showed decreased binding compared with the parental NV1, as shown in Figure 26.

EXAMPLE 22: Activatable anti-PD1 Pembrolizumab masking moieties

[000892] This example describes identification of masking moieties (MM) that reduce binding of anti-PD1 pembrolizumab antibody to its target.

[000893] Anti-PD1 Pembrolizumab (PM1) (see, e.g., US 8,354,509) was used to screen libraries using a method similar to that described in PCT International Publication Number WO 2010/081173, published 15 July 2010. The screening consisted of one round of MACS and three rounds of FACS sorting. For the initial MACS, approximately 1.6×10^{12} cells were incubated with PM1 antibody at a concentration of 200nM, and approximately 8×10^6 binders were collected using Protein-A Dynabeads (Life Technologies, Carlsbad, CA). FACS rounds were conducted labeling cells with DyLight 650 (Thermo-Fisher) labeled PM1 antibody and collecting cells with strongest fluorescence as follows: 100nM PM1-650 collecting brightest 10% for FACS round 1 (5.6×10^5 for F1), 10nM PM1-650 collecting brightest 1.5% for FACS round 2 (1.4×10^4 for F2), 1nM PM1-650 collecting all binders above background (4,000) and the top 0.2% (820) for FACS round 3 (F3). Individual peptide clones from the two F3 populations were identified by sequence analysis (Table 20), and selected peptide clones PM01-PM12 were subsequently verified for their ability to bind PM1 (masking moiety PM001 is also referred to

herein as PM01 and/or PM-01; masking moiety PM002 is also referred to herein as PM02 and/or PM-02, and so on).

Table 20

Mask	Amino Acid Sequence	SEQ ID NO:
PM01	GCDFTSAKHNCGS GW	1351
PM02	VGSNCWTGPACALTS	1352
PM03	FCAVMFDFLSDRCLH	1353
PM04	FCPPWLDYLGKNCMT	1354
PM05	MSCWDFSSAQGCGQH	1355
PM06	LMCADLHYNHYNCKY	1356
PM07	ELCGWQSFSGVCTSE	1357
PM08	WTYENCWASCQPHLE	1358
PM09	KLTEDFSSAA	1359
PM10	VGQSCFSGLVCDRQL	1360
PM11	ISHYCFSGKSCRD	1361
PM12	HCIPDFTSAAGDCMR	1362
PM13	RLVSAYSFS	1363
PM14	KFHHSHP LVHDFTSA	1364
PM15	ASYPDFSSANGVGLR	1365
PM16	GLATTLSNVDFTSAG	1366
PM17	DFTSANSASFSGDAST	1367
PM18	GRLPGHSVVDFTS AW	1368
PM19	SGSFYSSSAFDF TSA	1369
PM20	CDDFTSAQH SRINEC	1370
PM21	CDFTSAQGKKCR TAL	1371
PM22	YYIDKYQSPSYGPVL	1372
PM23	FSVARARSSADFTSS	1373
PM24	DSDFTSAGSADSR SR	1374
PM25	CDFTSAT SISKRC DH	1375

PM26	IESSASSWGLQASRN	1376
PM27	PRYHNLNFTTPALSPGS	1377
PM28	DLFARFPLDRDFTSA	1378
PM29	HCNFTTPPYCSSTLW	1379
PM30	NVPIILLTDRQLLSG	1380
PM31	NPTACDFTSSMATYC	1381
PM32	FVRTVRFSNSMFSVP	1382
PM33	YDFSSASNSSPSRQT	1383
PM34	AHPDFSSAMRGNLLG	1384
PM35	SSHVVHKDFTSANSR	1385
PM36	CPDFTSANGGGCWQM	1386
PM37	SLGQSYPTDFTCPGC	1387
PM38	ASMRSHEQRDFTSAY	1388
PM39	SCQFWFTLCSGGVFH	1389
PM40	PYPNNRTGMHDFTSA	1390
PM41	KPFPIDFTSAGTSGT	1391
PM42	SIKSFIPRDDFTSAA	1392
PM43	GIKNPATPFVDFTSA	1393
PM44	LSHTYPRGSSTIEAS	1394
PM45	PSLDFSSAT	1395
PM46	AFTPRIAPTFDVMKE	1396
PM47	LCGLQIPDCERS	1397
PM48	AAKMVSHSERDFTSA	1398
PM49	VSVECFSGMQCPHY	1399
PM50	ASKCRLPCMASTQIY	1400
PM51	GLRSCNIYFSIPCTY	1401
PM52	RGTS DGTL DFTTARS	1402
PM53	SMYPSASRLLHPQYP	1403
PM54	HCISCYDFTSAAGSF	1404
PM55	SSGRWGDAWACARIC	1405

PM56	RVFSDFTSASHSFGG	1406
PM57	TDRHSASGRDFTSAH	1407
PM58	AHCEDFSSAERIA TMGC	1408
PM59	ACDPYSFSIPCDDRL	1409
PM60	NSPFTLSHIYDR	1410
PM61	IGTNFTTPSAFVAFP	1411
PM62	RDAFPIYRNADFSTP	1412
PM63	SIPNASSYNFTSSSG	1413
PM64	AGIPDKRHTYDFTSA	1414
PM65	WPLAHDSDWNFTTP	1415
PM66	RHSPSSGHVDFTSAG	1416
PM67	SCFAWTDPVWNRCSW	1417
PM68	MPCDWTGPGKIWCGG	1418
PM69	RDCDFSTANFRSCNK	1419
PM70	LSCVVSPNYLHCNDH	1420
PM71	FVCGLYSFSGVCQGV	1421
PM72	IGLMCFSGLQCPMLA	1422
PM73	PGMNCFSGEICQMST	1423
PM74	GDVGSCWASCGLQGG	1424
PM75	SQFQDCWASC GASFT	1425
PM76	VGSLNCWYSCGDIWL	1426
PM77	MCESWLNFLGDQCGM	1427
PM78	RCMISQSSFSGMCGM	1428
PM79	NCAPWTSNMSNHCLK	1429
PM80	LCGVGSATGLELCGV	1430
PM81	GCDFSSLGGRQPCTP	1431
PM82	MGCNFTTYPYHTCNT	1432
PM83	GSCDFTSGAGKKCGS	1433
PM84	VSCDFTSSHARMCSR	1434
PM85	MRCTDFYYNHTNCIG	1435

PM86	RSCDFTSAANKYCAT	1436
PM87	LYCDSFSVPRPNCAP	1437
PM88	NSCDFTSARVSKCST	1438
PM89	STCSDNFTTPMPCNT	1439
PM90	DICNDRPNLTHCHYF	1440
PM91	LRCDDFTSAIGCRGY	1441
PM92	EGCDFTSALHSCNNY	1442
PM93	RKGC GDFTSASCFVV	1443
PM94	GMLCAGSSFGLCESM	1444
PM95	RESCFGSSLGLCTNK	1445
PM96	ILRCYDIPTNCMNFN	1446
PM97	NSECTFGAMYCRNKP	1447
PM98	ASGCFDEDIRCSGGA	1448
PM99	HYFCNQSNPSCQTAP	1449
PM100	AMGCELGGAGCIGSP	1450
PM101	TLKCHMPRKLCANDP	1451
PM102	RPACRDLPHNCITST	1452
PM103	QMSCHGNFTTCHSNP	1453
PM104	LTGCARGARPCRLRV	1454
PM105	WSELCLAGPSCGWVG	1455
PM106	VTKSCWQLPHCITAP	1456
PM107	KAASCPHNQICNMTA	1457
PM108	VSKNCFSGMMCPVFA	1458
PM109	NRSSCWGTGPTCHVLH	1459
PM110	ARTGCSGPVCLNDVS	1460
PM111	STRTCLAFTCINGNT	1461
PM112	MLDGNCWYACSYKNT	1462
PM113	FSRSDCWSACAPWRV	1463
PM114	GGRMDCWASCQPLSR	1464
PM115	NSPHSCMTNCDFTSA	1465

EXAMPLE 23: Activatable anti-PD1 Pembrolizumab Antibodies

[000894] This example describes examples of activatable anti-PD1 Pembrolizumab antibodies of the disclosure.

[000895] Activatable anti-PD1 PM1 antibodies comprising an anti-PD1 PM1 masking moiety, a cleavable moiety, and an anti-PD1 PM1 antibody of the disclosure were produced according to methods similar to those described in PCT Publication Nos. WO 2009/025846 and WO 2010/081173. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1514. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising a light chain comprising amino acid sequence SEQ ID NO: 638. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1514 and a light chain comprising amino acid sequence SEQ ID NO: 638. One embodiment of the disclosure is an anti-PD-1 activatable antibody comprising the CDRs of an anti-PD-1 activatable antibody comprising a heavy chain comprising amino acid sequence SEQ ID NO: 1514 and a light chain comprising amino acid sequence SEQ ID NO: 638.

[000896] The amino acid and nucleic acid sequences of several activatable anti-PD1 antibody variable domains of the disclosure are provided below. Antibodies were produced as hIgG4 containing a single amino acid substitution, S228P (Angal, et al. 1993. Mol Immunol 30:105-8.) HC and hK LC format.

PM1 light chain amino acid sequence:

EIVLTQSPATLSLSPGERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA⁵SYLESGVPA
RFSGSGSGTDFTLT¹ISSLEPEDFAVYYCQHSRDLPLT¹FGGGTKVEIK (SEQ ID NO: 638)

[spacer (SEQ ID NO: 362)][PM1 PM01 2001 (SEQ ID NO: 1466)] amino acid sequence:

[QGQSGQG] [CDFTSAKHNCGSGWGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGER
ATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA⁵SYLESGVPA⁵RFSGSGSGTDFTLT¹ISSL
EPEDFAVYYCQHSRDLPLT¹FGGGTKVEIK] (SEQ ID NO: 1467)

PM1 PM01 2001 amino acid sequence:

CDFTSAKHNCGSWGSSSGSISGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRASKG
VSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAVYYC
QHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1466)

[spacer (SEQ ID NO: 1126)][PM1 PM01 2001 (SEQ ID NO: 1468)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGA] [TGCGATTTTACTTCTGCCAAGCACAATTGCGGCTCTGGCT
GGGGCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAATCACGGCGG
ATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAAAGAGCCACC
CTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACTGGTATCAGC
AGAAGCCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAGCGGCGTGCC
CGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGCCTGGAACCC
GAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCCTGACATTTGGCGGAGGCA
CCAAGGTGGAAATCAAG] (SEQ ID NO: 1469)

[spacer (SEQ ID NO: 362)][PM1 PM02 2001 (SEQ ID NO: 1470)] amino acid sequence:

[QGQSGQG] [VGSNCWTGPACALTSGGGSSSGSISGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT I
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1471)

PM1 PM02 2001 amino acid sequence:

VGSNCWTGPACALTSGGGSSSGSISGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1470)

[spacer (SEQ ID NO: 918)][PM1 PM02 2001 (SEQ ID NO: 1472)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GTTGGGTCGAATTGCTGGACGGGGCCGGCGTGCGCTTTGA
CGTCGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG

CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1473)

[spacer (SEQ ID NO: 362)][PM1 PM03 2001 (SEQ ID NO: 1474)] amino acid sequence:

[QGQSGQG] [FCAVMFDLSDRCLHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTII
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1475)

PM1 PM03 2001 amino acid sequence:

FCAVMFDLSDRCLHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTIISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1474)

[spacer (SEQ ID NO: 918)][PM1 PM03 2001 (SEQ ID NO: 1476)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TTTTGCGCTGTGATGTTTGATTTTCTGTCTGATCGGTGCC
TGCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1477)

[spacer (SEQ ID NO: 362)][PM1 PM04 2001 (SEQ ID NO: 1478)] amino acid sequence:

[QGQSGQG] [FCPPWLDYLGKNCMTGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTII
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1479)

PM1 PM04 2001 amino acid sequence:

FCPPWLDYLGKNCMTGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1478)

[spacer (SEQ ID NO: 918)][PM1 PM04 2001 (SEQ ID NO: 1480)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TTTTGCCCCGCGTGGTTGGATTATTTGGGTAATAAGTGCA
TGACGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1481)

[spacer (SEQ ID NO: 362)][PM1 PM05 2001 (SEQ ID NO: 1482)] amino acid sequence:

[QGQSGQG] [MSCWDFSSAQGCGQHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRA SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT I
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1483)

PM1 PM05 2001 amino acid sequence:

MSCWDFSSAQGCGQHGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1482)

[spacer (SEQ ID NO: 918)][PM1 PM05 2001 (SEQ ID NO: 1484)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [ATGTCTTGCTGGGATTTTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
AGCATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG

CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1485)

[spacer (SEQ ID NO: 362)][PM1 PM06 2001 (SEQ ID NO: 1486)] amino acid sequence:

[QGQSGQG] [LMCADLHYNHYNCKYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTII
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1487)

PM1 PM06 2001 amino acid sequence:

LMCADLHYNHYNCKYGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTIISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1486)

[spacer (SEQ ID NO: 918)][PM1 PM06 2001 (SEQ ID NO: 1488)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TTGATGTGCGCTGATTTGCATTATAATCATTATAATTGCA
AGTATGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1489)

[spacer (SEQ ID NO: 362)][PM1 PM07 2001 (SEQ ID NO: 1490)] amino acid sequence:

[QGQSGQG] [ELCGWQSFSGVCTSEGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLTII
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1491)

PM1 PM07 2001 amino acid sequence:

ELCGWQSFSGVCTSEGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1490)

[spacer (SEQ ID NO: 918)][PM1 PM07 2001 (SEQ ID NO: 1492)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GAGTTGTGCGGTTGGCAGAGTTTTTCGGGGGTTTGCAC TA
GTGAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCG GACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1493)

[spacer (SEQ ID NO: 362)][PM1 PM08 2001 (SEQ ID NO: 1494)] amino acid sequence:

[QGQSGQG] [WTYENCWASCQPHLEGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRA SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT I
SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1495)

PM1 PM08 2001 amino acid sequence:

WTYENCWASCQPHLEGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1494)

[spacer (SEQ ID NO: 918)][PM1 PM08 2001 (SEQ ID NO: 1496)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [TGGACTTATGAGAATTGCTGGGCTTCGTGCCAGCCTCATT
TGGAGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCG GACTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG

CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1497)

[spacer (SEQ ID NO: 913)][PM1 PM09 2001 (SEQ ID NO: 1498)] amino acid sequence:

[QGQSGQ] [KLTEDFSSAAGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSC
RASKGVSTSGYSYLHWYQQKPGQAPRLLIYLAstyleSGVPARFSGSGSGTDFTLTISSLEPEDF
AVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1499)

PM1 PM09 2001 amino acid sequence:

KLTEDFSSAAGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRASKGVSTS
GYSYLHWYQQKPGQAPRLLIYLAstyleSGVPARFSGSGSGTDFTLTISSLEPEDFAVYYCQHSR
DLPLTFGGGTKVEIK (SEQ ID NO: 1498)

[spacer (SEQ ID NO: 1518)][PM1 PM09 2001 (SEQ ID NO: 1500)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAA] [AAGCTTACTGAGGATTTTCTAGCGCAGCAGGCTCGAGCGGCG
GCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAATCACGGCGGATCCGAGATCGTGCT
GACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAAAGAGCCACCCTGAGCTGTAGAGCC
TCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACTGGTATCAGCAGAAGCCCGGCCAGG
CCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAGCGGCGTGCCCGCCAGATTTTCTGG
CTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGCCTGGAACCCGAGGACTTCGCCGTG
TACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTGGCGGAGGCACCAAGGTGGAAATCA
AG] (SEQ ID NO: 1501)

[spacer (SEQ ID NO: 362)][PM1 PM10 2001 (SEQ ID NO: 1502)] amino acid sequence:

[QGQSGQG] [VGQSCFSGLVCDRLGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSP
GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLAstyleSGVPARFSGSGSGTDFTLT
ISSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1503)

PM1 PM10 2001 amino acid sequence:

VGQSCFSGLVCDRLGGGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
SKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAV
YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1502)

[spacer (SEQ ID NO: 918)][PM1 PM10 2001 (SEQ ID NO: 1504)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [GTGGGGCAGAGTTGCTTTTCTGGGCTGGTTTGCGATAGGC
AGCTGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAA
TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1505)

[spacer (SEQ ID NO: 913)][PM1 PM11 2001 (SEQ ID NO: 1506)] amino acid sequence:

[QGQSGQ] [ISHYCFSGKSCR DGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERAT
LSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEP
EDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1507)

PM1 PM11 2001 amino acid sequence:

ISHYCFSGKSCR DGSSGGSISSGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRASKGV
STSGYSYLHWYQQKPGQAPRLLIYLA SYLESGV PARFSGSGSGTDFTLT ISSLEPEDFAVYYCQ
HSRDLPLTFGGGTKVEIK (SEQ ID NO: 1506)

[spacer (SEQ ID NO: 1518)][PM1 PM11 2001 (SEQ ID NO: 1508)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAA] [ATCTCTCACTATTGTTTCAGTGGCAAATCCTGCAGGGACGGCT
CGAGCGGCGGCTCTATCTCTTCCGGA CTGCTGTCCGGCAGATCCGACAATCACGGCGGATCCGA
GATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAAAGAGCCACCCTGAGC
TGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACTGGTATCAGCAGAAGC
CCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAGCGGCGTGCCCGCCAG

ATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGCCTGGAACCCGAGGAC
 TTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTGGCGGAGGCACCAAGG
 TGGAAATCAAG] (SEQ ID NO: 1509)

[spacer (SEQ ID NO: 362)][PM1 PM12 2001 (SEQ ID NO: 1510)] amino acid sequence:

[QGQSGQG] [HCIPDFTSAAGDCMRGGGSSSGSISGLLSGRSDNHGGSEIVLTQSPATLSLSP
 GERATLSCRASKGVSTSGYSYLHWYQQKPGQAPRLLIYLAstyleSGVPARFSGSGSGTDFTLTII
 SSLEPEDFAVYYCQHSRDLPLTFGGGTKVEIK] (SEQ ID NO: 1511)

PM1 PM12 2001 amino acid sequence:

HCIPDFTSAAGDCMRGGGSSSGSISGLLSGRSDNHGGSEIVLTQSPATLSLSPGERATLSCRA
 SKGVSTSGYSYLHWYQQKPGQAPRLLIYLAstyleSGVPARFSGSGSGTDFTLTIISSLEPEDFAV
 YYCQHSRDLPLTFGGGTKVEIK (SEQ ID NO: 1510)

[spacer (SEQ ID NO: 918)][PM1 PM12 2001 (SEQ ID NO: 1512)] nucleotide sequence:

[CAAGGCCAGTCTGGCCAAGGT] [CATTGCATTCTGATTTTACTTCTGCTGCTGGTGATTGCA
 TGAGGGGAGGTGGCTCGAGCGGCGGCTCTATCTCTTCCGGACTGCTGTCCGGCAGATCCGACAA
 TCACGGCGGATCCGAGATCGTGCTGACACAGAGCCCTGCCACCCTGTCTCTGAGCCCTGGCGAA
 AGAGCCACCCTGAGCTGTAGAGCCTCTAAGGGCGTGTCCACCAGCGGCTACAGCTACCTGCACT
 GGTATCAGCAGAAGCCCGGCCAGGCCCCCAGACTGCTGATCTACCTGGCCTCCTACCTGGAAAG
 CGGCGTGCCCGCCAGATTTTCTGGCTCTGGCAGCGGCACCGACTTCACCCTGACAATCAGCAGC
 CTGGAACCCGAGGACTTCGCCGTGTACTACTGCCAGCACAGCAGGGACCTGCCCCTGACATTTG
 GCGGAGGCACCAAGGTGGAAATCAAG] (SEQ ID NO: 1513)

PM1 HC variable

QVQLVQSGVEVKKPGASVKVSCKASGYTFTNYMYWVRQAPGQGLEWMGGINPSNGGTNFKF
 KNRVTLTDSSTTTAYMELKSLQFDDTAVYYCARRDYRFDMGFYWGQGTITVTVSS (SEQ ID
 NO: 1514)

PM1 HC variable

CAGGTGCAGCTGGTGCAGTCTGGCGTGGAAGTGAAGAAACCAGGCGCCAGCGTGAAGGTGTCCT
 GCAAGGCCAGCGGCTACACCTTTACCAACTACTACATGTACTGGGTGCGCCAGGCCCCAGGCCA
 GGGACTGGAATGGATGGGCGGCATCAACCCCAGCAACGGCGGCACCAACTTCAACGAGAAGTTC
 AAGAACAGAGTGACCCTGACCACCGACAGCAGCACCACCACCGCCTACATGGAAGTGAAGTCCC
 TGCAGTTCGACGACACCGCCGTGTACTACTGCGCCAGACGGGACTACAGATTCGACATGGGCTT
 CGACTACTGGGGCCAGGGCACAACCGTGACCGTGTCTAGT (SEQ ID NO: 1515)

EXAMPLE 24: Characterization of activatable anti-PD1 PM1 antibodies

[000897] This example describes activatable anti-PD1 PM1 antibodies with reduced binding to hPD1.

[000898] Masking efficiencies were evaluated by standard plate ELISA. Briefly, human PD1-Fc (R and D systems, Minneapolis, MN) was adsorbed to the wells of a 96-well ELISA plate. Purified PM1 and activatable PM1 antibodies were applied to the plate in serial dilution and allowed to bind. Bound antibody and activatable antibodies were detected with anti-hu Fab-HRP conjugate (Sigma, St Louis, MO) and visualized with the chromogenic substrate TMB (Thermo Scientific, Rockford, IL). Plots were generated in Prism (Sigma Plot). All activatable PM1 antibodies showed decreased binding compared with the parental PM1, as shown in Figures 27A, 27B, and 27C.

EXAMPLE 25: Characterization of anti-PD1 activatable antibodies of the disclosure

[000899] The binding properties of several masking moieties of the disclosure identified for A1.5 (PD), NV1 (NV) and PM1 (PM) antibodies were evaluated. Eight peptide clones from each mask discovery effort were grown overnight at 37°C, 850rpm in 2mL deep well plates in LB + Chloramphenicol + 0.2% glucose, diluted 1:20 in LB + Chloramphenicol and grown 105 minutes at 37°C, 850rpm. Peptide expression was induced by the addition of 0.04% arabinose for 35 minutes, and clones were stained with 1nM A1.5-DyLight 650, 1nM PM1-DyLight 650 and 1nM NV1-DyLight 650 in cold PBS + 0.5% BSA. Clones were pelleted, the staining solution was removed, and cells were resuspended in PBS + 1% formaldehyde. Fluorescence was measured on a MACSQuant Analyzer 10 flow cytometer manufactured by Miltenyi Biotec, Inc. (San

Diego, CA). As shown in Figure 28, masking moieties bound to the anti-PD1 antibody used in the library screening from which they were identified.

EXAMPLE 26: Evaluation of masking moiety binding by effector negative activatable antibodies

[000900] This example describes additional activatable anti-PD-1 J43 antibodies that exhibit reduced binding to mouse PD-1.

[000901] Examples of additional activatable antibodies of the disclosure comprising an IgG2a effector negative (EN) anti-PD1 antibody J43 (J43 m2a EN) and a variety of mask and substrate combinations were produced using techniques as described herein. The amino acid and nucleic acid sequences of the constant region of the IgG2a EN antibody are provided below. A plasmid encoding this effector negative Fc region, pFUSE-mIgG2Ae1-Fc, is available from InvivoGen, San Diego, CA. The Fc region of mIgG2a was engineered by mutating the following amino acids to reduce binding to FcR and C1q: L235E and E318A/K320A/K322A.

> mIgG2a EN

AKTTAPSVYPLAPVCGD TTGSSVTLGCLVKGYFPEPVTLTWNSGSLSSGVHTFPAVLQSDLYTL
SSSVTVTSSTWPSQSITCNVAHPASSTKVDKKIEPRGPTIKPCPPCKCPAPNLEGGPSVFI FPP
KIKDVLMI SLSPIVTCVVVDVSEDDPDVQISW FVNNVEVHTAQTQTHREDYNSTLRVVSALPIQ
HQDWM SGKAFACAVNNKDLPAPIERTISKPKGSVRAPQVYVLPPEEEMTKKQVTLT CMVTDFM
PEDIYVEWTNNGKTELNYKNTEPVLDSDGSYFMYSKLRVEKKNWVERNSYSCSVVHEGLHNHHT
TKSFSRTPGK (SEQ ID NO: 1516)

> mIgG2a EN

GCCAAGACAACAGCCCCAGCGTGTACCCTCTGGCCCCTGTGTGTGGCGATACCACAGGCAGCT
CTGTGACCCTGGGCTGCCTCGTGAAGGGCTACTTCCCTGAGCCAGTGACCCTGACCTGGAACAG
CGGCTCTCTGTCTAGCGGCGTGCACACCTTTCCAGCCGTGCTGCAGAGCGACCTGTACACCCTG
AGCAGCAGCGTGACCGTGACCAGCAGCACATGGCCCAGCCAGAGCATCACCTGTAACGTGGCCC
ACCCTGCCAGCTCCACCAAGGTGGACAAGAAGATCGAGCCCAGAGGCCCCACCATCAAGCCTTG
CCCCCTTGCAAATGCCCTGCCCCCAATCTGGAAGGCGGCCCTAGCGTGTTTCATCTTCCCACCC
AAGATCAAGGACGTGCTGATGATCAGCCTGAGCCCCATCGTGACCTGCGTGGTGGTGGACGTGT

CCGAGGACGACCCCGATGTGCAGATCAGTTGGTTCGTGAACAACGTGGAAGTGCACACCGCCCA
 GACCCAGACACACAGAGAGGACTACAACAGCACCCCTGAGAGTGGTGTCCGCCCTGCCCATCCAG
 CACCAGGATTGGATGAGCGGCAAGGCCTTCGCCTGCGCTGTGAACAACAAGGACCTGCCAGCCC
 CCATCGAGCGGACCATCTCTAAGCCTAAGGGCAGCGTGCGGGCTCCCCAGGTGTACGTGCTGCC
 TCCTCCAGAGGAAGAGATGACCAAGAAACAAGTGACACTGACATGCATGGTCACCGACTTCATG
 CCCGAGGACATCTACGTGGAATGGACCAACAACGGCAAGACCGAGCTGAACTACAAGAACACCG
 AGCCCGTGCTGGACAGCGACGGCAGCTACTTCATGTACAGCAAGCTGCGGGTGGAAAAGAAAAA
 CTGGGTGGAACGGAACAGCTACAGCTGCAGCGTGGTGCACGAGGGCCTGCACAATCACCACACC
 ACCAAGAGCTTCAGCCGGACCCCTGGAAAA (SEQ ID NO: 1517)

[000902] Masking efficiencies of several of these activatable antibodies were determined as described herein. The results are shown in Figure 29A and Figure 29B.

J43m2a EN Activatable Antibody LCs:

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2001 (SEQ ID NO: 1808)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDF
 YPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEK
 SLSPAEL] (SEQ ID NO: 1809)

J43m2a EN MP8-2 2001 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNHGGGSYELTQPPSASVNVG
 ETVKITCSGDQLPKYFADWFHQRSQDTILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRA
 EDEGDYYCFSGYVDSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDF
 YPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEK
 SLSPAEL (SEQ ID NO: 1808)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2011 (SEQ ID NO: 1810)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNPGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1811)

J43m2a EN MP8-2 2011 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNPGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1810)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2012 (SEQ ID NO: 1812)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANPGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1813)

J43m2a EN MP8-2 2012 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANPGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1812)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2002 (SEQ ID NO: 1814)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSGNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVND FYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPA ECL] (SEQ ID NO: 1815)

J43m2a EN MP8-2 2002 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSGNHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVND FYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPA ECL (SEQ ID NO: 1814)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2003 (SEQ ID NO: 1816)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANPRGGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVND FYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPA ECL] (SEQ ID NO: 1817)

J43m2a EN MP8-2 2003 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANPRGGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVND FYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPA ECL (SEQ ID NO: 1816)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2006 (SEQ ID NO: 1818)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDDHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1819)

J43m2a EN MP8-2 2006 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDDHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1818)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2007 (SEQ ID NO: 1820)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDIHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1821)

J43m2a EN MP8-2 2007 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDIHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1820)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2008 (SEQ ID NO: 1822)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDQHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1823)

J43m2a EN MP8-2 2008 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDQHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1822)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2009 (SEQ ID NO: 1824)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDTHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1825)

J43m2a EN MP8-2 2009 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDTHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1824)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2010 (SEQ ID NO: 1826)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDYHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1827)

J43m2a EN MP8-2 2010 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDYHGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1826)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2013 (SEQ ID NO: 1828)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANIGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1829)

J43m2a EN MP8-2 2013 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSANIGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1828)

[spacer (SEQ ID NO: 362)] [J43m2a EN MP8-2 2014 (SEQ ID NO: 1830)] amino acid sequence:

[QGQSGQG] [ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNIGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L] (SEQ ID NO: 1831)

J43m2a EN MP8-2 2014 amino acid sequence:

ACRICQDHPATKWNSGGGSSGGSSISSGLLSGRSDNIGGGSYELTQPPSASVNVGETVKITCSGDQLPKYFADWFHQRS DQ TILQVIYDDNKRPSGIPERISGSSSGTTATLTIRDVRAEDEGDYYCFSGYVDS DSKLYVFGSGTQLTVLGGPKSSPKVTVFPPSPEELRTNKATLVCLVNDFYPGSATVTWKANGATINDGVKTTKPSKQGQNYMTSSYLSLTADQWKSHNRVSCQVTHEGETVEKSLSPAEC L (SEQ ID NO: 1830)

EXAMPLE 27: Activatable anti-mouse PD1 J43 m2a EN antibodies reduce incidence of diabetes in NOD mice

[000903] In this Example, anti-PD1 J43 m2a EN (effector negative) activatable antibodies were analyzed for the ability to protect from PD1 m2a EN-mediated induction of diabetes in NOD mice.

[000904] The NOD mice, substrain NOD/ShiLtJ, were obtained from Jackson Laboratory at 8 weeks and acclimated on site for 2 weeks. At 10 weeks, mice were checked for diabetes prior to enrollment, grouped, and dosed as set forth in Table 21.

Group	Count	Gender	Treatment	Dose (mg/kg)	Dose volume (mL/kg)	Schedule	Route
1	7	F	mIgG2a (C1.18.4)	10	10	q7dx1	IP
2	7	F	Anti-PD-1 J43 m2a EN	10	10	q7dx1	IP
3	7	F	Anti-PD-1 J43 m2a EN	3	10	q7dx1	IP
4	7	F	Anti-PD-1 J43 m2a EN	1	10	q7dx1	IP
5	7	F	J43 MP7-1 2001 m2a EN	10	10	q7dx1	IP

6	7	F	J43 MP7-1 2001 m2a EN	3	10	q7dx1	IP
7	7	F	J43 MP8-2 2001 m2a EN	10	10	q7dx1	IP
8	7	F	J43 MP8-2 2001 m2a EN	3	10	q7dx1	IP

[000905] Figure 30 which plots % non-diabetic versus number of days post dose, shows that anti-PD-1 J43 antibody induced diabetes in NOD mice with increased frequency as dosage increased. At day eleven post dose, the percentage of non-diabetic mice for the antibody-treated groups was 29%, 43% and 71% for the 10mg/kg, 3mg/kg and 1mg/kg dose groups, respectively. Activatable antibodies J43 MP7-1 2001 m2a EN and J43 MP8-2 2001 m2a EN required increased doses to induce diabetes at frequencies comparable to the parental antibody. At day eleven post dose with J43 MP7-1 2001 m2a EN, 71% of the 10mg/kg group remained non-diabetic and 86% of the 3mg/kg were non-diabetic. At day fourteen post dose with J43 MP8-2 2001 m2a EN, 86% of the 10 mg/kg group and 100% of the 3mg/kg group remained non-diabetic.

EXAMPLE 28. Activatable anti-mouse PD1 J43 m2a EN antibodies show efficacy in the MC38 syngeneic tumor model.

[000906] This Example demonstrates that activatable antibodies of the embodiments are able to reduce the growth of MC38 syngeneic tumors.

[000907] In this Example, anti-PD1 activatable antibodies J43 MP7-1 2001 m2a EN and J43 MP8-2 2001 m2a EN were analyzed for the ability to reduce the growth of MC38 syngeneic tumors.

[000908] The mouse colon carcinoma cell line MC38 was obtained from ATCC. MC38 were grown in RPMI-1640 supplemented with 10% fetal bovine serum at 37°C in an atmosphere of 5% CO₂ in air. Cells were harvested during the logarithmic growth period, resuspended in PBS with proper cell concentration, and kept on ice for tumor induction.

[000909] Each mouse was inoculated subcutaneously at the right flank with 1.5×10^6 of MC38 cells in PBS for tumor development. The treatments were started when the mean tumor size reached approximately 80 mm³ (60-120 mm³). Tumor sizes were measured twice weekly in two dimensions using a caliper, and the volume was expressed in mm³ using the formula: $V = 0.5 a \times b^2$ where a and b are the long and short diameters of the tumor, respectively.

[000910] The mice were grouped and dosed as set forth in Table 22.

Table 22:

Group	Count	Treatment	Dose (mg/kg)	Dose volume (mL/kg)	Schedule	Route
1	10	mIgG1 MOPC-21 + mIgG12b MPC-11	10 + 10	10	b.i.w. for 3 weeks	IP
2	10	Anti-CTLA4 9D9 mIgG2b	10	10	b.i.w. for 3 weeks	IP
3	10	Anti-PD-1 (J43 m2a EN)	10	10	b.i.w. for 3 weeks	IP
4	10	J43 MP8-2 2001 m2a EN	10	10	b.i.w. for 3 weeks	IP
5	10	J43 MP7-1 2001 m2a EN	10	10	b.i.w. for 3 weeks	IP
6	10	Anti-PD-1 (J43 m2a EN) + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP
7	10	J43 MP8-2 2001 m2a EN + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP
8	10	J43 MP7-1 2001 m2a EN + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP

[000911] Figures 31A and 31B, which plots tumor volume versus number of days post initial dose, demonstrates that anti-PD1 activatable antibodies MP7-1 2001 m2a EN and MP8-2 2001 m2a EN inhibit the growth of MC38 syngeneic tumors similar to positive control anti-PD1 antibody J43 m2a EN both as single agents and in combination with anti-CTLA4 antibody 9D9 mIgG2b (BioXCell, West Lebanon, NH).

EXAMPLE 29. Evaluation of anti-human PD1 antibodies and activatable anti-human PD1 antibodies in antigen recall assay.

[000912] This Example demonstrates that anti-PD1 antibodies and activated activatable anti-PD1 antibodies of the embodiments (i.e., activatable antibodies in which the CM has been

cleaved by a protease) are able to block PD-L1/PD-L2 binding to PD1 and to activate T cells in an antigen recall assay.

[000913] As shown in Figures 32A-32E, the anti-human PD1 antibody referred to herein as A1.5 Ab (i.e., VH of SEQ ID NO: 21 and VL of SEQ ID NO: 47) blocks PD-L1/PD-L2 binding to PD1 and potently activates T cells in an antigen recall assay. The binding of the anti-human PD-1 A1.5 Ab to immobilized human PD1 as detected by standard plate ELISA is shown in Figure 32A, and the binding of A1.5 Ab to cynomolgus PD1 (also referred to herein as cyno-PD1, Sino Biological Cat. #90311-C02H) as detected by ELISA is shown in Figure 32B. The inhibition of biotinylated human PD-L1 (also referred to herein as biotin-PD-L1) to immobilized PD1 by A1.5 Ab as determined by ELISA is shown in Figure 32C, and the inhibition of biotinylated human PD-L2 (also referred to herein as biotin-PD-L2) to immobilized PD1 as determined by ELISA is shown in Figure 32D.

[000914] As shown in Figure 32E, the A1.5 Ab enhances IFN- γ production in a CMV T cell restimulation assay. Briefly, CMV⁺ human PBMCs were plated at 250,000 cells/well with isotype control antibody or with the A1.5 Ab and stimulated with 5 μ g/mL CMV lysate for 4 days. Supernatant IFN- γ levels were measured by ELISA.

[000915] As shown in Figures 33A and 33B, the activatable anti-PD1 antibody referred to herein as A1.5 PD34 2001 (i.e., VH of SEQ ID NO: 21, VL of SEQ ID NO: 47, masking moiety of SEQ ID NO: 99, and cleavable moiety of SEQ ID NO: 214) binds human PD1 with decreased affinity relative to the parental PD-1 Ab, i.e., A1.5 Ab (Figure 33A), and the A1.5 PD34 2001 activatable antibody shows functional masking in a CMV T cell antigen recall assay (Figure 33B). Briefly, 4mg/mL of the A1.5 PD34 2001 activatable antibody was combined with 0 μ g/mL or 60 μ g/mL recombinant human urokinase (rh uPA, R&D, Cat. #1310-SE) and incubated at 37°C overnight. Binding of the activatable antibody incubated with uPA to hPD1 was assayed by standard plate ELISA. As shown in Figure 33, the activatable anti-PD1 antibodies of the disclosure regain full binding activated after activation by a protease.

EXAMPLE 30. Pharmacokinetic evaluation of activatable anti-human PD1 antibodies in non-human primate model.

[000916] This Example demonstrates pharmacokinetic and dose proportionality data for anti-PD-1 antibodies and anti-PD-1 activatable antibodies of the embodiments.

[000917] Briefly, female cynomolgus monkeys (n=2/group) were dosed with a single IV bolus dose of either the A1.5 antibody or the activatable antibody referred to herein as A1.5 PD34 2001 at 1 mg/kg or at 5 mg/kg. Plasma samples were analyzed for A1.5 and A1.5 PD34 2001 concentrations by a qualified anti-human sandwich ELISA. The results are shown in Figure 34, where each line represents one individual. Mean PK parameters shown for both A1.5 and A1.5 PD34 2001. Similar results were seen for activatable antibodies A1.5 PD34 2012 and A1.5 PD34 2011 administered to cynomolgus monkeys in a similar study.

EXAMPLE 31. Activatable anti-mouse PD-1 J43 antibodies reduce incidence of diabetes in NOD mice dosed with anti-CTLA4 antibody

[000918] In this Example, anti-PD-1 J43 activatable antibodies were analyzed for the ability to protect from anti-PD-1 induction of diabetes in NOD mice when dosed concurrently with anti-CTLA-4 9D9 mIgG2b antibody (BioXCell cat #BE0164). The NOD mice, substrain NOD/ShiLtJ, were obtained from Jackson Laboratory at 4 weeks and acclimated on site for 1 week. At 5 weeks, mice were checked for diabetes prior to enrollment, grouped, and dosed as set forth in Table 23. As used herein, the antibody referred to as “J43 m2a EN” (and variations thereof) comprises a heavy chain (HC) of SEQ ID NO: 546; a light chain (LC) of SEQ ID NO: 543; and a constant region of SEQ ID NO: 1516. As used herein, the activatable antibody referred to as “J43 MP8-2 2012 m2a EN” is an activatable antibody comprising the J43 m2a EN antibody, the MP8-2 masking moiety (SEQ ID NO: 549), and the 2012 substrate (SEQ ID NO: 1101), and the activatable antibody referred to as “J43 MP8-2 2001 m2a EN” is an activatable antibody comprising the J43 m2a EN antibody, the MP8-2 masking moiety (SEQ ID NO: 549), and the 2001 substrate (SEQ ID NO: 214).

Table 23. Dosing regimen

Group	Count	Treatment				Dose volume (mL/kg)	Schedule	Route
		Article #1	Dose (mg/kg)	Article #2	Dose (mg/kg)			
1	8	mIgG1 MOPC-21	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP

2	8	Anti-CTLA4 9D9 mIgG2b	10	mIgG1 MOPC-21	10	10	d0, d4, d7	IP
3	8	Anti-PD-1 (J43 m2a EN)	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP
4	8	Anti-PD-1 (J43 m2a EN)	1	mIgG2b MPC-11	10	10	d0, d4, d7	IP
5	8	Anti-PD-1 (J43 m2a EN)	10	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP
6	8	Anti-PD-1 (J43 m2a EN)	1	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP
7	8	J43 MP8-2 2012 m2a EN	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP
8	8	J43 MP8-2 2012 m2a EN	10	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP
9	8	J43 MP8-2 2001 m2a EN	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP
10	8	J43 MP8-2 2001 m2a EN	10	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP

[000919] Figure 35 which plots % non-diabetic versus number of days post initial dose, shows that the combination of 10 mg/kg anti-PD-1 J43 antibody plus 10 mg/kg anti-CTLA-4 antibody dosed on days 0, 4 and 7 induced diabetes in 50% of NOD mice by day eleven while the same dosing schedule of activatable anti-PD-1 J43 antibodies with CTLA-4 antibody resulted in no induction of diabetes to day eighteen.

EXAMPLE 32. Activatable anti-mouse PD1 J43 m2a EN antibodies show enhanced efficacy when coadministered with anti-CTLA4 antibody in the MC38 syngeneic tumor model.

[000920] This Example demonstrates that activatable antibodies of the embodiments are able to reduce the growth of MC38 syngeneic tumors as single agents and induce MC38 tumor regression when coadministered with anti-CTLA4 antibody 9D9 mIgG2b.

[000921] In this Example, anti-PD1 activatable antibodies J43 MP8-2 2011 m2a EN and J43 MP8-2 2012 m2a EN were analyzed for the ability to slow the growth of or, in combination with anti-CTLA4 antibody, induce regression in MC38 syngeneic tumors. As used herein, the antibody referred to as “J43 m2a EN” (and variations thereof) comprises a heavy chain (HC) of SEQ ID NO: 546; a light chain (LC) of SEQ ID NO: 543; and a constant region of SEQ ID NO: 1516. As used herein, the activatable antibody referred to as “J43 MP8-2 2011 m2a EN” is an activatable antibody comprising the J43 m2a EN antibody, the MP8-2 masking moiety (SEQ ID NO: 549), and the 2011 substrate (SEQ ID NO: 1100), and the activatable antibody referred to as “J43 MP8-2 2012 m2a EN” is an activatable antibody comprising the J43 m2a EN antibody, the MP8-2 masking moiety (SEQ ID NO: 549), and the 2012 substrate (SEQ ID NO: 1101).

[000922] The mouse colon carcinoma cell line MC38 was obtained from ATCC. MC38 cells were grown in RPMI-1640 supplemented with 10% fetal bovine serum at 37°C in an atmosphere of 5% CO₂ in air. Cells were harvested during the logarithmic growth period, resuspended in PBS with proper cell concentration, and kept on ice for tumor induction.

[000923] Each mouse was inoculated subcutaneously at the right flank with 1.5×10^6 MC38 cells in PBS for tumor development. The treatments were started when the mean tumor size reached approximately 60 mm³ (45-80 mm³). Tumor sizes were measured twice weekly in two dimensions using a caliper, and the volume was expressed in mm³ using the formula: $V = 0.5 \times a \times b^2$ where a and b are the long and short diameters of the tumor, respectively.

[000924] The mice were grouped and dosed as set forth in Table 24.

Table 24:

Group	Count	Treatment	Dose (mg/kg)	Dose volume (mL/kg)	Schedule	Route
1	10	mIgG1 MOPC-21 + mIgG12b MPC-11	10 + 10	10	b.i.w. for 3 weeks	IP
2	10	Anti-CTLA4 9D9 mIgG2b	10	10	b.i.w. for 3 weeks	IP

3	10	Anti-PD-1 (J43 m2a EN)	10	10	b.i.w. for 3 weeks	IP
4	10	J43 MP8-2 2012 m2a EN	10	10	b.i.w. for 3 weeks	IP
5	10	J43 MP8-2 2011 m2a EN	10	10	b.i.w. for 3 weeks	IP
6	10	Anti-PD-1 (J43 m2a EN) + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP
7	10	J43 MP8-2 2012 m2a EN + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP
8	10	J43 MP8-2 2011 m2a EN + Anti-CTLA4 9D9 mIgG2b	10+10	10	b.i.w. for 3 weeks	IP

[000925] Figure 36, which plots tumor volume versus number of days post initial dose, demonstrates that anti-PD1 activatable antibodies MP8-2 2012 m2a EN and MP8-2 2011 m2a EN inhibit the growth of MC38 syngeneic tumors similar to positive control anti-PD1 antibody J43 m2a EN both as single agents and in combination with anti-CTLA4 antibody 9D9 mIgG2b (BioXCell, West Lebanon, NH).

EXAMPLE 33. Activatable anti-mouse PD1 J43 m2a EN antibodies show durable anti-tumor activity when coadministered with anti-CTLA4 antibody in the MC38 syngeneic tumor model.

[000926] This Example demonstrates anti-PD-1 activatable antibodies of the embodiments in combination with anti-CTLA4 antibodies induce durable anti-tumor complete responses in the MC38 syngeneic tumor model.

[000927] In this Example, all animals treated with combinations of anti-CTLA4 and anti-PD1 agents from Example 28 that showed sustained tumor regression ((i) anti-PD-1 J43 m2aEN + anti-CTLA4 9D9 mIgG2b (referred to in Figure 37 as “anti-PD-1 J43 m2aEN Antibody” + “anti-CTLA4 9D9 mIgG2b Antibody”) (n=8), (ii) activatable antibody anti-PD-1 J43 MP8-2 2012 m2a EN + anti-CTLA4 9D9 mIgG2b (referred to in Figure 37 as “J43 MP8-2 2012 m2a EN AA” + “anti-CTLA4 9D9 mIgG2b Antibody” where AA stands for activatable antibody) (n=8), and activatable antibody anti-PD-1 J43 MP8-2 2011 m2a EN + anti-CTLA4 9D9 mIgG2b (referred to in Figure 37 as “J43 MP8-2 2011 m2a EN AA” + “anti-CTLA4 9D9 mIgG2b Antibody” where AA stands for activatable antibody) (n=6)) were implanted at day 38 with 1.5 x

10^6 MC38 tumor cells in the left flank opposite to the original MC38 implantation. Five untreated mice were implanted with 1.5×10^6 MC38 tumor cells to confirm tumorigenic activity of the MC38 cells.

[000928] Figure 37 shows that no mice previously treated with combinations of anti-PD1 and anti-CTLA4 agents showed regrowth of the original tumor sites (right flank) or growth of new tumors (left flank) while all five of the untreated control mice showed rapid tumor growth.

EXAMPLE 34. Activatable anti-mouse PD-1 J43 antibodies reduce incidence of diabetes in NOD mice co-dosed with anti-CTLA4 antibody

[000929] This Example demonstrates that anti-PD-1 activatable antibodies of the embodiments in combination with anti-CTLA4 antibodies protected NOD mice from PD1 m2a EN-mediated induction of diabetes.

[000930] In this Example, anti-PD-1 J43 activatable antibody MP8-2 2011 m2aEN was analyzed for the ability to protect from anti-PD-1 induction of diabetes in NOD mice when dosed concurrently with anti-CTLA-4 9D9 mIgG2b antibody (BioXcell cat #BE0164). The NOD mice, substrain NOD/ShiLtJ, were obtained from Jackson Laboratory at 4 weeks and acclimated on site for 1 week. At 5 weeks, mice were checked for diabetes prior to enrollment, grouped, and dosed as set forth in Table 25.

Table 25. Dosing regimen

Group	Count	Treatment					Schedule	Route
		Article #1	Dose (mg/kg)	Article #2	Dose (mg/kg)	Dose volume (mL/kg)		
1	8	Anti-CTLA4 9D9 mIgG2b	10	mIgG1 MOPC-21	10	10	d0, d4, d7	IP
2	8	Anti-PD-1 (J43 m2a EN)	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP
3	8	Anti-PD-1 (J43 m2a EN)	10	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP
4	8	J43 MP8-2 2011 m2a EN	10	mIgG2b MPC-11	10	10	d0, d4, d7	IP
5	8	J43 MP8-2 2011 m2a EN	10	Anti-CTLA4 9D9 mIgG2b	10	10	d0, d4, d7	IP

[000931] Figure 38, which plots % non-diabetic versus number of days post initial dose, shows that the single agent anti-PD-1 J43 antibody (referred to in the figure as “anti-PD1 J43

Ab”) and the combination of 10 mg/kg anti-PD-1 J43 antibody plus 10 mg/kg anti-CTLA4 antibody (referred to in the figure as “anti-PD1 J43 Ab” + “anti-CTLA4 9D9 mIgG2b”) dosed on days 0, 4 and 7 induced diabetes in 25% of NOD mice by day nine. In contrast, the same dosing schedule of activatable antibody anti-PD-1 J43 MP8-2 2011 m2a EN (referred to in the figure as “J43 MP8-2 2011 m2a EN AA” where AA stands for activatable antibody) as a single agent or in combination with anti-CTLA4 antibody (referred to in the figure as “J43 MP8-2 2011 m2a EN AA + “anti-CTLA4 9D9 mIgG2b” where AA stands for activatable antibody) resulted in no induction of diabetes to day fifteen.

EXAMPLE 35. Anti-PD-1 nivolumab activatable antibodies of the embodiments are functionally masked in a human T-cell restimulation assay

[000932] This Example describes the effect of masking moieties on the biological function of the anti-PD-1 nivolumab antibody.

[000933] PBMCs from a CMV-positive donor (Hemacare) were plated at 2×10^5 cells/well in the presence of CMV viral lysate (Astarte) and either anti-PD-1 antibody A1.5, anti-PD-1 activatable antibody A1.5 PD34 2001, anti-PD-1 antibody nivolumab (NV1), anti-PD-1 nivolumab activatable antibody NV1 NV07 2001, or an hIgG4 isotype control antibody. After four days, supernatant was removed from each well and IFN-gamma levels were assayed using an IFN-gamma ELISA kit (Life Technologies, Carlsbad, CA).

[000934] Figure 39 demonstrates that the anti-PD-1 activatable antibody A1.5 PD34 2001 (referred to in the figure as “A1.5 PD34 2001 AA” where AA stands for activatable antibody) and anti-PD1 nivolumab activatable antibody NV1 NV07 2001 (referred to in the figure as “NV1 NV07 2001 AA” where AA stands for activatable antibody) effected increased CMV-stimulated IFN-gamma secretion compared to the control hIgG4 antibody (referred to in the figure as hIgG4) but decreased potency relative to anti-PD-1 parental antibody A1.5 (referred to in the figure as A1.5) or anti-PD-1 parental antibody nivolumab (referred to in the figure as NV1).

EXAMPLE 36. Anti-PD-1 pembrolizumab activatable antibodies of the embodiments are functionally masked in a human T-cell restimulation assay

[000935] This example describes the effect of masking moieties on the biological function of the anti-PD-1 pembrolizumab antibody.

[000936] PBMCs from a CMV-positive donor (Hemacare) were plated at 2×10^5 cells/well in the presence of 4 ug/ml CMV viral lysate (Astarte) and either anti-PD-1 antibody A1.5, anti-PD-1 antibody pembrolizumab (PM1), anti-PD-1 pembrolizumab activatable antibody PM1 PM07 2001, or an hIgG4 isotype control antibody. After four days, supernatant was removed from each well and IFN-gamma levels were assayed using an IFN-gamma ELISA kit (Life Technologies, Carlsbad, CA).

[000937] Figure 40 demonstrates that anti-PD1 pembrolizumab activatable antibody PM1 PM07 2001 (referred to in the figure as “PM1 PM07 2001 AA” where AA stands for activatable antibody) effected increased CMV-stimulated IFN-gamma secretion compared to the control hIgG4 antibody (referred to in the figure as hIgG4) but decreased potency relative to anti-PD-1 parental antibody A1.5 (referred to in the figure as A1.5) or anti-PD-1 parental antibody pembrolizumab (referred to in the figure as PM1).

Other Embodiments

[000938] While the invention has been described in conjunction with the detailed description thereof, the foregoing description is intended to illustrate and not limit the scope of the invention, which is defined by the scope of the appended claims. Other aspects, advantages, and modifications are within the scope of the following.

CLAIMS

1. An isolated antibody or antigen binding fragment thereof (AB) that specifically binds to a human PD-1, wherein the AB comprises
 - a VH CDR1 amino acid sequence comprising GFTFSGYAMS (SEQ ID NO: 653),
 - a VH CDR2 amino acid sequence comprising YISNSGGNAH (SEQ ID NO: 658),
 - a VH CDR3 amino acid sequence comprising EDYGTSPFVY (SEQ ID NO: 664),
 - a VL CDR1 amino acid sequence comprising an amino acid sequence selected from the group consisting of
 - RASESVDSYGISFMN (SEQ ID NO: 675), and
 - RASESVDAYGISFMN (SEQ ID NO: 676)
 - a VL CDR2 amino acid sequence comprising AASNQGS (SEQ ID NO: 678), and
 - a VL CDR3 amino acid sequence comprising QQSKDVPWT (SEQ ID NO: 683).
2. The antibody of claim 1, wherein the AB blocks the ability of a human PDL1 or a human PDL2 to bind to human PD1 with an EC₅₀ of
 - 0.1 nM to 10 nM, 0.1 nM to 5 nM, 0.1 nM to 3 nM, 0.1 nM to 2 nM, 0.1 nM to 1 nM, 0.1 nM to 0.5 nM, 0.1 nM to 0.25 nM,
 - 0.25 nM to 10 nM, 0.25 nM to 5 nM, 0.25 nM to 3 nM, 0.25 nM to 2 nM, 0.25 nM to 1 nM, 0.25 nM to 0.5 nM,
 - 0.5 nM to 10 nM, 0.5 nM to 5 nM, 0.5 nM to 3 nM, 0.5 nM to 2 nM, 0.5 nM to 1 nM,
 - 1 nM to 10 nM, 1 nM to 5 nM, 1 nM to 3 nM, 1 nM to 2 nM,
 - 2 nM to 10 nM, 2 nM to 5 nM, 2 nM to 3 nM,
 - 3 nM to 10 nM, 3 nM to 5 nM, or
 - 5 nM to 10 nM.

3. The antibody of claim 1 or 2, wherein the antigen binding fragment thereof is selected from the group consisting of a Fab fragment, a F(ab')₂ fragment, a scFv, a scAb, a dAb, a single domain heavy chain antibody, and a single domain light chain antibody.

4. A conjugated antibody comprising the antibody of any one of claims 1 to 3 conjugated to an agent.

5. The conjugated antibody of claim 4, wherein the agent has one or more of the following characteristics selected from the group consisting of:

- (a) the agent is conjugated to the antibody via a linker;
- (b) the agent is conjugated to the antibody via a cleavable linker;
- (c) the agent is conjugated to the antibody via a non-cleavable linker;
- (d) the agent is a detectable moiety; and
- (e) the agent is a diagnostic agent.

6. A pharmaceutical composition comprising the antibody of any one of claims 1 to 3, or the conjugated antibody of claim 4 or 5 and a carrier.

7. A pharmaceutical composition comprising the antibody of any one of claims 1 to 3.

8. The pharmaceutical composition of claim 6 comprising an additional agent.

9. The pharmaceutical composition of claim 7 comprising an additional agent.

10. The pharmaceutical composition of claim 8, wherein the additional agent is a therapeutic agent.

11. The pharmaceutical composition of claim 9, wherein the additional agent is a therapeutic agent.

12. An isolated nucleic acid molecule encoding the antibody of any one of claims 1 to 3.

13. A vector comprising the isolated nucleic acid molecule of claim 12.

14. Use of an effective amount of the antibody of any one of claims 1 to 3, or the pharmaceutical composition of claims 7, 9, or 10 in the manufacture of a medicament for reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells, wherein the medicament is to be administered to a subject in need thereof.

15. Use of a therapeutically effective amount of the isolated antibody as in claim 1, or the pharmaceutical composition as in any of claims 7, 9, or 10 in the manufacture of a medicament for reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells, wherein the medicament is to be administered to a subject in need thereof.

16. Use of a therapeutically effective amount of the antibody of any one of claims 1 to 3 or the conjugated antibody of claim 4 or 5 or the pharmaceutical composition of any one of claims 7, 9, or 10 in the manufacture of a medicament for reducing immune suppression, wherein the medicament is to be administered to a subject in need thereof.

17. The use of claim 16, wherein the immune suppression is mediated by engagement of PD-1 on T cells to PDL1 or PDL2 on tumor or other immune cells.

18. The use of any one of claims 14 to 17, wherein the use comprises administering an additional agent, wherein the additional agent is a therapeutic agent.

19. An effective amount of the antibody of any one of claims 1 to 3, or the pharmaceutical composition of claims 7, 9, or 10 for use in reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells, wherein the antibody or the pharmaceutical composition is to be administered to a subject in need thereof.

20. A therapeutically effective amount of the isolated antibody as in claim 1, or the pharmaceutical composition as in any of claims 7, 9, or 10 for use in reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells, wherein the isolated antibody or the pharmaceutical composition is to be administered to a subject in need thereof.

21. A therapeutically effective amount of the antibody of any one of claims 1 to 3 or the conjugated antibody of claim 4 or 5 or the pharmaceutical composition of any one of claims 7, 9, or 10 for use in reducing immune suppression, wherein the antibody or the conjugated antibody or the pharmaceutical composition is to be administered to a subject in need thereof.

22. A therapeutically effective amount of the antibody of any one of claims 1 to 3 or the pharmaceutical composition of any one of claims 7, 9, or 10 for use in treating, alleviating a symptom of, or delaying the progression of a cancer, wherein the antibody or the pharmaceutical composition is to be administered to a subject in need thereof.

23. Use of an effective amount of the antibody of any one of claims 1 to 3, or the pharmaceutical composition of claims 7, 9, or 10 in the manufacture of a medicament for reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells in a subject in need of reduction thereof.

24. Use of a therapeutically effective amount of the isolated antibody as in claim 1, or the pharmaceutical composition as in any of claims 7, 9, or 10 in the manufacture of a

medicament for reducing binding of a ligand selected from the group consisting of PDL1 or PDL2 to PD-1 on T cells in a subject in need of reduction thereof.

25. Use of a therapeutically effective amount of the antibody of any one of claims 1 to 3 or the conjugated antibody of claim 4 or 5 or the pharmaceutical composition of any one of claims 7, 9, or 10 in the manufacture of a medicament for reducing immune suppression in a subject in need of reduction thereof.

26. The use of claim 25, wherein the immune suppression in a subject in need of reduction thereof is mediated by engagement of PD-1 on T cells to PDL1 or PDL2 on tumor or other immune cells.

27. Use of a therapeutically effective amount of the antibody of any one of claims 1 to 3 or the pharmaceutical composition of any one of claims 7, 9, or 10 in the manufacture of a medicament for treating, alleviating a symptom of, or delaying the progression of a cancer.

28. The use of any one of claims 23 to 27, further comprising an additional agent, wherein the additional agent is a therapeutic agent.

29. The antibody of claim 1, for use in therapy.